Cochrane
Library

Cochrane Database of Systematic Reviews

Haloperidol for psychosis-induced aggression or agitation (rapid tranquillisation) (Review)

Ostinelli EG, Brooke-Powney MJ, Li X, Adams CE

Ostinelli EG, Brooke-Powney MJ, Li X, Adams CE.
Haloperidol for psychosis-induced aggression or agitation (rapid tranquillisation).
Cochrane Database of Systematic Reviews 2017, Issue 7. Art. No.: CD009377.
DOI: [10.1002/14651858.CD009377.pub3](https://doi.org/10.1002/14651858.CD009377.pub3).

www.cochranelibrary.com

TABLE OF CONTENTS

HEADER	1
ABSTRACT	1
PLAIN LANGUAGE SUMMARY	2
SUMMARY OF FINDINGS	4
BACKGROUND	53
OBJECTIVES	53
METHODS	54
RESULTS	59
Figure 1.	60
Figure 2.	61
Figure 3.	65
DISCUSSION	80
AUTHORS' CONCLUSIONS	87
ACKNOWLEDGEMENTS	88
REFERENCES	89
CHARACTERISTICS OF STUDIES	104
DATA AND ANALYSES	178
Analysis 1.1. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 1 Tranquillisation or asleep - asleep.	183
Analysis 1.2. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 2 Repeated need for tranquillisation.	183
Analysis 1.3. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 3 Specific behaviour: 1a. Agitation - various measures. ..	183
Analysis 1.4. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 4 Specific behaviour: 1b. Agitation - average scores - i. up to 2 hours.	184
Analysis 1.5. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 5 Specific behaviour: 1c. Agitation - average scores - ii. up to 24 hours.	185
Analysis 1.6. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 6 Global outcome: 1. Not improved.	185
Analysis 1.7. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 7 Global outcome: 2. Need for benzodiazepine up to 24 hours.	186
Analysis 1.8. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 8 Global outcome: 3a. Average scores - i. up to 2 hours. ..	186
Analysis 1.9. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 9 Global outcome: 3b. Average scores - ii. up to 24 hours.	187
Analysis 1.10. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 10 Mental state: 1a. Average scores - i. up to 2 hours. ...	187
Analysis 1.11. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 11 Mental state: 1b. Average scores - ii. up to 24 hours. .	187
Analysis 1.12. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 12 Adverse effects: 1a. General.	188
Analysis 1.13. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 13 Adverse effects: 1b. General - serious.	188
Analysis 1.14. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 14 Adverse effects: 2a. Specific - arousal level.	189
Analysis 1.15. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 15 Adverse effects: 2b. Specific - cardiovascular - miscellaneous outcomes.	190
Analysis 1.16. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 16 Adverse effects: 2c. Specific - cardiovascular - QTc interval (average change at 24 hours).	191
Analysis 1.17. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 17 Adverse effects: 2d. Specific - movement disorders. .	191
Analysis 1.18. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 18 Adverse effects: 2e. Specific - movement disorders: i. average scores - i. up to 24 hours.	192
Analysis 1.19. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 19 Adverse effects: 2f. Specific - miscellaneous.	192
Analysis 1.20. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 20 Leaving the study early.	193
Analysis 2.1. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIRAZOLE, Outcome 1 Repeated need for rapid tranquillisation: needing additional injection.	198
Analysis 2.2. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIRAZOLE, Outcome 2 Specific behaviour: 1a. Agitation - PANSS-EC response up to 2 hours (at least 40% change on PANSS-EC from baseline).	199
Analysis 2.3. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIRAZOLE, Outcome 3 Specific behaviour: 1b. Agitation - average scores - up to 2 hours.	199
Analysis 2.4. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIRAZOLE, Outcome 4 Global outcome: 1. Need for benzodiazepine.	200
Analysis 2.5. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIRAZOLE, Outcome 5 Global outcome: 2. Average scores - up to 2 hours.	200

Analysis 2.6. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 6 Mental state: 1. Average scores - up to 2 hours.	201
Analysis 2.7. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 7 Adverse effects: 1a. General.	201
Analysis 2.8. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 8 Adverse effects: 1b. General - serious.	202
Analysis 2.9. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 9 Adverse effects: 2a. Specific - arousal level.	203
Analysis 2.10. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 10 Adverse effects: 2b. Specific - cardiac.	204
Analysis 2.11. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 11 Adverse effects: 2c. Specific - gastrointestinal.	204
Analysis 2.12. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 12 Adverse effects: 2d. Specific - movement disorder.	205
Analysis 2.13. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 13 Adverse effects: 2e. Specific - miscellaneous.	207
Analysis 2.14. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 14 Leaving the study early: 1. For specific reasons.	208
Analysis 3.1. Comparison 3 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: b. CHLORPROMAZINE, Outcome 1 Tranquillisation or asleep - asleep.	210
Analysis 3.2. Comparison 3 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: b. CHLORPROMAZINE, Outcome 2 Repeated need for rapid tranquillisation.	211
Analysis 3.3. Comparison 3 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: b. CHLORPROMAZINE, Outcome 3 Global outcome: 1. Not improved.	211
Analysis 3.4. Comparison 3 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: b. CHLORPROMAZINE, Outcome 4 Adverse effects: any serious or specific adverse effects.	212
Analysis 3.5. Comparison 3 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: b. CHLORPROMAZINE, Outcome 5 Leaving the study early.	213
Analysis 4.1. Comparison 4 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: c. DROPERIDOL, Outcome 1 Tranquillisation or asleep: 1. Not asleep.	215
Analysis 4.2. Comparison 4 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: c. DROPERIDOL, Outcome 2 Tranquillisation or asleep: 2. Time to sleep.	215
Analysis 4.3. Comparison 4 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: c. DROPERIDOL, Outcome 3 Repeated need for rapid tranquillisation.	215
Analysis 4.4. Comparison 4 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: c. DROPERIDOL, Outcome 4 Global outcome: 1. Need for benzodiazepine.	216
Analysis 4.5. Comparison 4 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: c. DROPERIDOL, Outcome 5 Adverse effects: 1. General. .	216
Analysis 4.6. Comparison 4 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: c. DROPERIDOL, Outcome 6 Adverse effects: 2a. Specific - arousal level.	217
Analysis 4.7. Comparison 4 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: c. DROPERIDOL, Outcome 7 Adverse effects: 2b. Specific - cardiovascular.	217
Analysis 4.8. Comparison 4 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: c. DROPERIDOL, Outcome 8 Adverse effects: 2c. Specific - movement disorders.	217
Analysis 4.9. Comparison 4 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: c. DROPERIDOL, Outcome 9 Adverse effects: 2d. Specific - miscellaneous.	218
Analysis 5.1. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 1 Tranquillisation or asleep - not asleep up to 24 hours.	222
Analysis 5.2. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 2 Specific behaviour: 2. Aggression - various measures.	222
Analysis 5.3. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 3 Global outcome: 1. General - no change - i. over 24 hours.	223
Analysis 5.4. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 4 Global outcome: 2a. Specific - not sedated - i. by 30 minutes.	223
Analysis 5.5. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 5 Global outcome: 2b. Specific - not sedated - ii. up to 2 hours.	224
Analysis 5.6. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 6 Mental state: 1. Average scores.	224

Analysis 5.7. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 7 Adverse effects: 1. General. ...	224
Analysis 5.8. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 8 Adverse effects: 2a. Specific - anticholinergic.	225
Analysis 5.9. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 9 Adverse effects: 2b. Specific - arousal level.	226
Analysis 5.10. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 10 Adverse effects: 2c. Specific - cardiovascular.	227
Analysis 5.11. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 11 Adverse effects: 2d. Specific - movement disorders.	228
Analysis 5.12. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 12 Adverse effects: 2e. Specific - miscellaneous.	231
Analysis 5.13. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 13 Leaving the study early: 1. For general reasons.	232
Analysis 5.14. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 14 Leaving the study early: 2. Specific reasons.	233
Analysis 6.1. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 1 Tranquillisation or asleep - asleep.	243
Analysis 6.2. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 2 Repeated need for tranquillisation - needing additional injection.	243
Analysis 6.3. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 3 Specific behaviour: 1a. Agitation - various measures.	244
Analysis 6.4. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 4 Specific behaviour: 1b. Agitation - average scores - i. by 30 minutes.	244
Analysis 6.5. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 5 Specific behaviour: 1c. Agitation - average scores - ii. up to 2 hours.	245
Analysis 6.6. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 6 Specific behaviour: 1d. Agitation - average scores - iii. up to 4 hours.	246
Analysis 6.7. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 7 Specific behaviour: 1e. Agitation - average scores - iv. up to 24 hours.	246
Analysis 6.8. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 8 Specific behaviour: 2a. Aggression - average scores - i. up to 2 hours.	247
Analysis 6.9. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 9 Specific behaviour: 2b. Aggression - average scores - ii. up to 4 hours.	247
Analysis 6.10. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 10 Specific behaviour: 2c. Aggression - average scores - iii. up to 24 hours.	248
Analysis 6.11. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 11 Specific behaviour: 3. Hostility.	248
Analysis 6.12. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 12 Global outcome: 1a. General - need for additional measures.	248
Analysis 6.13. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 13 Global outcome: 1b. General - time and doses.	249
Analysis 6.14. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 14 Global outcome: 1c. General - average scores - up to 2 hours.	249
Analysis 6.15. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 15 Global outcome: 1d. General - average scores - over 24 hours.	250
Analysis 6.16. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 16 Global outcome: 2a. Specific - alert - i. up to 2 hours.	250
Analysis 6.17. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 17 Global outcome: 2b. Specific - alert - ii. up to 4 hours.	250
Analysis 6.18. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 18 Global outcome: 2c. Specific - alert - iii. up to 24 hours.	251
Analysis 6.19. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 19 Global outcome: 2d. Specific - tranquil - i. up to 2 hours.	251
Analysis 6.20. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 20 Global outcome: 2e. Specific - tranquil - ii. up to 4 hours.	251

Analysis 6.21. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 21 Global outcome: 2f. Specific - tranquil - iii. up to 24 hours.	252
Analysis 6.22. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 22 Global outcome: 2g. Specific - sedated - i. up to 2 hours.	252
Analysis 6.23. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 23 Global outcome: 2h. Specific - sedated - ii. up to 4 hours.	253
Analysis 6.24. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 24 Global outcome: 2i. Specific - sedated - iii. up to 24 hours.	253
Analysis 6.25. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 25 Global outcome: 2j. Specific - asleep - i. up to 2 hours.	253
Analysis 6.26. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 26 Global outcome: 2k. Specific - asleep - ii. up to 4 hours.	254
Analysis 6.27. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 27 Global outcome: 2l. Specific - asleep - iii. up to 24 hours.	254
Analysis 6.28. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 28 Service use: 1. Average days to discharge.	255
Analysis 6.29. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 29 Service use: 2. Average patient-hours used during 24 hours (skewed data).	255
Analysis 6.30. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 30 Mental state: 1. Various outcomes reported as 'adverse events'.	255
Analysis 6.31. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 31 Mental state: 2a. Average scores - i. by 30 minutes.	256
Analysis 6.32. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 32 Mental state: 2b. Average scores - ii. up to 2 hours.	257
Analysis 6.33. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 33 Mental state: 2c. Average scores - iii. up to 24 hours.	257
Analysis 6.34. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 34 Mental state: 2d. Average scores - iv. over 24 hours.	258
Analysis 6.35. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 35 Adverse effects: 1a. General.	258
Analysis 6.36. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 36 Adverse effects: 1b. General - serious.	258
Analysis 6.37. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 37 Adverse effects: 2a. Specific - anticholinergic.	259
Analysis 6.38. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 38 Adverse effects: 2b. Specific - arousal level - i. various.	260
Analysis 6.39. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 39 Adverse effects: 2c. Specific - arousal level - ii. average scores - i. up to 2 hours.	260
Analysis 6.40. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 40 Adverse effects: 2d. Specific - arousal level - ii. average scores - ii. up to 4 hours.	261
Analysis 6.41. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 41 Adverse effects: 2e. Specific - arousal level - ii. average scores - iii. up to 24 hours.	261
Analysis 6.42. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 42 Adverse effects: 2f. Specific - cardiovascular i. binary.	262
Analysis 6.43. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 43 Adverse effects: 2g. Specific - cardiovascular ii. continuous.	262
Analysis 6.44. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 44 Adverse effects: 2h. Specific - gastrointestinal.	263
Analysis 6.45. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 45 Adverse effects: 2i. Specific - movement disorder - i. various.	263
Analysis 6.46. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 46 Adverse effects: 2j. Specific - movement disorder - ii. average scores - i. up to 24 hours.	264
Analysis 6.47. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 47 Adverse effects: 2k. Specific - movement disorder - iii. average scores - i. up to 24 hours.	265
Analysis 6.48. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 48 Adverse effects: 2l. Specific - movement disorder - vi. average scores - i. over 24 hours.	265

Analysis 6.49. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 49 Adverse effects: 2m. Specific - miscellaneous.	266
Analysis 6.50. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 50 Leaving the study early. ..	266
Analysis 7.1. Comparison 7 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: f. PERPHENAZINE, Outcome 1 Global outcome: No improvement.	269
Analysis 7.2. Comparison 7 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: f. PERPHENAZINE, Outcome 2 Adverse effects: 1. General.	269
Analysis 7.3. Comparison 7 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: f. PERPHENAZINE, Outcome 3 Adverse effects: 2. Specific.	269
Analysis 7.4. Comparison 7 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: f. PERPHENAZINE, Outcome 4 Leaving the study early: 1. Specific reasons.	270
Analysis 8.1. Comparison 8 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: g. QUETIAPINE, Outcome 1 Specific behaviour: 1a. Agitation - average scores - i. up to 24 hours.	272
Analysis 8.2. Comparison 8 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: g. QUETIAPINE, Outcome 2 Specific behaviour: 1b. Agitation - average scores - ii. over 24 hours.	272
Analysis 8.3. Comparison 8 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: g. QUETIAPINE, Outcome 3 Mental state: Average scores - i. over 24 hours.	273
Analysis 8.4. Comparison 8 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: g. QUETIAPINE, Outcome 4 Adverse effects: 1. General. .	273
Analysis 8.5. Comparison 8 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: g. QUETIAPINE, Outcome 5 Adverse effects: 2. Specific - movement disorder - i. average scores.	273
Analysis 8.6. Comparison 8 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: g. QUETIAPINE, Outcome 6 Leaving the study early.	273
Analysis 9.1. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 1 Tranquillisation or asleep - i. by 30 minutes.	277
Analysis 9.2. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 2 Tranquillisation or asleep - ii. up to 2 hours.	278
Analysis 9.3. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 3 Specific behaviour: 1a. Agitation - various measures.	278
Analysis 9.4. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 4 Specific behaviour: 1b. Agitation - average scores - i. up to 2 hours.	278
Analysis 9.5. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 5 Specific behaviour: 2a. Aggression - average scores - i. by 30 minutes.	279
Analysis 9.6. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 6 Specific behaviour: 2b. Aggression - average scores - ii. up to 2 hours.	279
Analysis 9.7. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 7 Global outcome: 1. Binary measures.	279
Analysis 9.8. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 8 Global outcome: 2. Continuous measures.	280
Analysis 9.9. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 9 Adverse effects: 1a. General - one or more adverse effects.	280
Analysis 9.10. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 10 Adverse effects: 2a. Specific - arousal.	281
Analysis 9.11. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 11 Adverse effects: 2b. Specific - cardiovascular - i. binary outcomes.	282
Analysis 9.12. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 12 Adverse effects: 2c. Specific - cardiovascular - ii. pulse rate - i. up to 2 hours.	282
Analysis 9.13. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 13 Adverse effects: 2d. Specific - cardiovascular - ii. pulse rate - ii. up to 24 hours.	283
Analysis 9.14. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 14 Adverse effects: 2e. Specific - movement disorder - i. binary outcomes.	283
Analysis 9.15. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 15 Adverse effects: 2f. Specific - movement disorders - ii. average scores - i. up to 24 hours.	284
Analysis 9.16. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 16 Adverse effects: 2g. Specific - movement disorder - ii. average scores - ii. over 24 hours.	284
Analysis 9.17. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 17 Adverse effects: 2h. Specific - miscellaneous.	285

Analysis 9.18. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 18 Leaving the study early - i. up to 24 hours.	286
Analysis 9.19. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 19 Leaving the study early - ii. over 24 hours.	287
Analysis 10.1. Comparison 10 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: i. THIOTHIXENE, Outcome 1 Repeated need for rapid tranquillisation.	288
Analysis 10.2. Comparison 10 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: i. THIOTHIXENE, Outcome 2 Specific behaviour: 1. Agitation - rated as 'adverse effect'.	289
Analysis 10.3. Comparison 10 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: i. THIOTHIXENE, Outcome 3 Global outcome.	289
Analysis 10.4. Comparison 10 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: i. THIOTHIXENE, Outcome 4 Adverse effects: 1. General - one or more adverse effects.	290
Analysis 10.5. Comparison 10 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: i. THIOTHIXENE, Outcome 5 Adverse effects: 2a. Specific - movement disorders.	290
Analysis 10.6. Comparison 10 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: i. THIOTHIXENE, Outcome 6 Adverse effects: 2b. Specific - others.	291
Analysis 11.1. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 1 Repeated need for tranquillisation - need for additional drugs for tranquillisation.	300
Analysis 11.2. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 2 Specific behaviour: 1a. Agitation - average scores - i. up to 2 hours.	300
Analysis 11.3. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 3 Specific behaviour: 1b. Agitation - average scores - ii. up to 4 hours.	301
Analysis 11.4. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 4 Specific behaviour: 1c. Agitation - average scores - iii. up to 24 hours.	301
Analysis 11.5. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 5 Specific behaviour: 1d. Agitation - average scores - iv. over 24 hours.	302
Analysis 11.6. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 6 Specific behaviour: 2a. Aggression - average scores - i. up to 2 hours.	302
Analysis 11.7. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 7 Specific behaviour: 2b. Aggression - average scores - ii. up to 4 hours.	303
Analysis 11.8. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 8 Specific behaviour: 2c. Aggression - average scores - iii. up to 24 hours.	303
Analysis 11.9. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 9 Global outcome: 1. General - need for additional measures.	303
Analysis 11.10. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 10 Global outcome: 2. Average scores - i. over 24 hours.	304
Analysis 11.11. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 11 Mental state: 1a. Average scores - i. up to 2 hours.	304
Analysis 11.12. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 12 Mental state: 1b. Average scores - ii. up to 4 hours.	305
Analysis 11.13. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 13 Mental state: 1c. Average scores - iii. up to 24 hours.	305
Analysis 11.14. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 14 Mental state: 1d. Average scores - iv. over 24 hours.	305
Analysis 11.15. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 15 Adverse effects: 1a. General - i. up to 24 hours.	306
Analysis 11.16. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 16 Adverse effects: 1b. General - ii. over 24 hours.	306
Analysis 11.17. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 17 Adverse effects: 1c. General - serious.	307
Analysis 11.18. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 18 Adverse effects: 2a. Specific - anticholinergic.	307
Analysis 11.19. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 19 Adverse effects: 2b. Specific - arousal level - i. various.	308
Analysis 11.20. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 20 Adverse effects: 2c. Specific - arousal level - ii. average scores - i. up to 2 hours.	309

Analysis 11.21. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 21 Adverse effects: 2d. Specific - arousal level - ii. average scores - ii. up to 4 hours.	309
Analysis 11.22. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 22 Adverse effects: 2e. Specific - arousal level - ii. average scores - iii. up to 24 hours.	309
Analysis 11.23. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 23 Adverse effects: 2f. Specific - cardiovascular.	310
Analysis 11.24. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 24 Adverse effects: 2g. Specific - gastrointestinal.	311
Analysis 11.25. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 25 Adverse effects: 2h. Specific - hematological tests.	311
Analysis 11.26. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 26 Adverse effects: 2i. Specific - movement disorders - i. binary measures.	313
Analysis 11.27. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 27 Adverse effects: 2j. Specific - movement disorders - ii. average scores - i. over 24 hours.	315
Analysis 11.28. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 28 Adverse effects: 2k. Specific - miscellaneous.	316
Analysis 11.29. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 29 Leaving the study early: 1. For any reason - i. over 24 hours.	316
Analysis 11.30. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 30 Leaving the study early: 2. For specific reasons - i. over 24 hours.	317
Analysis 12.1. Comparison 12 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: k. ZUCLOPENTHIXOL ACETATE, Outcome 1 Repeated need for tranquillisation: more than 3 injections.	318
Analysis 12.2. Comparison 12 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: k. ZUCLOPENTHIXOL ACETATE, Outcome 2 Adverse effects: 1a. Specific - movement disorders.	318
Analysis 12.3. Comparison 12 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: k. ZUCLOPENTHIXOL ACETATE, Outcome 3 Adverse effects: 1b. Specific - miscellaneous.	318
Analysis 12.4. Comparison 12 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: k. ZUCLOPENTHIXOL ACETATE, Outcome 4 Leaving the study early.	319
Analysis 13.1. Comparison 13 HALOPERIDOL vs BENZODIAZEPINE: a. FLUNITRAZEPAM, Outcome 1 Specific behaviour: 1. Aggression - binary measures - i. up to 2 hours.	319
Analysis 13.2. Comparison 13 HALOPERIDOL vs BENZODIAZEPINE: a. FLUNITRAZEPAM, Outcome 2 Global outcome: 1. Need for seclusion or restraint.	320
Analysis 13.3. Comparison 13 HALOPERIDOL vs BENZODIAZEPINE: a. FLUNITRAZEPAM, Outcome 3 Adverse effects: 1. Specific - movement disorders - i. binary measures - i. by 30 minutes.	320
Analysis 14.1. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b. LORAZEPAM, Outcome 1 Tranquillisation or asleep - not asleep.	322
Analysis 14.2. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b. LORAZEPAM, Outcome 2 Repeated need for rapid tranquillisation.	323
Analysis 14.3. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b. LORAZEPAM, Outcome 3 Global outcome: 1. No overall improvement.	323
Analysis 14.4. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b. LORAZEPAM, Outcome 4 Mental state: 1a. Average scores - i. up to 2 hours.	324
Analysis 14.5. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b. LORAZEPAM, Outcome 5 Mental state: 1b. Average scores - ii. up to 4 hours.	324
Analysis 14.6. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b. LORAZEPAM, Outcome 6 Adverse effects: 1a. General.	325
Analysis 14.7. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b. LORAZEPAM, Outcome 7 Adverse effects: 1b. General - serious.	325
Analysis 14.8. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b. LORAZEPAM, Outcome 8 Adverse effects: 2a. Specific - anticholinergic.	325
Analysis 14.9. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b. LORAZEPAM, Outcome 9 Adverse effects: 2b. Specific - arousal.	326
Analysis 14.10. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b. LORAZEPAM, Outcome 10 Adverse effects: 2c. Specific - cardiovascular.	326
Analysis 14.11. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b. LORAZEPAM, Outcome 11 Adverse effects: 2d. Specific - movement disorder.	326

Analysis 15.1. Comparison 15 HALOPERIDOL vs BENZODIAZEPINE: c. MIDAZOLAM, Outcome 1 Tranquillisation or asleep: average times in minutes - (skewed data).	328
Analysis 15.2. Comparison 15 HALOPERIDOL vs BENZODIAZEPINE: c. MIDAZOLAM, Outcome 2 Global outcome: 1. Need for rescue drug.	328
Analysis 15.3. Comparison 15 HALOPERIDOL vs BENZODIAZEPINE: c. MIDAZOLAM, Outcome 3 Adverse effects: 1. General - one or more drug related adverse effect.	329
Analysis 15.4. Comparison 15 HALOPERIDOL vs BENZODIAZEPINE: c. MIDAZOLAM, Outcome 4 Adverse effects: 2. Specific.	329
Analysis 16.1. Comparison 16 HALOPERIDOL vs COMBINATIONS: a. HALOPERIDOL + LEVOMEPROMAZINE, Outcome 1 Global outcome: 1. No overall improvement.	330
Analysis 17.1. Comparison 17 HALOPERIDOL vs COMBINATIONS: b. HALOPERIDOL + LORAZEPAM, Outcome 1 Tranquillisation or asleep - asleep.	331
Analysis 17.2. Comparison 17 HALOPERIDOL vs COMBINATIONS: b. HALOPERIDOL + LORAZEPAM, Outcome 2 Repeated need for rapid tranquillisation.	332
Analysis 17.3. Comparison 17 HALOPERIDOL vs COMBINATIONS: b. HALOPERIDOL + LORAZEPAM, Outcome 3 Global outcome: 1. No overall improvement.	332
Analysis 17.4. Comparison 17 HALOPERIDOL vs COMBINATIONS: b. HALOPERIDOL + LORAZEPAM, Outcome 4 Adverse effects: 1. General.	333
Analysis 17.5. Comparison 17 HALOPERIDOL vs COMBINATIONS: b. HALOPERIDOL + LORAZEPAM, Outcome 5 Adverse effects: 2a. Specific - anticholinergic.	333
Analysis 17.6. Comparison 17 HALOPERIDOL vs COMBINATIONS: b. HALOPERIDOL + LORAZEPAM, Outcome 6 Adverse effects: 2b. Specific - cardiovascular.	333
Analysis 17.7. Comparison 17 HALOPERIDOL vs COMBINATIONS: b. HALOPERIDOL + LORAZEPAM, Outcome 7 Adverse effects: 2c. Specific - movement disorders.	334
Analysis 18.1. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 1 Repeated need for tranquillisation.	337
Analysis 18.2. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 2 Specific behaviour: 1a. Agitation - average scores - i. up to 2 hours.	337
Analysis 18.3. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 3 Specific behaviour: 1b. Agitation - average scores - ii. up to 4 hours.	338
Analysis 18.4. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 4 Specific behaviour: 1c. Agitation - average scores - iii. up to 24 hours.	338
Analysis 18.5. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 5 Specific behaviour: 2a. Aggression - average scores - i. up to 2 hours.	339
Analysis 18.6. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 6 Specific behaviour: 2b. Aggression - average scores - ii. up to 4 hours.	339
Analysis 18.7. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 7 Specific behaviour: 2c. Aggression - average scores - iii. up to 24 hours.	339
Analysis 18.8. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 8 Global outcomes: 1. General - need for additional measures.	340
Analysis 18.9. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 9 Adverse effects: 1b. General - one or more adverse events.	340
Analysis 18.10. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 10 Adverse effects: 1a. General - serious.	340
Analysis 18.11. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 11 Adverse effects: 2a. Specific - arousal - i. various.	341
Analysis 18.12. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 12 Adverse effects: 2b. Specific - arousal - ii. average scores - i. up to 2 hours.	341
Analysis 18.13. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 13 Adverse effects: 2c. Specific - arousal - ii. average scores - ii. up to 4 hours.	341
Analysis 18.14. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 14 Adverse effects: 2d. Specific - arousal - iii. average scores - iii. up to 24 hours.	342
Analysis 18.15. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 15 Adverse effects: 2e. Specific - cardiovascular.	342
Analysis 18.16. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 16 Adverse effects: 2f. Specific - movement disorders.	342

Analysis 19.1. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 1 Tranquillisation or asleep - not tranquil or asleep - i. by 30 minutes.	346
Analysis 19.2. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 2 Tranquillisation or asleep - not tranquil or asleep - ii. up to 2 hours.	346
Analysis 19.3. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 3 Repeated need for tranquillisation - need for additional drugs for tranquillisation.	347
Analysis 19.4. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 4 Specific behaviour: 1a. Agitation - average scores - i. up to 2 hours.	347
Analysis 19.5. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 5 Specific behaviour: 1b. Agitation - average scores - ii. up to 4 hours.	348
Analysis 19.6. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 6 Specific behaviour: 1c. Agitation - average scores - iii. up to 24 hours.	348
Analysis 19.7. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 7 Specific behaviour: 2a. Aggression - average scores - i. up to 2 hours.	349
Analysis 19.8. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 8 Specific behaviour: 2b. Aggression - average scores - ii. up to 4 hours.	349
Analysis 19.9. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 9 Specific behaviour: 2c. Aggression - average scores - iii. up to 24 hours.	349
Analysis 19.10. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 10 Specific behaviour: 2d. Aggression - further aggressive episode - i. up to 24 hours.	350
Analysis 19.11. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 11 Global outcomes: 1. General - need for additional measures.	350
Analysis 19.12. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 12 Global outcomes: 2. Specific.	350
Analysis 19.13. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 13 Adverse effects: 1a. General.	351
Analysis 19.14. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 14 Adverse effects: 1b. General - serious.	351
Analysis 19.15. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 15 Adverse effects: 2a. Specific - arousal - i. various.	352
Analysis 19.16. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 16 Adverse effects: 2b. Specific - arousal - ii. average scores - i. up to 2 hours.	352
Analysis 19.17. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 17 Adverse effects: 2c. Specific - arousal - ii. average scores - ii. up to 4 hours.	352
Analysis 19.18. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 18 Adverse effects: 2d. Specific - arousal - ii. average scores - iii. up to 24 hours.	353
Analysis 19.19. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 19 Adverse effects: 2e. Specific - cardiovascular.	353
Analysis 19.20. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 20 Adverse effects: 2f. Specific - movement disorders.	353
Analysis 19.21. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 21 Service outcomes: 1. Not discharged by 14 days.	354
Analysis 19.22. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 22 Leaving the study early.	354
Analysis 20.1. Comparison 20 HALOPERIDOL vs COMBINATIONS: e. HALOPERIDOL + RISPERIDONE, Outcome 1 Specific behaviour: 1a. Agitation - average scores - i. up to 24 hours.	356
Analysis 20.2. Comparison 20 HALOPERIDOL vs COMBINATIONS: e. HALOPERIDOL + RISPERIDONE, Outcome 2 Specific behaviour: 1b. Agitation - average scores - ii. over 24 hours.	356
Analysis 20.3. Comparison 20 HALOPERIDOL vs COMBINATIONS: e. HALOPERIDOL + RISPERIDONE, Outcome 3 Global outcome: No improvement.	357
Analysis 20.4. Comparison 20 HALOPERIDOL vs COMBINATIONS: e. HALOPERIDOL + RISPERIDONE, Outcome 4 Adverse effects: 1a. Specific - arousal level - i. various.	357
Analysis 20.5. Comparison 20 HALOPERIDOL vs COMBINATIONS: e. HALOPERIDOL + RISPERIDONE, Outcome 5 Adverse effects: 1b. Specific - movement disorders - i. various.	357
Analysis 20.6. Comparison 20 HALOPERIDOL vs COMBINATIONS: e. HALOPERIDOL + RISPERIDONE, Outcome 6 Adverse effects: 1c. Specific - miscellaneous.	358

Analysis 21.1. Comparison 21 HALOPERIDOL vs COMBINATIONS: f. OLANZAPINE + MAGNESIUM VALPROATE, Outcome 1 Specific behaviour: 1. Agitation - average scores - over 24 hours.	359
Analysis 21.2. Comparison 21 HALOPERIDOL vs COMBINATIONS: f. OLANZAPINE + MAGNESIUM VALPROATE, Outcome 2 Global outcome: No improvement.	359
Analysis 21.3. Comparison 21 HALOPERIDOL vs COMBINATIONS: f. OLANZAPINE + MAGNESIUM VALPROATE, Outcome 3 Mental state: Average scores - i. over 24 hours.	359
Analysis 21.4. Comparison 21 HALOPERIDOL vs COMBINATIONS: f. OLANZAPINE + MAGNESIUM VALPROATE, Outcome 4 Adverse effects: General - over 24 hours.	360
Analysis 22.1. Comparison 22 HALOPERIDOL vs COMBINATIONS: g. QUETIAPINE + MAGNESIUM VALPROATE, Outcome 1 Specific behaviour: 1a. Agitation - binary measures.	361
Analysis 22.2. Comparison 22 HALOPERIDOL vs COMBINATIONS: g. QUETIAPINE + MAGNESIUM VALPROATE, Outcome 2 Specific behaviour: 1b. Agitation - average scores - i. over 24 hours.	361
Analysis 23.1. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 1 Tranquillisation or asleep.	364
Analysis 23.2. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 2 Specific behaviour: 1a. Agitation - average scores - i. up to 2 hours.	364
Analysis 23.3. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 3 Specific behaviour: 1b. Agitation - average scores - i. up to 4 hours.	364
Analysis 23.4. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 4 Specific behaviour: 1c. Agitation - average scores - iii. up to 24 hours.	365
Analysis 23.5. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 5 Specific behaviour: 1d. Agitation - average scores - iv. over 24 hours.	365
Analysis 23.6. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 6 Global outcome: No improvement.	366
Analysis 23.7. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 7 Mental state: 1. Average scores - i. over 24 hours.	366
Analysis 23.8. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 8 Adverse effects: 1. General.	366
Analysis 23.9. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 9 Adverse effects: 2a. Specific - arousal - i. various.	367
Analysis 23.10. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 10 Adverse effects: 2b. Specific - cardiovascular - i. various.	367
Analysis 23.11. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 11 Adverse effects: 2c. Specific - movement disorders - i. various.	368
Analysis 23.12. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 12 Adverse effects: 2d. Specific - miscellaneous.	368
Analysis 24.1. Comparison 24 HALOPERIDOL vs COMBINATIONS: i. ZIPRASIDONE + CLONAZEPAM, Outcome 1 Specific behaviour: 1a. Agitation - average scores - i. up to 24 hours.	370
Analysis 24.2. Comparison 24 HALOPERIDOL vs COMBINATIONS: i. ZIPRASIDONE + CLONAZEPAM, Outcome 2 Specific behaviour: 1b. Agitation - average scores - ii. over 24 hours.	370
Analysis 24.3. Comparison 24 HALOPERIDOL vs COMBINATIONS: i. ZIPRASIDONE + CLONAZEPAM, Outcome 3 Global outcome: 1. Average scores - i. over 24 hours.	370
Analysis 24.4. Comparison 24 HALOPERIDOL vs COMBINATIONS: i. ZIPRASIDONE + CLONAZEPAM, Outcome 4 Mental state: 1. Average scores - i. over 24 hours.	371
Analysis 24.5. Comparison 24 HALOPERIDOL vs COMBINATIONS: i. ZIPRASIDONE + CLONAZEPAM, Outcome 5 Leaving the study early.	371
ADDITIONAL TABLES	371
APPENDICES	378
WHAT'S NEW	378
HISTORY	378
CONTRIBUTIONS OF AUTHORS	379
DECLARATIONS OF INTEREST	379
SOURCES OF SUPPORT	379
DIFFERENCES BETWEEN PROTOCOL AND REVIEW	379
INDEX TERMS	379

[Intervention Review]

Haloperidol for psychosis-induced aggression or agitation (rapid tranquillisation)

Edoardo G Ostinelli¹, Melanie J Brooke-Powney², Xue Li³, Clive E Adams⁴

¹Department of Health Sciences, Università degli Studi di Milano, Milan, Italy. ²Department of Clinical Psychology, The University of Manchester, Manchester, UK. ³Systematic Review Solutions Ltd, Nottingham, UK. ⁴Cochrane Schizophrenia Group, The University of Nottingham, Nottingham, UK

Contact address: Edoardo G Ostinelli, Department of Health Sciences, Università degli Studi di Milano, Via Antonio di Rudinì 8, Milan, 20142, Italy. eg.ostinelli@gmail.com.

Editorial group: Cochrane Schizophrenia Group.

Publication status and date: New search for studies and content updated (no change to conclusions), published in Issue 7, 2017.

Citation: Ostinelli EG, Brooke-Powney MJ, Li X, Adams CE. Haloperidol for psychosis-induced aggression or agitation (rapid tranquillisation). *Cochrane Database of Systematic Reviews* 2017, Issue 7. Art. No.: CD009377. DOI: [10.1002/14651858.CD009377.pub3](https://doi.org/10.1002/14651858.CD009377.pub3).

Copyright © 2017 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.

ABSTRACT

Background

Haloperidol used alone is recommended to help calm situations of aggression or agitation for people with psychosis. It is widely accessible and may be the only antipsychotic medication available in limited-resource areas.

Objectives

To examine whether haloperidol alone is an effective treatment for psychosis-induced aggression or agitation, wherein clinicians are required to intervene to prevent harm to self and others.

Search methods

We searched the Cochrane Schizophrenia Group's Study-Based Register of Trials (26th May 2016). This register is compiled by systematic searches of major resources (including AMED, BIOSIS CINAHL, Embase, MEDLINE, PsycINFO, PubMed, and registries of clinical trials) and their monthly updates, handsearches, grey literature, and conference proceedings, with no language, date, document type, or publication status limitations for inclusion of records into the register.

Selection criteria

Randomised controlled trials (RCTs) involving people exhibiting aggression and/or agitation thought to be due to psychosis, allocated rapid use of haloperidol alone (by any route), compared with any other treatment. Outcomes of interest included tranquillisation or asleep by 30 minutes, repeated need for rapid tranquillisation within 24 hours, specific behaviours (threat or injury to others/self), adverse effects. We included trials meeting our selection criteria and providing useable data.

Data collection and analysis

We independently inspected all citations from searches, identified relevant abstracts, and independently extracted data from all included studies. For binary data we calculated risk ratio (RR), for continuous data we calculated mean difference (MD), and for cognitive outcomes we derived standardised mean difference (SMD) effect sizes, all with 95% confidence intervals (CI) and using a fixed-effect model. We assessed risk of bias for the included studies and used the GRADE approach to produce 'Summary of findings' tables which included our pre-specified main outcomes of interest.

Main results

We found nine new RCTs from the 2016 update search, giving a total of 41 included studies and 24 comparisons. Few studies were undertaken in circumstances that reflect real-world practice, and, with notable exceptions, most were small and carried considerable risk of bias. Due to the large number of comparisons, we can only present a summary of main results.

Compared with placebo, more people in the haloperidol group were asleep at two hours (2 RCTs, $n=220$, RR 0.88, 95%CI 0.82 to 0.95, very low-quality evidence) and experienced dystonia (2 RCTs, $n=207$, RR 7.49, 95%CI 0.93 to 60.21, very low-quality evidence).

Compared with aripiprazole, people in the haloperidol group required fewer injections than those in the aripiprazole group (2 RCTs, $n=473$, RR 0.78, 95%CI 0.62 to 0.99, low-quality evidence). More people in the haloperidol group experienced dystonia (2 RCTs, $n=477$, RR 6.63, 95%CI 1.52 to 28.86, very low-quality evidence).

Four trials ($n=207$) compared haloperidol with lorazepam with no significant differences with regard to number of participants asleep at one hour (1 RCT, $n=60$, RR 1.05, 95%CI 0.76 to 1.44, very low-quality of evidence) or those requiring additional injections (1 RCT, $n=66$, RR 1.14, 95%CI 0.91 to 1.43, very low-quality of evidence).

Haloperidol's adverse effects were not offset by addition of lorazepam (e.g. dystonia 1 RCT, $n=67$, RR 8.25, 95%CI 0.46 to 147.45, very low-quality of evidence).

Addition of promethazine was investigated in two trials ($n=376$). More people in the haloperidol group were not tranquil or asleep by 20 minutes (1 RCT, $n=316$, RR 1.60, 95%CI 1.18 to 2.16, moderate-quality evidence). Acute dystonia was too common in the haloperidol alone group for the trial to continue beyond the interim analysis (1 RCT, $n=316$, RR 19.48, 95%CI 1.14 to 331.92, low-quality evidence).

Authors' conclusions

Additional data from new studies does not alter previous conclusions of this review. If no other alternative exists, sole use of intramuscular haloperidol could be life-saving. Where additional drugs are available, sole use of haloperidol for extreme emergency could be considered unethical. Addition of the sedating promethazine has support from better-grade evidence from within randomised trials. Use of an alternative antipsychotic drug is only partially supported by fragmented and poor-grade evidence. Adding a benzodiazepine to haloperidol does not have strong evidence of benefit and carries risk of additional harm.

After six decades of use for emergency rapid tranquillisation, this is still an area in need of good independent trials relevant to real-world practice.

PLAIN LANGUAGE SUMMARY

Haloperidol as a means of calming people who are aggressive or agitated due to psychosis

Aim of review

This review looks at whether haloperidol is effective for treating people who are agitated and aggressive as a result of having psychosis.

Background

People with psychosis may experience hearing voices (hallucinations) or abnormal thoughts (delusions), which can make the person frightened, distressed and agitated (restless, excitable or irritable), leading sometimes to aggressive behaviour. This poses a significant challenge for mental health professionals, who have to quickly choose the best available treatment to prevent the risk of harm to both patient and/or others.

Haloperidol is a medication used for treating people with psychosis that can be taken by mouth or injected. As well as being an antipsychotic (preventing psychosis), it also calms people down or helps them to sleep.

Searches

In 2011 and 2016, the information specialist of the Cochrane Schizophrenia Group searched their specialised register for randomised trials that looked at the effects of giving haloperidol compared with either placebo or other treatments to people who are aggressive or agitated because they were experiencing psychosis.

Results

Forty-one studies are now included in the review but information in these is of low-quality. Main results show that when compared with placebo or no treatment, more people having haloperidol were asleep after two hours. However, evidence is not strong. Results are made more complex by large variety of available treatments (24 comparisons).

Conclusions

The review authors conclude there is weak evidence that haloperidol calms people down and helps manage difficult situations. However, this is not based on good-quality trials and therefore health professionals and people with mental health problems are left without clear evidence-based guidance. In some situations, haloperidol may be the only choice, but this is far from ideal because although haloperidol is effective at calming people down it has side effects (e.g. restlessness, shaking of the head, hands and body, heart problems). These side effects can be just as distressing as psychosis and may act as a barrier that stops people coming back for future treatment. More research is needed to help people consider and understand which medication is better at calming people down; has fewer side effects; works quickly and rapidly; and can be used at lower dosages (or less frequent injections).

This plain language summary is based on a summary written by a consumer Ben Gray from Rethink.

SUMMARY OF FINDINGS

Summary of findings for the main comparison. HALOPERIDOL compared to PLACEBO/NIL for psychosis-induced aggression or agitation

HALOPERIDOL compared with PLACEBO/NIL for psychosis-induced aggression or agitation

Patient or population: patients with psychosis-induced aggression or agitation

Setting: inpatients; multi-centre.

Intervention: HALOPERIDOL

Comparison: PLACEBO/NIL

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	PLACEBO/NIL	HALOPERIDOL				
Tranquillisation or asleep not asleep at 2 hours	Low¹		RR 0.88 (0.82 to 0.95)	220 (2 studies)	⊕⊕⊕⊕ very low 2,3,4	
	965 per 1000	849 per 1000 (791 to 917)				
	Moderate¹					
	980 per 1000	862 per 1000 (804 to 931)				
	High¹					
	995 per 1000	876 per 1000 (816 to 945)				
Repeated need for tranquillisation needing additional injection within 24 hours	Low¹		RR 0.51 (0.42 to 0.62)	660 (4 studies)	⊕⊕⊕⊕ low 2,4	
	400 per 1000	204 per 1000 (168 to 248)				
	Moderate¹					
	600 per 1000	306 per 1000 (252 to 372)				
	High¹					

	800 per 1000	408 per 1000 (336 to 496)			
Specific behaviour - threat or injury to self or others average change score (ABS, high = worse) at 2 hours		The mean specific behaviour - threat or injury to self or others in the intervention groups was 4.48 lower (5.78 to 3.18 lower)	474 (3 studies)	⊕⊕⊕⊕ very low 2,4,5	
Global outcome - overall improvement (not any improvement)	Low¹		RR 0.28 (0.08 to 1.07)	40 (1 study)	⊕⊕⊕⊕ low 2,6
	200 per 1000	56 per 1000 (8 to 414)			
	Moderate¹				
	350 per 1000	98 per 1000 (14 to 724)			
	High¹				
	500 per 1000	140 per 1000 (20 to 1000)			
Adverse effects: Specific - dystonia within 24 hours	Moderate¹		RR 7.49 (0.93 to 60.21)	207 (2 studies)	⊕⊕⊕⊕ very low 2,4,6
	0 per 1000	0 per 1000 (0 to 0)			
	High¹				
	10 per 1000	75 per 1000 (9 to 602)			
Economic outcome - not reported	See comment	See comment	Not estimable	-	See comment No study reported this outcome.

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.
Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.
Very low quality: We are very uncertain about the estimate.

- 1 Control risk - moderate risk is roughly equal to that of the control group.
- 2 Risk of bias: rated 'serious' - method of randomisation not described, allocation concealment not stated, incomplete outcome data, sponsored by drug company.
- 3 Indirectness: rated 'serious' - no measure for tranquil or asleep at 30 minutes, therefore had to use not asleep at 2 hours.
- 4 Publication bias: rated 'strongly suspected' - sponsored by drug company.
- 5 Indirectness: rated 'serious' - not threat of injury to self or others, scale derived data.
- 6 Imprecision: rated 'serious' - 95% confidence interval is wide.

Summary of findings 2. HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE for psychosis-induced aggression or agitation

HALOPERIDOL compared with OTHER ANTIPSYCHOTICS: a. ARIPIPRAZOLE for psychosis-induced aggression or agitation

Patient or population: patients with psychosis-induced aggression or agitation

Setting: multi-centre.

Intervention: HALOPERIDOL

Comparison: OTHER ANTIPSYCHOTICS: a. ARIPIPRAZOLE

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	OTHER ANTIPSY-CHOTICS: a. ARIPIPRAZOLE	HALOPERIDOL				
Tranquillisation or asleep - not reported	See comment	See comment	Not estimable	-	See comment	No study re-ported this out-come.
Repeated need for rapid tranquillisation - needing additional injection	Low ¹		RR 0.78 (0.62 to 0.99)	473 (2 studies)	⊕⊕⊕⊖ low ^{2,3}	
	200 per 1000	156 per 1000 (124 to 198)				
	Moderate ¹					
	400 per 1000	312 per 1000 (248 to 396)				

	High¹			
	650 per 1000	507 per 1000 (403 to 644)		
Specific behaviours - agitation: CABS average change score at 2 hours		The mean specific behaviours - agitation: average change score at 2 hours in the intervention groups was 0.55 lower (2.10 lower to 1.01 higher)	470 (2 studies)	⊕⊕⊕⊕ very low 1,2,4
Global outcomes: need for benzodiazepine ²	Low¹		RR 1.26 (0.74 to 2.16)	477 (2 studies)
	50 per 1000	63 per 1000 (37 to 108)		⊕⊕⊕⊕ very low 2,3,5
	Moderate¹			
	100 per 1000	126 per 1000 (74 to 216)		
	High¹			
	200 per 1000	252 per 1000 (148 to 432)		
Adverse effects: any serious, specific - dystonia during 24 hours	Low⁶		RR 6.63 (1.52 to 28.86)	477 (2 studies)
	0 per 1000	0 per 1000 (0 to 0)		⊕⊕⊕⊕ very low 3,7,8
	Moderate⁶			
	50 per 1000	332 per 1000 (76 to 1000)		
	High⁶			
	100 per 1000	663 per 1000 (152 to 1000)		

Economic outcome - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
--	-------------	-------------	---------------	---	-------------	---------------------------------

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Moderate risk roughly equates with that of the trial control group.

² Risk of bias: rated 'severe' - method of randomisation not described, possible that blinding was broken in Bristol Myers 2005, adverse effects only reported when reported by 5% of participants, sponsored by drug company.

³ Publication bias: rated 'strongly suspected' - sponsored by drug company.

⁴ Indirectness: rated 'serious' - scale-derived data, not threat or injury to self or others as stated in protocol.

⁵ Indirectness: rated 'serious' - no measure for global outcome, overall improvement, therefore global improvement is inferred by need for benzodiazepine.

⁶ Low risk equates with that of trial control group.

⁷ Risk of bias: rated 'serious' - method of randomisation not described, incomplete outcome data, adverse effects only reported when reported by at least 5% of participants, sponsored by drug company.

⁸ Imprecision: rated 'very serious' - 95% confidence intervals are wide.

Summary of findings 3. HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: b. CHLORPROMAZINE for psychosis-induced aggression or agitation

HALOPERIDOL compared with OTHER ANTIPSYCHOTIC: b. CHLORPROMAZINE for psychosis-induced aggression or agitation

Patient or population: patients with psychosis-induced aggression or agitation

Setting: inpatients.

Intervention: HALOPERIDOL

Comparison: OTHER ANTIPSYCHOTIC: b. CHLORPROMAZINE

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	OTHER ANTIPSYCHOTIC: b. CHLORPROMAZINE	HALOPERIDOL

Tranquillisation or asleep Not tranquil or asleep	Low¹	RR 1.93 (1.04 to 3.60)	39 (1 study)	⊕⊕⊕⊕ very low 2,3,4	
	200 per 1000				
	386 per 1000 (208 to 720)				
	Moderate¹				
	500 per 1000				
	965 per 1000 (520 to 1000)				
Repeated need for rapid tranquillisation - more than 1 injection	High¹	RR 1.07 (0.89 to 1.28)	30 (1 study)	⊕⊕⊕⊕ very low 2,3,4	
	700 per 1000				
	1000 per 1000 (728 to 1000)				
	Low¹				
	800 per 1000				
	856 per 1000 (712 to 1000)				
Specific behaviour - threat or injury of harm to self or others - not reported	Moderate¹	Not estimable	-	See comment	No study reported this outcome.
	900 per 1000				
	963 per 1000 (801 to 1000)				
	High¹				
	990 per 1000				
	1000 per 1000 (881 to 1000)				
Global outcome - not any improvement	See comment	RR 0.15 (0.05 to 0.49)	89 (2 studies)	⊕⊕⊕⊕ very low 2,3	
	See comment				
	Low¹				
	1000 per 1000				
	150 per 1000 (50 to 490)				
	Moderate¹				
	300 per 1000				
	45 per 1000 (15 to 147)				

	High¹				
	500 per 1000	75 per 1000 (25 to 245)			
Adverse effects: specific - cardiovascular - hypotension	Low¹		RR 0.51 (0.01 to 2.60)	30 (1 study)	⊕⊕⊕⊕ very low 2,3,4
	50 per 1000	10 per 1000 (0 to 192)			
	Moderate¹				
	150 per 1000	30 per 1000 (1 to 577)			
	High¹				
	250 per 1000	50 per 1000 (2 to 962)			
Adverse effects: specific - central nervous system - seizures	Low¹		RR 0.33 (0.01 to 7.58)	30 (1 study)	⊕⊕⊕⊕ very low 2,3,4
	10 per 1000	3 per 1000 (0 to 76)			
	Moderate¹				
	50 per 1000	17 per 1000 (0 to 379)			
	High¹				
	150 per 1000	50 per 1000 (1 to 1000)			
Economic outcome - not reported	See comment	See comment	Not estimable	-	See comment No study reported this outcome.

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

1 Control risk - moderate risk roughly equates to that of the control group

2 Risk of bias: rated 'serious' - not explicitly described as randomised, missing data was not imputed using appropriate methods such as LOCF, small study.

3 Imprecision: rated 'serious' - small study.

4 Publication bias: rated 'strongly suspected' - small study (15 participants per group).

Summary of findings 4. HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: c. DROPERIDOL for psychosis-induced aggression or agitation

HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: c. DROPERIDOL for psychosis-induced aggression or agitation

Patient or population: psychosis-induced aggression or agitation

Setting: inpatients; emergency room.

Intervention: HALOPERIDOL

Comparison: OTHER ANTIPSYCHOTIC: c. DROPERIDOL

Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	Nº of participants (studies)	Quality of the evidence (GRADE)	Comments
	Risk with OTHER ANTIPSYCHOTIC: c. DROPERIDOL	Risk with HALOPERIDOL				
Tranquillisation or asleep - not asleep up to 2 hours	Study population		RR 1.07 (0.44 to 2.60)	228 (1 RCT)	⊕⊕⊕⊖ moderate ¹	
	82 per 1.000	88 per 1.000 (36 to 213)				
Repeated need for rapid tranquillisation - more than 1 injection	Low		RR 2.38 (1.27 to 4.47)	255 (2 RCTs)	⊕⊕⊕⊖ moderate ²	
	50 per 1.000	119 per 1.000 (64 to 224)				
	Moderate					
	70 per 1.000	167 per 1.000 (89 to 313)				
	High					
	360 per 1.000	857 per 1.000				

	(457 to 1.000)				
Specific behaviour - threat or injury to self or others - not reported	-	-	-	-	No study reported this outcome.
Global outcome - need for benzodiazepine	Study population	RR 0.31 (0.07 to 1.44)	228 (1 RCT)	⊕⊕⊕⊖ moderate ¹	
	59 per 1.000 18 per 1.000 (4 to 85)				
Adverse effects: specific - dystonia	Study population	RR 0.86 (0.11 to 6.56)	255 (2 RCTs)	⊕⊕⊕⊖ moderate ⁵	
	8 per 1.000 7 per 1.000 (1 to 51)				
Economic outcome - not reported	-	-	-	-	No study reported this outcome.

***The risk in the intervention group** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio

GRADE Working Group grades of evidence

High quality: We are very confident that the true effect lies close to that of the estimate of the effect

Moderate quality: We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low quality: Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect

Very low quality: We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect

¹ CI include no effect

² Risk of bias: rated 'serious' - method of randomisation not reported, described as double blind but no further information given regarding rater blinding, small short study.

³ Publication bias: rated 'strongly suspected' - small study.

⁴ Moderate risk roughly equates with that of the trial control group,

⁵ Adverse effects - imprecision - 95% confidence interval.

⁶ Adverse effects - publication bias - small study.

Summary of findings 5. HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: d. LOXAPINE for psychosis-induced aggression or agitation

HALOPERIDOL compared with OTHER ANTIPSYCHOTICS: d. LOXAPINE for psychosis-induced aggression or agitation

Patient or population: patients to psychosis-induced aggression or agitation
Setting: inpatients; emergency room.
Intervention: HALOPERIDOL
Comparison: OTHER ANTIPSYCHOTICS: d. LOXAPINE

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	OTHER ANTIPSY- CHOTICS: d. LOXAPINE	HALOPERIDOL				
Tranquillisation or asleep - not asleep by 12 hours	Low¹		RR 4.31 (0.54 to 34.48)	54 (1 study)	⊕⊕⊕⊕ very low 2,3,4	
	10 per 1000	43 per 1000 (5 to 345)				
	Moderate¹					
	50 per 1000	215 per 1000 (27 to 1000)				
	High¹					
	100 per 1000	431 per 1000 (54 to 1000)				
Global outcome: specific Not sedated at 60 minutes *	Low¹		RR 3.50 (0.86 to 14.18)	30 (1 study)	⊕⊕⊕⊕ very low 2,4,5,6	* data for pre-specified outcome Repeated need for rapid tranquillisation were not reported
	10 per 1000	57 per 1000 (9 to 345)				
	Moderate¹					
	100 per 1000	569 per 1000 (94 to 1000)				
	High¹					
	200 per 1000	1000 per 1000 (188 to 1000)

Specific behaviour - aggression - no overall improvement	Low¹	RR 1.10 (0.69 to 1.76)	30 (1 study)	⊕○○○ very low 2,4,6,7
	500 per 1000	550 per 1000 (345 to 880)		
	Moderate¹			
	650 per 1000	715 per 1000 (448 to 1000)		
	High¹			
	800 per 1000	880 per 1000 (552 to 1000)		
Global outcome: no change at 48 hours CGI	Low¹	RR 0.93 (0.14 to 6.15)	56 (1 study)	⊕⊕○○ low 2,4
	10 per 1000	9 per 1000 (1 to 62)		
	Moderate¹			
	70 per 1000	65 per 1000 (10 to 431)		
	High¹			
	140 per 1000	130 per 1000 (20 to 861)		
Adverse effects: Specific - dystonia between 0-3 days (IM phase)	Low¹	RR 0.94 (0.06 to 13.93)	35 (1 study)	⊕○○○ very low 2,4
	10 per 1000	9 per 1000 (1 to 139)		
	Moderate¹			
	50 per 1000	47 per 1000 (3 to 697)		
	High¹			
	100 per 1000	94 per 1000		

	(6 to 1000)				
Adverse effects: Specific - rigidity between 0-3 days (IM phase)	Low¹		RR 0.88	35	⊕⊕⊕⊕
	750 per 1000	660 per 1000 (487 to 893)	(0.65 to 1.19)	(1 study)	very low ^{2,4,6}
	Moderate¹				
	850 per 1000	748 per 1000 (552 to 1000)			
	High¹				
	950 per 1000	836 per 1000 (617 to 1000)			
Economic outcome - not reported	See comment	See comment	Not estimable	-	See comment No study reported this outcome.

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Moderate risk is roughly equal to that of the control group.

² Risk of bias: rated 'serious' - method of randomisation not described, allocation concealment not stated, staff administering medication were not blind to the intervention, the number of participants reported as leaving the study at certain time points is inconsistent.

³ Indirectness: rated 'serious' - reported at 12 hours rather than 30 minutes as stated in protocol.

⁴ Imprecision: rated 'serious' - 95% confidence interval is wide.

⁵ Indirectness: rated 'serious' - no data on repeated need for rapid tranquillisation, therefore inferred from the data 'not sedated at 60 minutes'.

⁶ Publication bias: rated 'strongly suspected' - small study.

⁷ Indirectness: rated 'serious' - not able to use threat or injury to self or others as stated in the protocol because this was not reported.

Summary of findings 6. HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: e. OLANZAPINE for psychosis induced aggression or agitation

HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: e. OLANZAPINE for psychosis induced aggression or agitation

Patient or population: psychosis induced aggression or agitation

Setting: Emergency room; multi-centre.

Intervention: HALOPERIDOL

Comparison: OTHER ANTIPSYCHOTIC: e. OLANZAPINE

Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	N° of participants (studies)	Quality of the evidence (GRADE)	Comments
	Risk with OTHER ANTIPSYCHOTIC: e. OLANZAPINE	Risk with HALOPERIDOL				
Tranquilisation or asleep assessed with: not asleep up to 2 hours	Low		RR 1.16 (1.02 to 1.32)	257 (1 RCT)	⊕⊕⊕⊕ VERY LOW ^{2 3}	
	200 per 1.000	232 per 1.000 (204 to 264)				
	Moderate					
	700 per 1.000	812 per 1.000 (714 to 924)				
	High					
Repeated need for tranquillisation assessed with: needing an additional injection by 24 hours	Low		RR 1.06 (0.75 to 1.51)	392 (3 RCTs)	⊕⊕⊕⊕ VERY LOW ²	
	50 per 1.000	53 per 1.000 (38 to 76)				
	Moderate					
	200 per 1.000	212 per 1.000 (150 to 302)				
	High					
	350 per 1.000	371 per 1.000 (262 to 529)

Specific behaviour - threat or injury to self or others assessed with: less than 40% reduction in PANSS-EC rated at 2 hours	Low		RR 0.96 (0.58 to 1.58)	45 (1 RCT)	⊕⊕⊕⊕ VERY LOW ^{2 5}
	200 per 1.000	192 per 1.000 (116 to 316)			
	Moderate				
	400 per 1.000	384 per 1.000 (232 to 632)			
	High				
	600 per 1.000	576 per 1.000 (348 to 948)			
Global outcome - need for benzodiazepine during 24 hours	Low		RR 1.05 (0.63 to 1.74)	343 (2 RCTs)	⊕⊕⊕⊕ VERY LOW ^{6 7}
	50 per 1.000	53 per 1.000 (32 to 87)			
	Moderate				
	150 per 1.000	158 per 1.000 (95 to 261)			
	High				
	250 per 1.000	263 per 1.000 (158 to 435)			
Adverse effects: Specific - dystonia during 24 hours	Low		RR 12.92 (1.67 to 99.78)	343 (2 RCTs)	⊕⊕⊕⊕ VERY LOW ^{6 7}
	0 per 1.000	0 per 1.000 (0 to 0)			
	Moderate				
	0 per 1.000	0 per 1.000 (0 to 0)			
	High				
	5 per 1.000	65 per 1.000			

	(8 to 499)					
Economic outcome - not reported	-	-	-	-	-	No studies reported this outcome.

***The risk in the intervention group** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio; **OR:** Odds ratio;

GRADE Working Group grades of evidence

High quality: We are very confident that the true effect lies close to that of the estimate of the effect

Moderate quality: We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low quality: Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect

Very low quality: We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect

¹ Moderate risk is roughly equal to that of the control group.

² Risk of bias: rated 'very serious' - method of randomisation not reported, allocation concealment not stated, described as double blind but no further details are given regarding blinding, selective reporting, sponsored by drug company.

³ Indirectness: rated 'serious' - tranquil at 30 minutes is not measured, therefore used not asleep at 2 hour.

⁴ Publication bias: rated 'strongly suspected' - sponsored by drug company.

⁵ Indirectness: rated 'serious' - not threat or injury to self or others as stated in protocol, therefore inferred further aggressive behaviour by less than a 40% reduction in PANSS-EC score.

⁶ Risk of bias: rated 'serious' - method of randomisation is not reported, allocation concealment not stated, described as double blind but no further information given, incomplete outcome data, selective reporting, sponsored by drug company.

⁷ Imprecision: rated 'serious' - 95% confidence interval is wide.

Summary of findings 7. HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: f. PERPHENAZINE for psychosis-induced aggression or agitation

HALOPERIDOL compared with OTHER ANTIPSYCHOTIC: f. PERPHENAZINE for psychosis-induced aggression or agitation

Patient or population: patients with psychosis-induced aggression or agitation

Setting: inpatients.

Intervention: HALOPERIDOL

Comparison: OTHER ANTIPSYCHOTIC: f. PERPHENAZINE

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				

	OTHER ANTIPSY- CHOTIC: f. PER- PHENAZINE	HALOPERIDOL				
Tranquillisation or asleep - not reported	See comment	See comment	Not estimable	-	See comment	No study re- ported this out- come.
Repeated need for rapid tran- quillisation - not reported	See comment	See comment	Not estimable	-	See comment	No study re- ported this out- come.
Specific behaviour - injury or threat of self to others - not re- ported	See comment	See comment	Not estimable	-	See comment	No study re- ported this out- come.
Global outcome: general - no improvement	Low¹		RR 0.46 (0.04 to 4.68)	44 (1 study)	⊕⊕⊕⊕ very low ^{2,3}	
	10 per 1000	5 per 1000 (0 to 47)				
	Moderate¹					
	100 per 1000	46 per 1000 (4 to 468)				
	High¹					
	200 per 1000	92 per 1000 (8 to 936)				
Adverse effect: Specific - hy- potensive episode	Low¹		RR 0.31 (0.01 to 7.12)	44 (1 study)	⊕⊕⊕⊕ very low ^{2,3,4}	
	10 per 1000	3 per 1000 (0 to 71)				
	Moderate¹					
	50 per 1000	15 per 1000 (0 to 356)				
	High¹					
	100 per 1000	31 per 1000				

	(1 to 712)				
Adverse effect - discontinued due to EPS	Low¹		RR 1.83	44	⊕○○○
			(0.18 to 18.7)	(1 study)	very low 2,3,5
	10 per 1000	18 per 1000			
		(2 to 187)			
	Moderate¹				
	50 per 1000	92 per 1000			
		(9 to 935)			
	High¹				
	100 per 1000	183 per 1000			
		(18 to 1000)			
Economic outcome - not re-reported	See comment	See comment	Not estimable	-	See comment
					No study re-reported this outcome.

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Moderate risk is roughly equal to that of the trial control group.

² Risk of bias: rated 'serious' - method or randomisation not described, allocation concealment not stated, described as double-blind but no further details given regarding rater blinding, author suspects that only the adverse effects severe enough to warrant discontinuation were reported.

³ Publication bias: rated 'strongly suspected' - small study.

⁴ Imprecision: rated 'serious' - 95% confidence interval is wide, small study.

⁵ Indirectness: rated 'serious' - not specific adverse effect - only have data for the number of people discontinued due to EPS, not the number of people who had EPS in general.

Summary of findings 8. HALOPERIDOL compared to OTHER ANTIPSYCHOTICS: g. QUETIAPINE for psychosis-induced aggression or agitation

HALOPERIDOL compared to OTHER ANTIPSYCHOTICS: g. QUETIAPINE for psychosis-induced aggression or agitation

Patient or population: psychosis-induced aggression or agitation

Setting: inpatients.
Intervention: HALOPERIDOL
Comparison: OTHER ANTIPSYCHOTICS: g. QUETIAPINE

Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	Nº of participants (studies)	Quality of the evidence (GRADE)	Comments
	Risk with OTHER ANTIPSYCHOTICS: g. QUETIAPINE	Risk with HALOPERIDOL				
Tranquillisation or asleep - not reported	-	-	-	-	-	No study reported this outcome.
Repeated need for tranquillisation - not reported	-	-	-	-	-	No study reported this outcome.
Specific behaviour - agitation assessed with: change score at 24 hours on the PANSS-EC		MD 0.10 higher (0.56 lower to 0.76 higher)	-	80 (1 RCT)	⊕⊕⊕⊕ low ¹	
Global outcome - not reported	-	-	-	-	-	No study reported this outcome.
Adverse effects assessed with: one or more drug-related adverse events	Low		RR 1.81 (1.19 to 2.77)	80 (1 RCT)	⊕⊕⊕⊕ low ¹	
	200 per 1.000	362 per 1.000 (238 to 554)				
	Moderate					
	400 per 1.000	724 per 1.000 (476 to 1.000)				
	High					
	600 per 1.000	1000 per 1.000 (714 to 1.000)				
Economic outcome - not reported	-	-	-	-	-	No study reported this outcome.

*The risk in the intervention group (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: We are very confident that the true effect lies close to that of the estimate of the effect

Moderate quality: We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low quality: Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect

Very low quality: We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect

¹ Imprecision: rated 'very serious' - small study, 95% CI are wide.

Summary of findings 9. HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: h. RISPERIDONE for psychosis-induced aggression or agitation

HALOPERIDOL compared with OTHER ANTIPSYCHOTIC: g. RISPERIDONE for psychosis-induced aggression or agitation

Patient or population: patients with psychosis-induced aggression or agitation

Setting: inpatients; emergency room; multi-centre.

Intervention: HALOPERIDOL

Comparison: OTHER ANTIPSYCHOTIC: g. RISPERIDONE

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	OTHER ANTIPSYCHOTIC: g. RISPERIDONE	HALOPERIDOL				
Tranquillisation or asleep - not asleep	Low ¹		RR 0.84 (0.74 to 0.95)	162 (1 study)	⊕⊕⊕⊖ moderate ²	
	700 per 1000	588 per 1000 (518 to 665)				
	Moderate ¹					
	900 per 1000	756 per 1000 (666 to 855)				
	High ¹					
	1000 per 1000	840 per 1000 (740 to 950)

Repeated need for rapid tranquillisation - needing additional benzodiazepine	Low	RR 0.98 (0.65 to 1.47)	286 (2 studies)	⊕⊕○○ low 2,3
	100 per 1000			
	98 per 1000 (65 to 147)			
	Moderate			
	200 per 1000			
	196 per 1000 (130 to 294)			
Specific behaviours - agitation >50% reduction in PANSS-EC score Follow-up: 1 days	High	RR 1.15 (0.6 to 2.22)	124 (1 study)	⊕⊕○○ low 2,4
	300 per 1000			
	294 per 1000 (195 to 441)			
	Low¹			
	50 per 1000			
	58 per 1000 (30 to 111)			
Global outcome - rated as severe - CGI-S Follow-up: 1 days	Moderate¹	RR 0.89 (0.51 to 1.58)	162 (1 study)	⊕○○○ very low 2,3,4
	200 per 1000			
	230 per 1000 (120 to 444)			
	High¹			
	350 per 1000			
	402 per 1000 (210 to 777)			
	Low¹	RR 0.89 (0.51 to 1.58)	162 (1 study)	⊕○○○ very low 2,3,4
	100 per 1000			
	89 per 1000 (51 to 158)			
	Moderate¹			
	200 per 1000			
	178 per 1000 (102 to 316)			
	High¹			
	300 per 1000			
	267 per 1000			

(153 to 474)					
Adverse effects: Specific - EPS during 24 hours	Low¹		RR 1.6 (0.55 to 4.62)	124 (1 study)	⊕⊕⊕⊕ low 2,4
	10 per 1000	16 per 1000 (6 to 46)			
	Moderate¹				
	100 per 1000	160 per 1000 (55 to 462)			
	High¹				
	200 per 1000	320 per 1000 (110 to 924)			
Adverse effects: Specific - acute dystonia during 24 hours	Low¹		RR 5 (0.24 to 102.07)	286 (2 studies)	⊕⊕⊕⊕ very low 2,3,4
	10 per 1000	50 per 1000 (2 to 1000)			
	Moderate¹				
	20 per 1000	100 per 1000 (5 to 1000)			
	High¹				
	30 per 1000	150 per 1000 (7 to 1000)			
Economic outcome - not reported	See comment	See comment	Not estimable	-	See comment No study reported this outcome.

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

- ¹ Moderate risk is roughly equal to that of the trial control group.
- ² Risk of bias: rated 'serious' - method of randomisation not described, single blind, sponsored by drug company.
- ³ Imprecision: rated 'serious' - few data and confidence intervals wide.
- ⁴ Publication bias: rated 'strongly suspected' - sponsored by drug company.

Summary of findings 10. HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: i. THIOTHIXENE for psychosis-induced aggression or agitation

HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: h. THIOTHIXENE for psychosis-induced aggression or agitation

Patient or population: patients with psychosis-induced aggression or agitation

Setting: inpatients; emergency room.

Intervention: HALOPERIDOL

Comparison: OTHER ANTIPSYCHOTIC: h. THIOTHIXENE

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	OTHER ANTIPSYCHOTIC: h. THIOTHIXENE	HALOPERIDOL				
Tranquillisation or asleep - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Repeated need for rapid tranquillisation	Low ¹		RR 1.07 (0.89 to 1.28)	30 (1 study)	⊕⊕⊕⊕ very low ^{2,3}	
	850 per 1000	910 per 1000 (756 to 1000)				
	Moderate ¹					
	900 per 1000	963 per 1000 (801 to 1000)				
	High ¹					
	950 per 1000	1000 per 1000 (845 to 1000)				

Specific behaviour - threat or injury to self or others - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Global outcome No improvement at 1 hour	Low¹		RR 0.50 (0.14 to 1.84)	44 (1 study)	⊕⊕⊕⊕ very low 2,3,4	
	50 per 1000	25 per 1000 (7 to 92)				
	Moderate¹					
	250 per 1000	125 per 1000 (35 to 460)				
	High¹					
	450 per 1000	225 per 1000 (63 to 828)				
Adverse effects: specific - hypotensive episode	Low		Not estimable	30 (1 study)	⊕⊕⊕⊕ very low 2,3,4	
	0 per 1000	0 per 1000 (0 to 0)				
Adverse effects: specific - tachycardia	Low¹		RR 0.28 (0.01 to 6.52)	44 (1 study)	⊕⊕⊕⊕ very low 2,3,4	
	10 per 1000	3 per 1000 (0 to 65)				
	Moderate¹					
	50 per 1000	14 per 1000 (0 to 326)				
	High¹					
	100 per 1000	28 per 1000 (1 to 652)				
Economic outcome - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Control risk - moderate risk roughly equates to that of the trial control group.

² Risk of bias: rated 'serious' - method of sequence generation not clear, allocation concealment not reported, small study, sponsored by drug company.

³ Publication bias: rated 'strongly suspected' - small study, sponsored by drug company.

⁴ Imprecision: rated 'serious' - small study, wide confidence intervals.

Summary of findings 11. HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE for psychosis-induced aggression or agitation

HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: i. ZIPRASIDONE for psychosis-induced aggression or agitation

Patient or population: psychosis-induced aggression or agitation

Setting: inpatients; emergency room; multi-centre.

Intervention: HALOPERIDOL

Comparison: OTHER ANTIPSYCHOTIC: i. ZIPRASIDONE

Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	N° of participants (studies)	Quality of the evidence (GRADE)	Comments
	Risk with OTHER ANTIPSYCHOTIC: i. ZIPRASIDONE	Risk with HALOPERIDOL				
Tranquilisation or asleep - not reported	-	-	-	-	-	No trial reported this outcome.
Repeated need for tranquilisation assessed with: need for additional drugs for tranquillisation up to 24 hours	Study population		RR 1.64 (1.07 to 2.53)	60 (1 RCT)	⊕⊕⊕⊕ very low ^{1 2}	
	467 per 1.000	765 per 1.000 (499 to 1.000)				
Specific behaviour - agitation assessed with: average endpoint scores at 2 hours on the PANSS-EC		MD 0.06 higher (1.13 lower to 1.25 higher)	-	231 (1 RCT)	⊕⊕⊕⊕ low ^{3 4}	

follow up: 72 hours					
Global outcome assessed with: CGI-S - average change score at 72 hours.	MD 0.34 higher (0.13 higher to 0.55 higher)	-	132 (1 RCT)	⊕⊕⊕⊕ very low ⁵	
Adverse effects: Specific - dystonia during 72 hours	Low	RR 10.26 (1.67 to 63.17)	508 (2 RCTs)	⊕⊕⊕⊕ very low ^{5 7}	
	2 per 1.000 21 per 1.000 (3 to 126)				
	Moderate				
	4 per 1.000 41 per 1.000 (7 to 253)				
	High				
Adverse effects: Specific - clinically significant abnormal ECG during 72 hours	Study population	RR 1.01 (0.60 to 1.71)	376 (1 RCT)	⊕⊕⊕⊕ very low ^{3 9}	
	127 per 1.000 128 per 1.000 (76 to 217)				
Economic outcome - not reported	-	-	-	-	No trial reported this outcome.

***The risk in the intervention group** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio

GRADE Working Group grades of evidence

High quality: We are very confident that the true effect lies close to that of the estimate of the effect

Moderate quality: We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low quality: Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect

Very low quality: We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect

¹ Risk of bias: rated 'serious' - method of allocation concealment not stated, blinding of participants ineffective

² Imprecision: rated 'very serious' - less than 100 patients, 95% CI extends beyond the no effect point

- ³ Risk of bias: rated 'serious' - method of randomisation is not described, allocation concealment is not stated, single-blind.
⁴ Indirectness: rated 'serious' - not threat or injury to self or others, therefore had to use scale derived data for agitation.
⁵ Risk of bias: rated 'very serious' - allocation of concealment not stated, open trial, adverse effects only reported where they occurred in $\geq 10\%$ of people, sponsored by drug company.
⁶ Publication bias: rated 'strongly suspected' - sponsored by drug company.
⁷ Imprecision: rated 'serious' - 95% confidence intervals are wide.
⁸ Moderate risk roughly is roughly equal to that of the control group.
⁹ Indirectness: rated 'serious' - abnormal ECG - not necessarily a serious adverse effect.

Summary of findings 12. HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: k. ZUCLOPENTHIXOL ACETATE for psychosis-induced aggression or agitation

HALOPERIDOL compared with OTHER ANTIPSYCHOTICS: j. ZUCLOPENTHIXOL ACETATE for psychosis-induced aggression or agitation

Patient or population: patients with psychosis-induced aggression or agitation

Setting: inpatients.

Intervention: HALOPERIDOL

Comparison: OTHER ANTIPSYCHOTICS: j. ZUCLOPENTHIXOL ACETATE

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	OTHER ANTIPSYCHOTICS: j. ZUCLOPENTHIXOL ACETATE	HALOPERIDOL				
Tranquillisation or asleep - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Repeated need for rapid tranquillisation - more than 3 injections	Low¹		RR 2.54 (1.19 to 5.46)	70 (1 study)	⊕⊕⊕⊕ very low ^{2,3,4}	
	50 per 1000	127 per 1000 (60 to 273)				
	Moderate¹					
	200 per 1000	508 per 1000 (238 to 1000)				
	High¹					

	350 per 1000	889 per 1000 (417 to 1000)				
Specific behaviour - threat or injury to self or others - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Global outcome - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Adverse effects: specific - tremor by 7 days	Low¹		RR 4.16 (0.93 to 18.62)	70 (1 study)	⊕⊕⊕⊕ very low 2,4,5	
	10 per 1000	42 per 1000 (9 to 186)				
	Moderate¹					
	50 per 1000	208 per 1000 (47 to 931)				
	High¹					
	100 per 1000	416 per 1000 (93 to 1000)				
Economic outcome - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Moderate risk is roughly equal to that of the control group.

² Risk of bias: rated 'serious' - method of randomisation not reported, open label, author suspects that rater was only blind for some outcomes.

³ Indirectness: rated 'serious' - measures more than 3 injections, not just more than 1 injection.

⁴ Imprecision: rated 'serious' - small study.

⁵ Indirectness: rated 'serious' - no serious specific adverse effect, therefore used tremor.

Summary of findings 13. HALOPERIDOL compared to BENZODIAZEPINE: a. FLUNITRAZEPAM for psychosis-induced aggression or agitation

HALOPERIDOL compared with BENZODIAZEPINE: a. FLUNITRAZEPAM for psychosis-induced aggression or agitation

Patient or population: patients with psychosis-induced aggression or agitation

Setting: inpatients.

Intervention: HALOPERIDOL

Comparison: BENZODIAZEPINE: a. FLUNITRAZEPAM

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	BENZODIAZEPINE: a. FLUNITRAZEPAM	HALOPERIDOL				
Tranquillisation or asleep - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Repeated need for tranquillisation within 24 hours - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Specific behaviours - aggression 50% reduction in OAS score at 90 minutes	Low¹		RR 1.15 (0.86 to 1.55)	28 (1 study)	⊕⊕⊕⊕ very low 2,3,4	
	650 per 1000	747 per 1000 (559 to 1000)				
	Moderate¹					
	800 per 1000	920 per 1000 (688 to 1000)				
	High¹					
	950 per 1000	1000 per 1000 (817 to 1000)				
Global outcome - need for restraint or seclusion	See comment	See comment	Not estimable	28 (1 study)	⊕⊕⊕⊕ very low 2,4,5,6	No events in either group.

Adverse effects: specific - EPS	See comment	See comment	Not estimable	28 (1 study)	⊕⊕⊕⊕ very low 2,4,6,7	No events in either group.
Economic outcome - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

1 Moderate risk is roughly equal to that of the control group.

2 Risk of bias: rated 'serious' - allocation concealment not stated, described as double-blind, however the blinding of participants and personnel are not reported, it is not clear whether all side effects are reported, small short study.

3 Indirectness: rated 'serious' - not threat or injury to self or others, inferred from OAS rating scale.

4 Publication bias: rated 'strongly suspected' - small study.

5 Indirectness: rated 'serious' - there is no measure for overall improvement, therefore inferred no overall improvement by need for restraints.

6 Imprecision: rated 'serious' - small study.

7 Indirectness: rated 'serious' - EPS can refer to a number of different symptoms rather than one specific side effect.

Summary of findings 14. HALOPERIDOL compared to BENZODIAZEPINE: b. LORAZEPAM for psychosis-induced aggression or agitation

HALOPERIDOL compared with BENZODIAZEPINE: b. LORAZEPAM for psychosis-induced aggression or agitation

Patient or population: patients with psychosis-induced aggression or agitation

Setting: inpatients; emergency room.

Intervention: HALOPERIDOL

Comparison: BENZODIAZEPINE: b. LORAZEPAM

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	BENZODIAZEPINE: b. LORAZEPAM	HALOPERIDOL				
Tranquillisation or asleep	Low ¹		RR 1.05	60	⊕⊕⊕⊕

not asleep at 60 minutes	500 per 1000	525 per 1000 (380 to 720)	(0.76 to 1.44)	(1 study)	very low 2,3,4,5	
	Moderate ¹					
	700 per 1000	735 per 1000 (532 to 1000)				
	High ¹					
	900 per 1000	945 per 1000 (684 to 1000)				
Repeated need for rapid tranquillisation more than 1 injection	Low ¹		RR 1.14 (0.91 to 1.43)	66 (1 study)	⊕⊕⊕⊕ very low 4,5,6	
	500 per 1000	570 per 1000 (455 to 715)				
	Moderate ¹					
	750 per 1000	855 per 1000 (683 to 1000)				
	High ¹					
	900 per 1000	1000 per 1000 (819 to 1000)				
Specific behaviour - threat or injury to self or others within 24 hours - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Global outcome: no overall improvement - at 60 minutes	Low ¹		RR 1.64 (0.54 to 5.03)	44 (1 study)	⊕⊕⊕⊕ very low 2,4	
	50 per 1000	82 per 1000 (27 to 252)				
	Moderate ¹					
	150 per 1000	246 per 1000 (81 to 755)				
	High ¹					

	100 per 1000	164 per 1000 (54 to 503)				
Adverse effects: specific - dys- tonia (only reported if occurred in ≥9% during 24 hours)	Low¹		RR 3.54 (0.42 to 30.03)	66 (1 study)	⊕⊕⊕⊕ very low 4,5,6	
	10 per 1000	35 per 1000 (4 to 300)				
	Moderate¹					
	30 per 1000	106 per 1000 (13 to 901)				
	High¹					
	50 per 1000	177 per 1000 (21 to 1000)				
Adverse effects: specific - hy- pertonia/rigidity (only report- ed if occurred in ≥9%) during 24 hours	Low⁷		RR 6.22 (0.33 to 115.91)	66 (1 study)	⊕⊕⊕⊕ very low 4,5,6	
	0 per 1000	0 per 1000 (0 to 0)				
	Moderate⁷					
	50 per 1000	311 per 1000 (17 to 1000)				
	High⁷					
	100 per 1000	622 per 1000 (33 to 1000)				
Economic outcome - not re- ported	See comment	See comment	Not estimable	-	See comment	No study re- ported this out- come.

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.
Very low quality: We are very uncertain about the estimate.

- 1 Moderate risk is roughly equal to that of the trial control group.
- 2 Risk of bias: rated 'very serious' - not explicitly described as randomised, allocation concealment not stated, incomplete outcome data, sponsored by drug company.
- 3 Indirectness: rated 'serious' - no measure for tranquillisation or asleep at 30 minutes, therefore had to use 60 minutes.
- 4 Imprecision: rated 'serious' - small study.
- 5 Publication bias: rated 'strongly suspected' - small study, sponsored by drug company.
- 6 Risk of bias: rated 'serious' - allocation concealment is not stated, incomplete outcome data, selective reporting, sponsored by drug company.
- 7 Low risk is roughly equal to that of the trial control group.

Summary of findings 15. HALOPERIDOL compared to BENZODIAZEPINE: c. MIDAZOLAM for psychosis-induced aggression or agitation

HALOPERIDOL compared with BENZODIAZEPINE: c. MIDAZOLAM for psychosis-induced aggression or agitation

Patient or population: patients with psychosis-induced aggression or agitation

Setting: emergency room.

Intervention: HALOPERIDOL

Comparison: BENZODIAZEPINE: c. MIDAZOLAM

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	BENZODIAZEPINE: c. MIDAZOLAM	HALOPERIDOL				
Tranquillisation or asleep - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Repeated need for rapid tranquillisation - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Specific behaviour - threat or injury to self or others - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Global outcome Need for rescue drug	Low¹		RR 1.14 (0.46 to 2.87)	84 (1 study)	⊕⊕⊕⊖ low 2,3	
	50 per 1000	57 per 1000				

	(23 to 143)				
	Moderate¹				
	150 per 1000	171 per 1000 (69 to 430)			
	High¹				
	250 per 1000	285 per 1000 (115 to 717)			
Adverse effects - general - one or more drug-related adverse effect	Low⁴		RR 5.00 (0.25 to 101.11)	84 (1 study)	⊕⊕⊕⊖ low 3,5
	0 per 1000	0 per 1000 (0 to 0)			
	Moderate⁴				
	50 per 1000	250 per 1000 (12 to 1000)			
	High⁴				
	100 per 1000	500 per 1000 (25 to 1000)			
Economic outcome - not reported	See comment	See comment	Not estimable	-	See comment No study re-ported this out-come.

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Medium risk is roughly equal to that of the trial control group.

² Indirectness: rated 'serious' - no measure for overall improvement, no overall improvement inferred from need for rescue drug.

- ³ Imprecision: rated 'serious' - 95% confidence interval is wide.
⁴ Low risk is roughly equal to that of the trial control group.
⁵ Indirectness: rated 'serious' - no serious adverse effect, therefore used one or more adverse effects.

Summary of findings 16. HALOPERIDOL compared to COMBINATIONS: a. HALOPERIDOL + LEVOMEPRMAZINE for psychosis-induced aggression or agitation

HALOPERIDOL compared with 1. COMBINATIONS - HALOPERIDOL + LEVOMEPRMAZINE for psychosis-induced aggression or agitation

Patient or population: patients with psychosis-induced aggression or agitation

Setting: inpatients.

Intervention: HALOPERIDOL

Comparison: COMBINATIONS: a. HALOPERIDOL + LEVOMEPRMAZINE

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	COMBINATIONS: a. HALOPERIDOL + LEVOMEPRMAZINE	HALOPERIDOL				
Tranquillisation or asleep - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Repeated need for tranquillisation - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Specific behaviours - threat or injury to self or others - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Global outcome: general - no overall improvement	Low¹		RR 8.18 (0.5 to 133.66)	19 (1 study)	⊕⊕⊕⊕ very low 2,3,4	
	0 per 1000	0 per 1000 (0 to 0)				
	Moderate¹					
	200 per 1000	1000 per 1000 (100 to 1000)				
	High¹					

	400 per 1000	1000 per 1000 (200 to 1000)				
Adverse effect: any serious, specific - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Economic outcome - not measured	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Low risk is roughly equal to that of the trial control group.

² Risk of bias: rated 'very serious' - method of randomisation not described, allocation concealment not stated, open trial, no reporting of the presence or absence of adverse effects in the results, small study.

³ Imprecision: rated 'very serious' - very small study (n = 19), 95% confidence interval is wide.

⁴ Publication bias: rated 'strongly suspected' - small study.

Summary of findings 17. HALOPERIDOL compared to COMBINATIONS: b. HALOPERIDOL + LORAZEPAM for psychosis-induced aggression or agitation

HALOPERIDOL compared with COMBINATIONS: b. HALOPERIDOL + LORAZEPAM for psychosis-induced aggression or agitation

Patient or population: patients with psychosis-induced aggression or agitation

Setting: inpatients; emergency room.

Intervention: HALOPERIDOL

Comparison: COMBINATIONS: b. HALOPERIDOL + LORAZEPAM

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	COMBINATIONS: b. HALOPERIDOL + LO- RAZEPAM	HALOPERIDOL

Tranquillisation or asleep - not asleep by 3 hours	Low¹	RR 1.83 (1.11 to 3.02)	67 (1 study)	⊕⊕⊕⊕ very low 2,3,4,5
	200 per 1000			
	366 per 1000 (222 to 604)			
	Moderate¹			
	400 per 1000			
Repeated need for rapid tranquillisation - needing additional injection during 24 hours	732 per 1000 (444 to 1000)	RR 1.05 (0.87 to 1.27)	67 (1 study)	⊕⊕⊕⊕ very low 2,4,5
	High¹			
	600 per 1000			
	1000 per 1000 (666 to 1000)			
	Low¹			
Specific behaviour - threat or injury to self or others - not reported	750 per 1000	Not estimable	-	See comment
	788 per 1000 (653 to 952)			
	Moderate¹			
	850 per 1000			
	892 per 1000 (740 to 1000)			
Global outcome: no overall improvement - at 30 minutes	High¹	RR 2.67 (1.25 to 5.68)	45 (1 study)	⊕⊕⊕⊕ very low 4,6
	950 per 1000			
	997 per 1000 (827 to 1000)			
	Low¹			
	50 per 1000			
	134 per 1000 (62 to 284)			
	Moderate¹			
	250 per 1000			
	668 per 1000 (312 to 1000)			

	High¹				
	450 per 1000	1000 per 1000 (562 to 1000)			
Adverse effects: Specific - dystonia (only reported if occurred in ≥9%) during 24 hours	Low⁷		RR 8.25 (0.46 to 147.45)	67 (1 study)	⊕○○○ very low 2,4,5,8
	0 per 1000	0 per 1000 (0 to 0)			
	Moderate⁷				
	50 per 1000	412 per 1000 (23 to 1000)			
	High⁷				
	100 per 1000	825 per 1000 (46 to 1000)			
Adverse effects: Specific - hypertonia/rigidity (only reported if occurred in ≥9%) during 24 hours	Low¹		RR 2.74 (0.30 to 25.05)	67 (1 study)	⊕○○○ very low 2,4,5,8
	10 per 1000	27 per 1000 (3 to 250)			
	Moderate¹				
	30 per 1000	82 per 1000 (9 to 751)			
	High¹				
	50 per 1000	137 per 1000 (15 to 1000)			
Economic outcome - not reported	See comment	See comment	Not estimable	-	See comment No study reported this outcome.

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Moderate risk is roughly equal to that of the control group.

² Risk of bias: rated 'serious' - allocation concealment not stated, 2 people excluded from the efficacy analysis following randomisation, adverse effects only reported if they occurred in ≥9% of participants, sponsored by drug company.

³ Indirectness: rated 'serious' - there are no data for 30 minutes, only 3 hours.

⁴ Imprecision: rated 'serious' - relatively small number of participants in each treatment arm.

⁵ Publication bias: rated 'strongly suspected' - sponsored by drug company.

⁶ Risk of bias: rated 'very serious' - method of randomisation not described, allocation concealment not described, open trial, no reporting regarding the presence or absence of adverse effects, duration of the study is not reported.

⁷ Low risk is roughly equal to that of the control group.

⁸ Indirectness: rated 'severe' - not reported as 'severe' in this study.

Summary of findings 18. HALOPERIDOL compared to COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM for psychosis-induced aggression or agitation

HALOPERIDOL compared to COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM for psychosis-induced aggression or agitation

Patient or population: psychosis-induced aggression or agitation

Setting: emergency room.

Intervention: HALOPERIDOL

Comparison: COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM

Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	N° of participants (studies)	Quality of the evidence (GRADE)	Comments
	Risk with COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM	Risk with HALOPERIDOL				
Tranquillisation or asleep - not reported	-	-	-	-	-	No study reported this outcome.
Repeated need for tranquillisation assessed with: need for additional drugs for tranquillisation by 12 hours	Study population		RR 0.88 (0.69 to 1.13)	60 (1 RCT)	⊕⊕⊕⊕ very low ^{1 2}	
	867 per 1.000	763 per 1.000 (598 to 979)				
Specific behaviour - agitation	MD 11.20 lower		-	60 (1 RCT)	⊕⊕⊕⊕ very low ^{1 2}	

assessed with: average endpoint scores at 12 hours on OASS	(12.24 lower to 10.16 lower)				
Global outcome assessed with: additional restraint, seclusion or special observation during 12 hours	Low		RR 0.29 (0.13 to 0.61)	60 (1 RCT)	⊕⊕⊕⊕ very low ^{1 2}
	350 per 1.000	102 per 1.000 (46 to 214)			
	Moderate				
	700 per 1.000	203 per 1.000 (91 to 427)			
	High				
	950 per 1.000	275 per 1.000 (124 to 580)			
Adverse effects assessed with: EPS by 12 hours	Study population		RR 1.67 (0.44 to 6.36)	60 (1 RCT)	⊕⊕⊕⊕ very low ^{1 2}
	100 per 1.000	167 per 1.000 (44 to 636)			
Economic outcome - not reported	-	-	-	-	- No study reported this outcome.

***The risk in the intervention group** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio

GRADE Working Group grades of evidence

High quality: We are very confident that the true effect lies close to that of the estimate of the effect

Moderate quality: We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low quality: Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect

Very low quality: We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect

¹ Risk of bias: rated 'very serious' - method of allocation not stated, patients instructed to not reveal their allocation although the trial is described as double blind, selective reporting, small study.

² Imprecision: rated 'very serious' - small study, 95% CI cross the no effect line.

Summary of findings 19. HALOPERIDOL compared to COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE for psychosis-induced aggression or agitation

HALOPERIDOL compared to COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE for psychosis-induced aggression or agitation

Patient or population: psychosis-induced aggression or agitation

Setting: emergency room.

Intervention: HALOPERIDOL

Comparison: COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE

Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	N° of participants (studies)	Quality of the evidence (GRADE)	Comments
	Risk with COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE	Risk with HALOPERIDOL				
Tranquillisation or asleep assessed with: Not tranquil or asleep at 20 minutes	Study population		RR 1.60 (1.18 to 2.16)	316 (1 RCT)	⊕⊕⊕⊖ moderate ¹	
	281 per 1.000	450 per 1.000 (332 to 608)				
Repeated need for tranquillisation assessed with: need for additional drugs for tranquillisation by 2 hours	Low		RR 0.78 (0.43 to 1.41)	376 (2 RCTs)	⊕⊕⊖⊖ very low ^{1 3 4}	
	50 per 1.000	39 per 1.000 (22 to 71)				
	Moderate					
	120 per 1.000	94 per 1.000 (52 to 169)				
	High					
	450 per 1.000	351 per 1.000 (194 to 635)				
Specific behaviours - aggression assessed with: Further aggressive episode within 24 hours	Study population		RR 1.06 (0.68 to 1.65)	316 (1 RCT)	⊕⊕⊕⊖ moderate ¹	
	194 per 1.000	205 per 1.000 (132 to 320)				
Global outcomes assessed with: restraints needed by 120 minutes	Study population		RR 1.21 (0.84 to 1.76)	316 (1 RCT)	⊕⊕⊕⊖ moderate ^{1 5}	

	238 per 1.000	287 per 1.000 (199 to 418)			
Adverse effects: specific - acute dys- tonia	Low		RR 19.48 (1.14 to 331.92)	316 (1 RCT)	⊕⊕⊕⊕ low ^{1 6}
	0 per 1.000	0 per 1.000 (0 to 0)			
	Moderate				
	50 per 1.000	974 per 1.000 (57 to 1.000)			
	High				
	100 per 1.000	1000 per 1.000 (114 to 1.000)			
Adverse effects: specific - seizure	Study population		RR 2.05 (0.19 to 22.39)	316 (1 RCT)	⊕⊕⊕⊕ low ^{1 6}
	6 per 1.000	13 per 1.000 (1 to 140)			
Economic outcome - not reported	-	-	-	-	- No trial report- ed this out- come.

***The risk in the intervention group** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio

GRADE Working Group grades of evidence

High quality: We are very confident that the true effect lies close to that of the estimate of the effect

Moderate quality: We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low quality: Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect

Very low quality: We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect

¹ Risk of bias: rated 'serious' - open trial.

² Moderate risk is roughly equal to that of the trial control group.

³ Inconsistency: rated 'very serious' - I² 86%.

⁴ Publication bias: rated 'strongly suspected' - visual inspection of the funnel plot.

- ⁵ Indirectness: overall improvement not reported, therefore global improvement inferred from need for restraints at by 120 minutes.
⁶ Imprecision: rated 'serious' - 95% confidence intervals are wide.
⁷ Low risk is roughly equal to that of the trial control group.

Summary of findings 20. HALOPERIDOL compared to COMBINATIONS: e. HALOPERIDOL + RISPERIDONE for psychosis-induced aggression or agitation

HALOPERIDOL compared to COMBINATIONS: e. HALOPERIDOL + RISPERIDONE for psychosis-induced aggression or agitation

Patient or population: psychosis-induced aggression or agitation

Setting: inpatients.

Intervention: HALOPERIDOL

Comparison: COMBINATIONS: e. HALOPERIDOL + RISPERIDONE

Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	Nº of participants (studies)	Quality of the evidence (GRADE)	Comments
	Risk with COMBINATIONS: e. HALOPERIDOL + RISPERIDONE	Risk with HALOPERIDOL				
Tranquillisation or asleep - not reported	-	-	-	-	-	No study reported this outcome.
Repeated need for tranquillisation - not reported	-	-	-	-	-	No study reported this outcome.
Specific behaviour - agitation assessed with: average endpoint scores at 24 hours on PANSS-EC		MD 3.82 higher (1.35 higher to 6.29 higher)	-	100 (1 RCT)	⊕⊕⊕⊕ low ^{1 2}	
Global outcome assessed with: no improvement at 72 hours	Study population		RR 0.38 (0.11 to 1.33)	100 (1 RCT)	⊕⊕⊕⊕ low ^{1 2}	
	160 per 1.000	61 per 1.000 (18 to 213)				
Adverse effects: specific assessed with akathisia during 14 days	Study population		RR 0.88 (0.48 to 1.60)	100 (1 RCT)	⊕⊕⊕⊕ low ^{1 2}	
	320 per 1.000	282 per 1.000 (154 to 512)				

Economic outcome - not reported

-

-

-

-

-

No study reported this outcome.

***The risk in the intervention group** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio

GRADE Working Group grades of evidence

High quality: We are very confident that the true effect lies close to that of the estimate of the effect

Moderate quality: We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low quality: Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect

Very low quality: We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect

¹ Risk of bias: rated 'serious' - allocation concealment procedures not stated, blinding procedures not stated.

² Imprecision: rated 'serious' - 95% CIs are wide.

Summary of findings 21. HALOPERIDOL compared to COMBINATIONS: f. OLANZAPINE + MAGNESIUM VALPROATE for psychosis-induced aggression or agitation

HALOPERIDOL compared to COMBINATIONS: f. OLANZAPINE + MAGNESIUM VALPROATE for psychosis-induced aggression or agitation

Patient or population: psychosis-induced aggression or agitation

Setting: inpatients.

Intervention: HALOPERIDOL

Comparison: COMBINATIONS: f. OLANZAPINE + MAGNESIUM VALPROATE

Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	Nº of participants (studies)	Quality of the evidence (GRADE)	Comments
	Risk with COMBINATIONS: f. OLANZAPINE + MAGNESIUM VALPROATE	Risk with HALOPERIDOL				
Tranquillisation or asleep - not reported	-	-	-	-	-	No study reported this outcome.

Repeated need for tranquillisation - not reported	-	-	-	-	No study reported this outcome.
Specific behaviour - agitation assessed with: average endpoint scores at 72 hours on PANSS-EC	MD 0.04 lower (1.33 lower to 1.25 higher)		-	64 (1 RCT)	⊕⊕⊕⊕ very low ^{1 2}
Global outcome assessed with: no improvement	Study population		RR 1.06 (0.38 to 2.95)	64 (1 RCT)	⊕⊕⊕⊕ very low ^{1 2}
	182 per 1.000	193 per 1.000 (69 to 536)			
Adverse effects assessed with: one or more drug-related adverse events	Low		RR 3.06 (1.62 to 5.79)	64 (1 RCT)	⊕⊕⊕⊕ very low ^{1 2}
	100 per 1.000	306 per 1.000 (162 to 579)			
	Moderate				
	240 per 1.000	734 per 1.000 (389 to 1.000)			
	High				
	500 per 1.000	1000 per 1.000 (810 to 1.000)			
Economic outcome - not reported	-	-	-	-	No study reported this outcome.

***The risk in the intervention group** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio

GRADE Working Group grades of evidence

High quality: We are very confident that the true effect lies close to that of the estimate of the effect

Moderate quality: We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low quality: Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect

Very low quality: We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect

- ¹ Risk of bias: rated 'serious' - allocation concealment procedures not stated, blinding procedures not stated, small study.
² Imprecision: rated 'very serious' - small study, 95% CI cross the no effect area.

Summary of findings 22. HALOPERIDOL compared to COMBINATIONS: g. QUETIAPINE + MAGNESIUM VALPROATE for psychosis-induced aggression or agitation

HALOPERIDOL compared with COMBINATIONS: d. QUETIAPINE + MAGNESIUM VALPROATE for psychosis-induced aggression or agitation

Patient or population: patients with psychosis-induced aggression or agitation

Setting: inpatients.

Intervention: HALOPERIDOL

Comparison: COMBINATIONS: d. QUETIAPINE + MAGNESIUM VALPROATE

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	COMBINATIONS: d. QUETIAPINE + MAGNESIUM VALPROATE	HALOPERIDOL				
Tranquillisation or asleep - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Repeated need for tranquillisation - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Specific behaviours - agitation at 3 days average endpoint score (PANSS-EC) Follow-up: 8 weeks		The mean specific behaviours - agitation at 3 days in the intervention groups was 0.02 higher (2.31 lower to 2.35 higher)		60 (1 study)	⊕⊕⊕⊕ very low ^{1,2}	
Global effect Specific behaviour - agitation - no effect based on PANSS-EC (< 25% decrease)	Low ³		RR 1.17 (0.44 to 3.06)	60 (1 study)	⊕⊕⊕⊕ very low ^{1,4}	
	50 per 1000	58 per 1000 (22 to 153)				
	Moderate ³					
	200 per 1000	234 per 1000				

	(88 to 612)					
	High³					
	350 per 1000	409 per 1000 (154 to 1000)				
Adverse effects - any specific, serious - not reported	See comment	See comment	Not estimable	-	See comment	No study reported this outcome.
Economic outcome - not reported	See comment	See comment	Not estimable	0 (0)	See comment	No study reported this outcome.

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Risk of bias: rated 'very serious' - method of randomisation is not reported, allocation concealment not stated, open-label, selective reporting.

² Indirectness: rated 'very serious' - no data for aggressive outburst with 24 hours, therefore had to use scale-derived data for agitation at 72 hours.

³ Moderate risk is roughly equal to that of the trial control group.

⁴ Indirectness: rated 'serious' - no measure for overall improvement, therefore inferred no overall improvement from "no effect" based on the PANSS-EC.

Summary of findings 23. HALOPERIDOL compared to COMBINATIONS: h. RISPERIDONE + CLONAZEPAM for psychosis induced aggression or agitation

HALOPERIDOL compared to COMBINATIONS: h. RISPERIDONE + CLONAZEPAM for psychosis induced aggression or agitation

Patient or population: psychosis induced aggression or agitation

Setting: inpatients.

Intervention: HALOPERIDOL

Comparison: COMBINATIONS: h. RISPERIDONE + CLONAZEPAM

Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	Nº of participants (studies)	Quality of the evidence (GRADE)	Comments
	Risk with COMBINATIONS: h.	Risk with HALOPERIDOL

	RISPERIDONE + CLONAZEPAM				
Tranquillisation or asleep assessed with: not asleep at 4 hours	Study population		RR 0.82 (0.66 to 1.03)	205 (1 RCT)	⊕⊕⊕⊕ very low ¹²
	673 per 1.000	552 per 1.000 (444 to 693)			
Repeated need for tranquillisation - not reported	-	-	-	-	- No study re-ported this out-come.
Specific behaviour - agitation assessed with: average change scores at 2 hours on PANSS-EC	MD 0.50 lower (3.8 lower to 2.8 higher)		-	108 (1 RCT)	⊕⊕⊕⊕ very low ¹²
Global outcome assessed with: no improvement at 72 hours	Study population		RR 0.60 (0.15 to 2.38)	100 (1 RCT)	⊕⊕⊕⊕ low ^{3 4}
	100 per 1.000	60 per 1.000 (15 to 238)			
Adverse effects assessed with: akathisia during 5 days (only reported if occurred ≥50%)	Low		RR 1.63 (1.12 to 2.38)	205 (1 RCT)	⊕⊕⊕⊕ very low ¹⁵
	150 per 1.000	244 per 1.000 (168 to 357)			
	Moderate				
	300 per 1.000	489 per 1.000 (336 to 714)			
	High				
	500 per 1.000	815 per 1.000 (560 to 1.000)			
Economic outcome - not reported	-	-	-	-	- No study re-ported this out-come.
<p>*The risk in the intervention group (and its 95% confidence interval) is based on the assumed risk in the comparison group and the relative effect of the intervention (and its 95% CI).</p> <p>CI: Confidence interval; RR: Risk ratio</p>					

GRADE Working Group grades of evidence

High quality: We are very confident that the true effect lies close to that of the estimate of the effect

Moderate quality: We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low quality: Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect

Very low quality: We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect

- ¹ Risk of bias: rated 'very serious' - open study, evidence of reporting bias, sponsored trial.
- ² Imprecision: rated 'serious' - 95% CI cross the no effect area.
- ³ Risk of bias: rated 'serious' - allocation concealment and blinding procedures not stated.
- ⁴ Imprecision: rated 'serious' - 95% CI are wide.
- ⁵ Imprecision: rated 'serious' - 95% CI are wide.

Summary of findings 24. HALOPERIDOL compared to COMBINATIONS: i. ZIPRASIDONE + CLONAZEPAM for psychosis-induced aggression or agitation

HALOPERIDOL compared to COMBINATIONS: i. ZIPRASIDONE + CLONAZEPAM for psychosis-induced aggression or agitation

Patient or population: psychosis-induced aggression or agitation

Setting: inpatients.

Intervention: HALOPERIDOL

Comparison: COMBINATIONS: i. ZIPRASIDONE + CLONAZEPAM

Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	Nº of participants (studies)	Quality of the evidence (GRADE)	Comments
	Risk with COM- BINATIONS: i. ZIPRASIDONE + CLONAZEPAM	Risk with HALOPERIDOL				
Tranquillisation or asleep - not reported	-	-	-	-	-	No study reported this outcome.
Repeated need for tranquillisation - not reported	-	-	-	-	-	No study reported this outcome.
Specific behaviour - agitation assessed with: average endpoint scores at 12 hours on PANSS-EC		MD 0.28 lower (1.80 lower to 1.24 higher)	-	71 (1 RCT)	⊕⊕⊕⊕ very low ^{1 2}	
Global outcome - no improvement - not reported	-	-	-	-	-	No study reported this outcome.

Adverse effects - not reported	-	-	-	-	-	No study reported this outcome.
Economic outcome - not reported	-	-	-	-	-	No study reported this outcome.

***The risk in the intervention group** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio

GRADE Working Group grades of evidence

High quality: We are very confident that the true effect lies close to that of the estimate of the effect

Moderate quality: We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low quality: Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect

Very low quality: We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect

¹ Risk of bias: rated 'very serious' - randomisation, allocation and blinding procedures not stated, evidence of attrition bias, small study.

² Imprecision: rated 'very serious' - small study, 95% CI are wide.

BACKGROUND

Description of the condition

Psychosis is associated with a number of mental disorders, including schizophrenia and bipolar disorder. Symptoms of psychosis include delusions and hallucinations, which can lead some people to become confused, frightened or agitated. Often they can suffer a combination of these distressing emotions (APA 2006). Agitation is characterised by restlessness, excitability and irritability, and for some people, this can result in verbal and physical aggressive behaviour (Mohr 2005). Agitation and aggression within the psychiatric setting imposes a significant challenge to clinicians who, while attempting to make an accurate diagnosis and formulation (Schleifer 2011), have to intervene quickly in order to manage the risk that the service user may present to themselves, other service users and staff (NICE 2005, NICE 2015).

Description of the intervention

Haloperidol was developed in 1958 by the Belgian company Janssen Pharmaceutica, and with the earlier development of chlorpromazine was considered a "psychopharmacological revolution" for the treatment of schizophrenia (López-Munoz 2009). Newer anti-psychotic medication has been developed for the day-to-day management of symptoms of schizophrenia, however, haloperidol continues to be in wide use, particularly for the management of psychosis-induced agitation. In clinical practice, where de-escalation techniques have not been appropriate nor sufficient in managing a person's agitation and where that person poses a threat to themselves or others, it may be necessary to use medication for rapid tranquillisation (Pratt 2008). Although there may be occasions where it is appropriate to induce sleep, the aim of rapid tranquillisation should usually be to sedate agitated or aggressive people to a point where they are still able to engage in further assessment and be responsive to communication (Parker 2010). If clinically feasible, oral administration is preferred. However, if the person refuses medication or there is no improvement after 30 to 60 minutes, haloperidol in single doses of 5 mg, administered intramuscularly (IM) is a commonly used treatment option (Pereira 2007).

Cardiac problems and extrapyramidal side effects (EPS) such as dystonia and akathisia are associated with conventional antipsychotics such as haloperidol. Experiencing distressing adverse effects may act as a barrier for future engagement with services or treatment (Pratt 2008). The UK's NICE guidelines recommend that when using haloperidol IM, an antimuscarinic agent such as procyclidine or benztropine should be administered to counter the potential for EPS. Subsequent to the publication of these guidelines, the manufacturers of Haldol (Janssen-Cilag Ltd 2010) recommended that a baseline electrocardiogram (ECG) should be undertaken prior to treatment with haloperidol. In addition, they have discontinued the license for intravenous (IV) use and reduced the recommended maximum dose from 90 mg/day to 30 mg/day orally, and 18 mg/day IM (Parker 2010). When Haldol is prescribed outside of these recommendations, this would be considered 'off label' and may increase the prescriber's professional responsibility and potential liability (Pratt 2008). Despite these adverse effects, haloperidol alone or combined with a benzodiazepine was recommended by the NICE guidelines (NICE 2005) and by the American Psychiatric Association (APA 2006), and

haloperidol remains on the World Health Organization's Essential Medicine list (WHO 2011, WHO 2015). With the last updated NICE guidelines, recommendation changed indicating either the use of haloperidol combined with promethazine or lorazepam alone for rapid tranquillisation (NICE 2015).

How the intervention might work

Haloperidol is mainly indicated for schizophrenia or other psychoses, mania, violent or dangerously impulsive behaviour, excitement and for short-term adjunctive management of psychomotor agitation (BNF 2011). Haloperidol is from the butyrophenone family of antipsychotics (neuroleptics) (López-Munoz 2009). It is thought that haloperidol prevents the occurrence of delusions and hallucinations by blocking the dopamine D2 receptors in the meso-cortico-limbic system. In one study, while the D2 dopamine receptor binding was high in the temporal cortex for both haloperidol and atypical antipsychotics, haloperidol induced a significantly higher binding index in the thalamus and striatum than atypical antipsychotics. It is thought that this antidopaminergic activity in the dorsolateral striatum may contribute to the extra pyramidal effects that are associated with typical antipsychotics such as haloperidol (Xiberas 2001).

Why it is important to do this review

The UK's 2005 NICE guidelines recommended that where haloperidol is administered via injection it should be combined with lorazepam when given IM. However, these guidelines also recommended, that, in exceptional circumstances, haloperidol can be given alone by IV injection (NICE 2005). Despite this recommendation, it is difficult to administer IV injections safely in restraint situations and there is evidence that haloperidol alone continues to be given via IM injection (Choudhury 2011). Surveys undertaken with consultant psychiatrists have shown that following the withdrawal of droperidol, haloperidol was the preferred alternative for rapid tranquillisation (Reid 2003) and that over 33% of respondents would be prepared to use doses above the BNF limits (Pereira 2007). Further, since the NICE guideline on the management of disturbed and violent behaviour was published, Janssen-Cilag Ltd has withdrawn the license for IV use. With the increased prescribing of haloperidol in the UK by 16% (excluding depot injections) between 2004 and 2009 (NHS 2009), the authors felt it timely and justified to examine the evidence regarding the efficacy and safety of prescribing haloperidol alone for treating psychosis-induced agitation and aggression. This is one of a series of linked completed and maintained reviews and other reviews that are underway (Table 1) that will create a body of evidence assessing the effectiveness of various drugs and their preferred routes of administration for both short- and long-term psychosis-induced aggression or agitation.

In the UK's 2015 NICE guidelines, either use of haloperidol combined with promethazine or lorazepam alone is now recommended, with the latter preferred when in presence of cardiovascular disease or in case a recent ECG is unavailable (NICE 2015).

OBJECTIVES

To examine whether haloperidol alone, administered orally, intramuscularly or intravenously, is an effective treatment for psychosis-induced aggression or agitation, wherein clinicians are required to intervene to prevent harm to self and others.

METHODS

Criteria for considering studies for this review

Types of studies

All relevant randomised controlled trials. If a trial was described as 'double-blind' but implied randomisation, we included such trials in a sensitivity analysis (see [Sensitivity analysis](#)). If their inclusion did not result in a substantive difference, they remained in the analyses. If their inclusion resulted in statistically significant differences, we did not add the data from these lower-quality studies to the results of the better trials, but presented such data within a subcategory. We excluded quasi-randomised studies, such as those allocating by alternate days of the week.

Types of participants

People exhibiting agitation or aggression (or both) thought to be due to psychotic illness, regardless of age or sex. We included studies that involved participants with other diagnoses, such as drug or alcohol intoxication, organic problems including dementia, learning disability and non-psychotic mental illness, providing that the number of participants in these groups did not exceed the number of participants with psychosis.

Types of interventions

1. Rapid use of haloperidol

Alone, any dose via any route of administration; compared with rapid use of the following.

a. Placebo or no intervention

We are aware that, in a situation where haloperidol is available, the option of not giving treatment would be problematic. These are, however, valuable and pioneering studies which give us some insight to the absolute value of haloperidol in the emergency situation.

b. Other antipsychotic

Any dose via any route of administration.

c. Benzodiazepine

Any dose via any route of administration.

d. Anticonvulsant alone

Any dose via any route of administration.

e. Combination of drugs

Any dose via any route of administration.

Rapid use of the interventions is defined as administration deemed necessary to calm behaviour or prevent harm to the participant or others.

Types of outcome measures

All outcomes grouped by time: by 30 minutes, up to two hours, up to four hours, up to 24 hours and over 24 hours.

Primary outcomes

1. Tranquilisation or asleep

- 1.1 Not tranquil or asleep - by up to 30 minutes
- 1.2 Repeated need for rapid tranquillisation

Secondary outcomes

1. Tranquillisation or asleep

- 1.1 Not tranquil - over 30 minutes
- 1.2 Not asleep - over 30 minutes
- 1.3 Time to tranquillisation/sleep
- 1.4 Time to tranquillisation
- 1.5 Time to sleep

2. Specific behaviours

- 2.1 Self-harm, including suicide
- 2.2 Injury to others
- 2.3 Agitation
 - 2.3.1 Another episode of agitation by 24 hours
 - 2.3.2 No clinically important change in agitation
 - 2.3.3 No change in agitation
 - 2.3.4 Average endpoint in agitation score
 - 2.3.5 Average change in agitation scores
- 2.4 Aggression
 - 2.4.1 Another episode of aggression by 24 hours
 - 2.4.2 No clinically important change in aggression
 - 2.4.3 No change in aggression
 - 2.4.4 Average endpoint in aggression score
 - 2.4.5 Average change in aggression scores

3. Global outcomes

- 3.1 No overall improvement
- 3.2 Use of restraints/seclusion
- 3.3 Relapse - as defined by each study
- 3.4 Recurrence of violent incidents
- 3.5 Needing extra visits from the doctor
- 3.6 Refusing oral medication
- 3.7 Not accepting treatment
- 3.8 Average endpoint score
- 3.9 Average change score
- 3.10 Average dose of drug

4. Service outcomes

- 4.1 Duration of hospital stay
- 4.2 Re-admission
- 4.3 No clinically important engagement with services
- 4.4 No engagement with services
- 4.5 Average endpoint engagement score
- 4.6 Average change in engagement scores

5. Mental state

- 5.1 No clinically important change in general mental state
- 5.2 No change in general mental state
- 5.3 Average endpoint general mental state score
- 5.4 Average change in engagement scores

6. Adverse effects

- 6.1 Death
- 6.2 Any non-serious general adverse effects
- 6.3 Any serious, specific adverse effects

- 6.4 Average endpoint general adverse effect score
- 6.5 Average change in general adverse effect scores
- 6.6 No clinically important change in specific adverse effects
- 6.7 No change in specific adverse effects
- 6.8 Average endpoint specific adverse effects
- 6.9 Average change in specific adverse effects

7. Leaving the study early

- 7.1 For specific reasons
- 7.2 For general reasons

8. Satisfaction with treatment

- 8.1 Recipient of treatment not satisfied with treatment
- 8.2 Recipient of treatment average satisfaction score
- 8.3 Recipient of treatment average change in satisfaction scores
- 8.4 Informal treatment provider not satisfied with treatment
- 8.5 Informal treatment providers' average satisfaction scores
- 8.6 Informal treatment providers' average change in satisfaction scores
- 8.7 Professional providers' not satisfied with treatment
- 8.8 Professional providers' average satisfaction score
- 8.9 Professional providers' average change in satisfaction scores

9. Acceptance of treatment

- 9.1 Not accepting treatment
- 9.2 Average endpoint acceptance score
- 9.3 Average change in acceptance scores

10. Quality of life

- 10.1 No clinically important change in quality of life
- 10.2 Not any change in quality of life
- 10.3 Average endpoint quality of life score
- 10.4 Average change in quality of life score
- 10.5 No clinically important change in specific aspects of quality of life
- 10.6 No change in specific aspects of quality of life
- 10.7 Average endpoint specific aspects of quality of life
- 10.8 Average change in specific aspects of quality of life

11. Economic outcomes

- 11.1 Direct costs
- 11.2 Indirect costs

Outcomes used for 'Summary of findings' table

We used the GRADE approach to interpret findings (Schünemann 2008) and used the GRADE profiler (GRADE PRO) to import data from RevMan 5 (Review Manager) to create 'Summary of findings' tables. These tables provide outcome-specific information concerning the overall quality of evidence from each included study in the comparison, the magnitude of effect of the interventions examined, and the sum of available data on all outcomes that we rated as important to patient-care and decision making. We selected the following main outcomes for inclusion in the 'Summary of findings' tables.

- 1. Tranquillisation or asleep - not tranquil or asleep by 30 minutes.
- 2. Tranquillisation or asleep - repeated need for rapid tranquillisation - within 24 hours.
- 3. Specific behaviours - threat or injury to others/self.
- 4. Global outcomes - overall improvement.

- 5. Adverse effects - any serious, specific adverse effects.
- 6. Economic outcomes.

Search methods for identification of studies

Electronic searches

Cochrane Schizophrenia Group's Study-Based Register of Trials

On May 26, 2016, the information specialist searched the register using the following search strategy which has been developed based on literature review and consulting with the authors of the review:

(*Agitation* OR *Aggression*) in Healthcare Condition AND (*Haloperidol*) in Intervention of STUDY

In such study-based register, searching the major concept retrieves all the synonym and relevant studies because all the studies have already been organised based on their interventions and linked to the relevant topics.

This register is compiled by systematic searches of major resources (including AMED, BIOSIS CINAHL, Embase, MEDLINE, PsycINFO, PubMed, and registries of clinical trials) and their monthly updates, handsearches, grey literature, and conference proceedings (see [Group's Module](#)). There is no language, date, document type, or publication status limitations for inclusion of records into the register.

For previous searches, please see [Appendix 1](#).

Searching other resources

1. Reference searching

We inspected references of all included studies for further relevant studies.

2. Personal contact

We attempted to contact the first author of each included study for information regarding unpublished trials.

Data collection and analysis

Selection of studies

Review author MJP inspected citations from the searches and identified relevant abstracts. A random 20% sample was independently re-inspected by CEA to ensure reliability. Where disputes arose, we acquired the full report for more detailed scrutiny. We obtained full reports of the abstracts that met the review criteria and these were inspected by MJP. Again, a random 20% of reports were re-inspected by CEA, in order to ensure reliable selection. Where it was not possible to resolve disagreement by discussion, we attempted to contact the authors of the study for clarification.

Review authors EGO and XL screened results from 2016 search to identify relevant abstracts. We obtained and inspected full reports of the abstracts meeting the review criteria.

Data extraction and management

1. Extraction

Review author MJP (original search), EGO (2016 search) extracted data from all included studies. In addition, to ensure reliability, CEA independently extracted data from a random sample of these studies, comprising 10% of the total. Again, any disagreement was discussed, decisions documented and, if necessary, we contacted the authors of studies for clarification. If there were any remaining problems, HJ helped clarify issues and these final decisions were documented. We extracted data presented only in graphs and figures whenever possible, but we only included the data if two review authors independently had the same result. We attempted to contact authors through an open-ended request in order to obtain missing information or for clarification whenever necessary. If studies were multi-centre, where possible, we extracted data relevant to each component centre separately.

2. Management

2.1 Forms

We extracted data onto standard, simple forms.

2.2 Scale-derived data

We included continuous data from rating scales only if:

- the psychometric properties of the measuring instrument have been described in a peer-reviewed journal ([Marshall 2000](#)); and
- the measuring instrument has not been written or modified by one of the trialists for that particular trial.

Ideally, the measuring instrument should either be i. a self-report or ii. completed by an independent rater or relative (not the therapist). We realise that this is not often reported clearly, therefore, in [Description of studies](#), we recorded whether or not this was the case.

2.3 Endpoint versus change data

There are advantages of both endpoint and change data. Change data can remove a component of between-person variability from the analysis. On the other hand, calculation of change needs two assessments (baseline and endpoint), which can be difficult in unstable and difficult to measure conditions such as schizophrenia. We decided primarily to use endpoint data, and only use change data if the former were not available. We combined endpoint and change data in the analysis and we used mean differences (MD) rather than standardised mean differences (SMD) throughout ([Higgins 2011](#)).

2.4 Skewed data

Continuous data on clinical and social outcomes are often not normally distributed. To avoid the pitfall of applying parametric tests to non-parametric data, we aimed to apply the following standards to all data before inclusion: a) standard deviations (SDs) and means are reported in the paper or obtainable from the authors; b) when a scale starts from the finite number zero, the SD, when multiplied by two, is less than the mean (as otherwise the mean is unlikely to be an appropriate measure of the centre of the distribution, ([Altman 1996](#)); c) if a scale started from a positive value (such as the Positive and Negative Syndrome Scale (PANSS), ([Kay 1986](#))), which can have values from 30 to 210), the calculation described above was modified to take the scale starting point into

account. In these cases skew is present if $2\text{SD} > (\text{S} - \text{S min})$, where S is the mean score and 'S min' is the minimum score. Endpoint scores on scales often have a finite start and end point and these rules can be applied. Skewed data pose less of a problem when looking at means if the sample size is large (> 200) and we entered these into the syntheses. We present skewed data from studies of less than 200 participants in 'Additional tables' rather than in analyses.

When continuous data are presented on a scale that includes a possibility of negative values (such as change data), it is difficult to tell whether data are skewed or not. These were entered into analyses.

2.5 Common measure

To facilitate comparison between trials, if required, we converted variables that can be reported in different metrics, such as time to tranquillisation (e.g. minutes, hours) to a common metric (e.g. minutes).

2.6 Conversion of continuous to binary

Where possible, efforts were made to convert outcome measures to dichotomous data. This was done by identifying cut-off points on rating scales and dividing participants accordingly into 'clinically improved' or 'not clinically improved'. It is generally assumed that if there is a 50% reduction in a scale-derived score such as the Brief Psychiatric Rating Scale (BPRS, [Overall 1962](#)) or the PANSS scale ([Kay 1986](#)), this could be considered as a clinically significant response ([Leucht 2005](#); [Leucht 2005a](#)). If data based on these thresholds were not available, we used the primary cut-off presented by the original authors.

2.7 Direction of graphs

Where possible, we entered data in such a way that the area to the left of the line of no effect indicated a favourable outcome for haloperidol for psychosis-induced agitation. Where keeping to this made it impossible to avoid outcome titles with clumsy double-negatives (e.g. 'Not improved'), we reported data where the left of the line indicates an unfavourable outcome. This was noted in the relevant graphs.

Assessment of risk of bias in included studies

Review authors MJP and EGO assessed risk of bias by using the criteria described in the *Cochrane Handbook for Systematic Reviews of Interventions* ([Higgins 2011](#)) to assess trial quality. This set of criteria is based on evidence of associations between overestimate of effect and high risk of bias of the article such as sequence generation, allocation concealment, blinding, incomplete outcome data and selective reporting. HJ re-rated 10% of included studies.

If the raters disagreed, or if either rater was in doubt, we decided the final rating by consensus, with the involvement of another member of the review group. Where inadequate details of randomisation and other characteristics of trials were provided, we contacted authors of the studies in order to obtain further information. We reported non-concurrence in quality assessment, but if disputes arose as to which category a trial was to be allocated, again, we resolved these by discussion.

We noted the level of risk of bias in both the text of the review and in the 'Summary of findings' tables.

Measures of treatment effect

1. Binary data

For binary outcomes, we calculated a standard estimation of the risk ratio (RR) and its 95% confidence interval (CI). It has been shown that RR is more intuitive (Boissel 1999) than odds ratios and that odds ratios tend to be interpreted as RR by clinicians (Deeks 2000).

2. Continuous data

For continuous outcomes, we estimated mean difference (MD) between groups. We preferred not to calculate effect size measures (SMD). However, if scales of very considerable similarity were used, we presumed there was a small difference in measurement, and we calculated the effect size and transformed the effect back to the units of one or more of the specific instruments.

Unit of analysis issues

1. Cluster trials

Studies increasingly employ 'cluster randomisation' (such as randomisation by clinician or practice) but analysis and pooling of clustered data poses problems. Firstly, authors often fail to account for intra-class correlation in clustered studies, leading to a 'unit of analysis' error (Divine 1992), whereby P values are spuriously low, CIs unduly narrow and statistical significance overestimated. This causes type I errors (Bland 1997; Gulliford 1999).

Where clustering was not accounted for in primary studies, we presented data in a table, with a (*) symbol to indicate the presence of a probable unit of analysis error. We attempted to contact the first authors of studies to obtain intra-class correlation coefficients (ICCs) for their clustered data and to adjust for this by using accepted methods (Gulliford 1999). Where clustering had been incorporated into the analysis of primary studies, we presented these data as if from a non-cluster randomised study, but adjusted for the clustering effect.

We sought statistical advice and have been advised that the binary data as presented in a report should be divided by a 'design effect'. This is calculated using the mean number of participants per cluster (m) and the ICC [Design effect = $1 + (m - 1) \times \text{ICC}$] (Donner 2002). If the ICC was not reported it was assumed to be 0.1 (Ukoumunne 1999).

Where cluster studies had been appropriately analysed taking into account ICCs and relevant data documented in the report, synthesis with other studies was possible using the generic inverse variance technique.

2. Cross-over trials

A major concern of cross-over trials is the carry-over effect. It occurs if an effect (e.g. pharmacological, physiological or psychological) of the treatment in the first phase is carried over to the second phase. As a consequence, on entry to the second phase the participants can differ systematically from their initial state despite a wash-out phase. For the same reason cross-over trials are not appropriate if the condition of interest is unstable (Elbourne 2002). As both effects are very likely in severe mental illness, if we had included cross-over trials, we would only have used data from the first phase of cross-over studies.

3. Studies with multiple treatment groups

Where a study involved more than two treatment arms, where relevant, the additional treatment arms were presented in comparisons. If data were binary these were simply added and combined within the two-by-two table. If data were continuous, we combined data following the formula in section 7.7.3.8 (Combining groups) of the *Cochrane Handbook for Systematic Reviews of Interventions*. Where the additional treatment arms were not relevant, we did not reproduce these data.

Dealing with missing data

1. Overall loss of credibility

At some degree of loss of follow-up, data must lose credibility (Xia 2009). We chose that, for any particular outcome, should more than 50% of data be unaccounted for, we did not reproduce these data or use them within analyses, (except for the outcome 'leaving the study early'). Where more than 50% of those in one arm of a study were lost, but the total loss was less than 50%, we marked such data with (*) to indicate that such a result may well be prone to bias.

2. Binary

In the case where attrition for a binary outcome was between 0% and 50% and where these data were not clearly described, we presented data on a 'once-randomised-always-analyse' basis (an intention-to-treat (ITT) analysis). Those leaving the study early were all assumed to have the same rates of negative outcome as those who completed, with the exception of the outcome of death and adverse effects. For these outcomes, the rate of those who stayed in the study - in that particular arm of the trial - were used for those who did not. We undertook a sensitivity analysis to test how prone the primary outcomes were to change when 'completer' data only were compared with the ITT analysis using the above assumptions.

3. Continuous

3.1 Attrition

In the case where attrition for a continuous outcome was between 0% and 50% and completer-only data were reported, we reproduced these.

3.2 Standard deviations (SDs)

If SDs were not reported, we first tried to obtain the missing values from the authors. If not available, where there were missing measures of variance for continuous data, but an exact standard error (SE) and confidence intervals (CIs) were available for group means, and either a P value or 't' value were available for differences in mean, we calculated them according to the rules described in the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011): When only the SE was reported, SDs were calculated by the formula $\text{SD} = \text{SE} \times \text{square root } (n)$. Chapters 7.7.3 and 16.1.3 of the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011) present detailed formulae for estimating SDs from P values, t or F values, CIs, ranges or other statistics. If these formulae did not apply, we calculated the SDs according to a validated imputation method which is based on the SDs of the other included studies (Furukawa 2006). Although some of these imputation strategies can introduce error, the alternative would be to exclude a given study's outcome and

thus to lose information. Nevertheless, we examined the validity of the imputations in a sensitivity analysis excluding imputed values.

3.3 Last observation carried forward (LOCF)

We anticipated that in some studies the method of LOCF would be employed within the study report. As with all methods of imputation to deal with missing data, LOCF introduces uncertainty about the reliability of the results (Leucht 2007). Therefore, where LOCF data were used in the trial, if less than 50% of the data had been assumed, we reproduced these data and indicated that they were the product of LOCF assumptions.

Assessment of heterogeneity

1. Clinical heterogeneity

We considered all included studies initially, without seeing comparison data, to judge clinical heterogeneity. We inspected all studies for clearly outlying people or situations which we had not predicted would arise. When such situations or participant groups arose, these were discussed.

2. Methodological heterogeneity

We considered all included studies initially, without seeing comparison data, to judge methodological heterogeneity. We inspected all studies for clearly outlying methods which we had not predicted would arise. When such methodological outliers arose, these were discussed.

3. Statistical heterogeneity

3.1 Visual inspection

We visually inspected graphs to investigate the possibility of statistical heterogeneity.

3.2 Employing the I^2 statistic

We investigated heterogeneity between studies by considering the I^2 statistic alongside the χ^2 P value. The I^2 statistic provides an estimate of the percentage of inconsistency thought to be due to chance (Higgins 2003). The importance of the observed value of I^2 depends on i. magnitude and direction of effects and ii. strength of evidence for heterogeneity (e.g. P value from χ^2 test, or a CI for I^2). We interpreted an I^2 estimate greater than or equal to around 50% accompanied by a statistically significant χ^2 statistic, as evidence of substantial levels of heterogeneity (Section 9.5.2 - Higgins 2011). When substantial levels of heterogeneity were found in the primary outcome, we explored reasons for heterogeneity (Subgroup analysis and investigation of heterogeneity).

Assessment of reporting biases

1. Protocol versus full study

Reporting biases arise when the dissemination of research findings is influenced by the nature and direction of results. These are described in section 10.1 of the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011). We tried to locate protocols of included randomised trials. If the protocol was available, outcomes in the protocol and in the published report were compared. If the protocol was not available, outcomes listed in the methods section of the trial report were compared with actual reported results.

2. Funnel plot

Reporting biases arise when the dissemination of research findings is influenced by the nature and direction of results (Egger 1997). These are again described in Section 10 of the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011). We are aware that funnel plots may be useful in investigating reporting biases but are of limited power to detect small-study effects. We did not use funnel plots for outcomes where there were 10 or fewer studies, or where all studies were of similar sizes. In other cases, where funnel plots were possible, we sought statistical advice in their interpretation.

Data synthesis

We understand that there is no closed argument for preference for use of fixed-effect or random-effects models. The random-effects method incorporates an assumption that the different studies are estimating different, yet related, intervention effects. This often seems to be true to us and the random-effects model takes into account differences between studies even if there is no statistically significant heterogeneity. There is, however, a disadvantage to the random-effects model. It puts added weight onto small studies which often are the most biased ones. Depending on the direction of effect, these studies can either inflate or deflate the effect size. We chose the fixed-effect model for all analyses. The reader is, however, able to choose to inspect the data using the random-effects model.

Subgroup analysis and investigation of heterogeneity

1. Subgroup analyses

We did not anticipate any subgroup analyses.

2. Investigation of heterogeneity

If inconsistency was high, we reported it. First, we investigated whether data were entered correctly. Second, if data were correct, we visually inspected the graph and successively removed outlying studies to see if homogeneity was restored. For this review, we decided that should this occur with data contributing to the summary finding of no more than around 10% of the total weighting, we would present the data. If not, we would not pool data but would discuss the issues. We know of no supporting research for this 10% cut-off but are investigating the use of prediction intervals as an alternative to this unsatisfactory state.

When unanticipated clinical or methodological heterogeneity were obvious, we stated hypotheses regarding these for future reviews or versions of this review. We did not anticipate undertaking analyses relating to these.

Sensitivity analysis

1. Implication of randomisation

We aimed to include trials in a sensitivity analysis if they were described in some way as to imply randomisation. For the primary outcomes we included these studies, and if there was no substantive difference when the implied randomised studies were added to those with better description of randomisation, then we used all data from these studies.

2. Assumptions for lost binary data

Where assumptions had to be made regarding people lost to follow-up (see [Dealing with missing data](#)), we compared the findings of the primary outcomes when we used our assumption compared with 'completer' data only. Where there was a substantial difference, we reported results and discussed them, but continued to employ our assumption.

Where assumptions had to be made regarding missing SDs data (see [Dealing with missing data](#)), we compared the findings on primary outcomes when we used our assumption compared with 'completer' data only. We undertook a sensitivity analysis to test how prone results were to change when 'completer' data only were compared with the imputed data using the above assumption. If there was a substantial difference, we reported results and discussed them, but continued to employ our assumption.

3. Risk of bias

We analysed the effects of excluding trials that were judged to be at high risk of bias across one or more of the domains of randomisation (implied as randomised with no further details available) allocation concealment, blinding and outcome reporting for the meta-analysis of the primary outcome. If the exclusion of trials at high risk of bias did not substantially alter the direction of effect or the precision of the effect estimates, then we included data from these trials in the analysis.

4. Imputed values

We also undertook a sensitivity analysis to assess the effects of including data from trials where we used imputed values for ICC in calculating the design effect in cluster-randomised trials.

If substantial differences were noted in the direction or precision of effect estimates in any of the sensitivity analyses listed above,

we did not pool data from the excluded trials with the other trials contributing to the outcome, but presented them separately.

5. Fixed and random effects

All data was be synthesised using a fixed-effect model, however, we also synthesised data for the primary outcome using a random-effects model to evaluate whether this altered the significance of the result.

RESULTS

Description of studies

For substantive descriptions of studies please see the [Characteristics of included studies](#), [Characteristics of excluded studies](#), [Characteristics of studies awaiting classification](#) and [Characteristics of ongoing studies](#).

Results of the search

The electronic search (June 2011) yielded 665 citations of potentially eligible studies. After checking titles and abstracts, 209 full-text papers were obtained for a second assessment. These publications related to 100 different studies (29 included studies, 55 excluded, 15 awaiting classification and one ongoing). The reference lists of the 29 studies to be included were checked, through which we identified another three potential included studies. Again, full-text papers were obtained and on closer inspection, [Reschke 1974](#) and [Fitzgerald 1969](#) were included and [Goldstein 1966](#) excluded because, although participants, were experiencing psychosis, they were not specifically aggressive or agitated. The reference lists of related Cochrane systematic reviews were also searched, from which we identified [Bailine 1987](#) for inclusion (2011 search, [Figure 1](#)).

Figure 1. Study flow diagram.

The update search run in 2016 yielded 34 citations, of which nine new studies were included ([Figure 2](#)).

Figure 2. Study flow diagram.

Included studies

Details of the 41 studies included in the review are provided in the [Characteristics of included studies](#) table.

1. Length of studies

The duration of the studies ranged from two hours ([Calver 2015](#); [Dorevitch 1999](#)) to eight weeks ([Higashima 2004](#)). The majority were

≤ 72 hours, apart from [Garza-Trevino 1989](#) and [Nobay 2004](#) which did not report the duration of the study.

2. Participants

2.1 Clinical state

Participants in the studies presented with acute exacerbation of psychotic symptoms. Some studies focused on people whose psychosis had primarily triggered agitation ([Battaglia 2002](#); [Breier](#)

2002; Bristol Myers 2004a; Bristol-Myers 2005b; Brook 2000; Eli Lilly 2004; Fang 2012; Garza-Trevino 1989; Guo 2007; Higashima 2004; Kinon 2001d; Liang 2013; Lim 2010; Liu 2012; Resnick 1984; Shu 2010; Walther 2014; Yin 2012; Zhou 2014), and other trials included people with both agitation and aggression (Bailine 1987; Baldacara 2011; Battaglia 1997; Calver 2015; Currier 2004; Dorevitch 1999; Fitzgerald 1969; Foster 1997; Fruensgaard 1977; Huf 2007; Kewala 1984; Li 2006; Man 1973; Nobay 2004; Paprocki 1977; Reschke 1974; Ritter 1972; Salzman 1991; Shi 2010; Stotsky 1977; Taymeeyapradit 2002; Tuason 1986). All participants were so disturbed that clinicians felt that they required rapid tranquillisation.

2.2 Diagnosis

Over 80% of participants had a diagnosis of schizophrenia. Other diagnoses included schizoaffective disorder, schizophreniform disorder, bipolar disorder, and 'psychosis not otherwise specified'. A small minority of participants had a diagnosis of organic mental disorder and substance-induced psychosis, but in no trial was this subgroup the majority or the focus of the study.

2.3. Exclusions

Where exclusion criteria were reported, these often included pregnant or lactating women, people with serious medical illnesses, people who had used certain medications (e.g. antipsychotics/anti-depressants) within a specified period prior to enrolment, people with a known allergy/hypersensitivity to the study drugs and people with alcohol or drug dependency.

2.4 Age

In the 23 studies that reported an age range, participants ranged from 18 to 73 years. Nine studies reported a mean age (mostly around 30), with Garza-Trevino 1989 being the only study not to report the age of participants.

2.5 Sex

There were 2236 male participants and 1310 participants were women; the sex of 1387 people was not reported.

3. Study size

The study sizes varied with the smallest study having only 19 participants (Higashima 2004) and the largest randomising 448 people (Bristol Myers 2004a).

4. Setting

Where reported, the majority of participants presented at psychiatric emergency departments and were newly admitted inpatients. Interestingly, one study included outpatients (Stotsky 1977).

5. Interventions

It is unsurprising that this old and well-established drug has been compared with many other interventions. For all but one study (Kinon 2001d - see below), haloperidol and the other drugs were administered by the intramuscular (IM) route. What was surprising, however, was how poorly reported the true doses were. Dose ranges were commonly reported but to stipulate their mean or median was the exception rather than the rule. In several comparisons we have had to make educated guesses and the likely end doses.

5.1 Versus placebo

Trials comparing haloperidol with placebo largely used around 10 mg of the drug given by IM injection.

5.2 Versus other antipsychotics

The aripiprazole comparison involved around 10 mg of haloperidol compared with up to 45 mg of the control drug. The older trials involved in the chlorpromazine comparison used larger doses of haloperidol (10 mg to 20 mg with one trial allowing the option of up to 100 mg/day) with between 100 mg and up to 500 mg of chlorpromazine. In the droperidol trial 10 mg of haloperidol was compared with around 15 mg of droperidol. Although one loxapine versus haloperidol trial did allow for up to 100 mg of haloperidol, more commonly used doses across trials were in the 10 mg to 20 mg range compared with 50 mg to 250 mg of loxapine. The olanzapine dose tended to be around 10 mg to 15 mg, and in these trials haloperidol was given around the 10 mg dose. The Kinon 2001d trial (olanzapine versus haloperidol) is notable for it being the only one that used oral drugs rather than IM route of administration. The perphenazine comparison used doses of both drugs in the mid 20's mg given by IM injection. The quetiapine trial compared a maximum of 12 mg of haloperidol with up to 750 mg of quetiapine. In the risperidone comparison, haloperidol was used at between about 5 mg and 15 mg and the risperidone between 2 mg and 6 mg oral liquid and oral velotab. The thiothixene trial used higher doses of both drugs (~30 mg) given by IM injection. The ziprasidone studies used rather high dose haloperidol (20 mg up to 40 mg) and compared this with around 40 mg of ziprasidone oral and IM. Haloperidol dose in the zuclopenthixol acetate trial was also high (~30 mg) compared with up to ~100 mg IM of the longer-acting compound).

5.3 Versus benzodiazepine

The one trial comparing IM flunitrazepam with IM haloperidol used about 1 mg of the benzodiazepine and 5 mg of the antipsychotic. The lorazepam trials, using a combination of oral and IM administration employed doses of around 4 mg to 6 mg of the drug. The comparison dose was around 10 mg of haloperidol. The single midazolam study compared 5 mg with 5 mg (both IM).

5.4 Versus combinations

The haloperidol versus haloperidol plus levomepromazine study were oral treatments, but we remain unclear as to the doses. These seem to be around 5 mg for each compound. For the haloperidol versus haloperidol plus lorazepam comparison, the antipsychotic tended to be given at around the 5 mg level for both groups and the benzodiazepine at 4 mg each. In the haloperidol versus haloperidol plus midazolam study, the antipsychotic at 5 mg was compared with the adjunction of 15 mg of midazolam. In the haloperidol versus haloperidol plus promethazine trial, all treatments were IM. The haloperidol was up to 10 mg and the promethazine up to 50 mg. The haloperidol versus haloperidol plus risperidone study compared 10 mg to 20 mg of haloperidol with 3 mg to 6 mg of risperidone. For the haloperidol versus olanzapine plus magnesium valproate comparison, the first treatment tended to be given at around 15 mg and the second one at respectively 10 mg plus 1.25 mg. In the haloperidol versus quetiapine (IM) plus magnesium valproate (oral) comparison, the dose of haloperidol was up to 30 mg/day and the quetiapine's 800 mg/day. The additional magnesium valproate was dosed at about 0.5 g to 1 g/orally/day. In the haloperidol versus risperidone plus clonazepam comparison,

haloperidol was administered at 10 mg to 20, mg whilst risperidone plus clonazepam doses ranged from, respectively, 2 mg to 6 mg plus 0 mg to 8 mg. Finally, the haloperidol versus ziprasidone plus clonazepam study compared up to 30 mg of haloperidol versus up to 160 mg of ziprasidone and up to 6 mg of clonazepam.

6. Outcomes

For the majority of trials, binary data were available with respect to repeated need for tranquillisation, not tranquil or asleep, need for adjunctive medication, leaving the study early and adverse effects. The majority of trials employed continuous scales to measure other clinical outcomes. However, the results of these scale-derived findings were often presented in graphs, as P values alone, or as means without standard deviations. Much data were unusable for synthesis. Requests for further information from the majority of authors remain unanswered. Dr Dubin (Dubin 1985) kindly answered our request, however was unable to provide us with the standard deviation or standard error.

The various rating scales, from which we were able to obtain usable data are listed below.

6.1 Specific behaviour - Agitation

a. Agitated Behavior Scale (ABS/CABS)

The ABS (Corrigan 1989) is a 14-item scale used for measuring agitation levels. The ABS was originally designed to monitor agitated behaviour in the recovery period after stroke and there are three factor-based sub-scores: i. aggression, ii. disinhibition; and iii. lability. High scores indicated higher levels of aggression.

b. Agitation-Calmness Evaluation Scale (ACES)

The ACES is a single-item rating scale developed by Eli Lilly and company (Breier 2002). On this scale, 1 = marked agitation, 4 = normal, 9 = unable to be aroused.

c. Positive and Negative Syndrome Scale Excited Component (PANSS-EC)

The PANSS-EC is a five-item scale (excitement, tension, hostility, uncooperativeness, and poor impulse control). The items are rated from one (not present) to seven (extremely severe). Scores range from five to 35, with mean scores ≥ 20 indicating agitation. A high score indicates high levels of agitation (Montoya 2011).

d. Overt Agitation Severity scale (OASS)

The OASS (Yudofsky 1997) is a 12-item scale used for identification and measurement of agitated behaviours, rated with a intensity score (ranging from one to four); the items are rated in terms of frequency from zero (not present) to four (always present). Total OASS score is the sum of all the Severity Scores (intensity*frequency).

6.2 Specific behaviour - Aggression

a. Overt Aggression Scale (OAS)

The OAS (Yudofsky 1986) is a 16-item rating scale which aims to measure the intensity of verbal and physical aggression. Clinicians are required to comment on the duration of the aggressive incident as well as the intervention required to control it. High scores are indicative of higher levels of aggression.

6.3 Global state

a. Clinical Global Impression (CGI)

The CGI (Guy 1976) is not a diagnostic tool but rather, enables clinicians to quantify the severity of symptoms of any mental health problem at one point in time. Clinicians are then able to use this to track whether there has been any improvement or worsening of symptoms over time. A seven-point rating scale is used with high scores indicating increased severity or less recovery.

b. Clinical Global Impression - Severity (CGI-S)

The CGI-S (Guy 1976) requires clinicians to consider the severity of a person's symptoms in relation to the clinicians past experience of people with the same diagnosis. Clinicians then have to give a rating from one = normal to seven = extremely ill. High scores indicate increased severity.

c. Clinical Global Impression - Improvement (CGI-I)

The CGI-I (Guy 1976) enables clinicians to assess whether a person's symptoms have improved or worsened following an intervention. Based on the clinicians judgement, a rating on a seven-point scale is given from one (= very much improved) to seven (= very much worse). Therefore, low scores indicate greater improvement.

6.4 Mental state

a. Brief Psychiatric Rating Scale (BPRS)

The Brief Psychiatric Rating Scale was originally developed by Overall and Gorham (Overall 1962) as a 14-item scale to measure the severity of a range of psychiatric symptoms, including psychosis. This rating scale items evolved over time and now consists of 24 items which can be rated on a seven-point scale from 'not present' to 'extremely severe'. A high score would suggest poor mental health. It is not clear for the majority of the studies included in this review, which version of the BPRS was used.

b. Positive and Negative Syndrome Scale (PANSS)

The PANSS was developed and published by Kay, Flszbein and Opler (Kay 1987). The PANSS is designed as a brief interview, whereby the severity of 30 symptoms of schizophrenia can be assessed on a scale of one to seven. A high score would indicate more severe symptoms. The PANSS can be divided into separate sub-scales by focusing on the statements relating to positive symptoms (e.g. hallucinations), negative symptoms (e.g. social withdrawal) or general psychopathology (e.g. anxiety and uncooperativeness).

6.5 Adverse effects

a. Barnes Akathisia Scale (BAS)

The BAS scale was developed by Barnes 1989 and includes items which aim to rate both the observable symptoms which characterise akathisia such as restless movements and also the person's subjective experience, including any distress. The items are rated from zero = normal to three = severe. There is also an item for rating global severity from zero = absent to five = severe. A high score indicates high levels of akathisia.

b. Simpson-Angus Scale (SAS)

The SAS (Simpson 1970) is a 10-item scale which measures drug-induced parkinsonism (extrapyramidal side effects). Each item is

scored from zero to four. A high score would indicate increased levels of parkinsonism.

c. Treatment Emergent Symptom Scale (TESS)

The TESS (Guy 1976) is a six-item scale which is used to assess the occurrence and intensity of treatment-related adverse effects. A high score indicates worse symptoms. This scale was not reported as a continuous score but studies did glean binary data using this measure (e.g. Li 2006).

7. Missing outcomes

Not one of the studies evaluated satisfaction with care, acceptance of treatment, quality of life or economic outcomes.

8. Funders

Fifteen of the 41 included studies received sponsorship from pharmaceutical companies.

Excluded studies

In total we excluded 66 studies.

Of these, 20 studies had to be excluded based on their method of allocation. NCT00189995 2005 was not conducted because ethical approval was not granted and Daniel 2000, Jones 2001 and Mendelowitz 2004 were literature reviews. Chen 2004, Chen 2008, Fan 2012; Huang 2004, Jiang 2009; Li 2007, Liang 2003; Pan 2005, Pei 2009; Wang 2004 and Wang 2005 were all quasi-randomised whereby randomisation was based on methods such as allocating to group by hospital number. It is possible that these studies were randomised if allocation of hospital numbers were somehow, designated by a procedure involving chance alone. We were, however, not reassured by the text of the trials, thus, we have excluded this quasi-randomised group. Currier 2000, Freeman 2009, Kinon 2003, Mantua 2006 and Pedros 2004 used research methods other than a randomised controlled trial, for example a naturalistic observational study or cohort study.

We had to exclude 34 studies on the basis of the characteristics of participants. Singh 1981 and Zapletalek 1986 were not explicitly described as randomised, however, they were reported to be double-blind and thus remained included on the basis of allocation. They were eventually excluded because although participants were suspected to have psychosis, these people were not overtly agitated or aggressive requiring rapid tranquillisation. We excluded a group of clearly randomised trials for the same

reason (Addington 1996; Allen 2007; Beasley 1998; Brook 1998; Chouinard 1991; Colonna 1998; Crandall 2005; Daniel 2003; Daniel 2004; Goldstein 1966; Kane 2000; Kinon 2001a; Mandel 2008; McEvoy 1994; NCT00631722 2008; Pathiraja 1995; Puech 1998; Smith 1974; Teja 1975; Ungvari 1982; Veser 2006). Participants included in Harvey 2004 had mild symptoms of schizophrenia and, again, were not overtly aggressive. We identified two studies where participants were children. We excluded these because either, participants did not display aggression (Farettra 1970), or because less than half of participants were suspected to have psychosis (Asadollahi 2015; Campbell 1982).

In Citrome 2001, Krakowski 2006, Krakowski 2008 and Lewis 2006, participants had a diagnosis of schizophrenia and displayed aggressive behaviour. However, the studies had to be excluded because participants displayed long-term aggression, rather than aggression within an emergency situation requiring rapid tranquillisation. Participants in Romeo 2009 and Vaisanen 1981 were people with intellectual disability who presented with challenging behaviour rather than psychosis. Thomas 1992 focused on rapid tranquillisation in the emergency setting, but the majority of participants were intoxicated and therefore did not necessarily have psychosis-induced aggression.

For 10 studies the intervention under investigation did not meet our inclusion criteria. Alexander 2004, Bieniek 1998, Conde 2011, TREC CG, NCT00797277 2008, Simpson 2010 and Srinath 2010 did not include a comparison group which involved giving of haloperidol alone; Vives 2015 did not include a comparison group which involved giving a drug out of haloperidol; Liu 2012b compared haloperidol with a non-drug methodology of intervention; in Li 2005 there was no specific outcome for agitation.

Finally, two relevant studies were excluded because we found no usable data. Participants were randomised and had a diagnosis of schizophrenia with agitation/aggression. However, in Wan 2005 the scales that were used to assess outcome were not validated and results for haloperidol and chlorpromazine were not presented separately. In Wyant 1990 they used a "modified CGI" for which we could find no evidence of validation and then reported mean data graphically rather than numerically.

Risk of bias in included studies

Although each 'Summary of findings' table describes bias by outcome, an overview is reported here and a graphical impression is seen in Figure 3.

Figure 3. 'Risk of bias' summary: review authors' judgements about each risk of bias item for each included study.

	Random sequence generation (selection bias)	Allocation concealment (selection bias)	Blinding of participants and personnel (performance bias)	Blinding of outcome assessment (detection bias)	Incomplete outcome data (attrition bias)	Selective reporting (reporting bias)	Other bias
Bailine 1987	+	?	?	?	?	-	-
Baldacara 2011	?	?	-	-	+	-	+
Battaglia 1997	+	?	+	+	-	-	-
Battaglia 2002	?	?	+	+	-	+	-
Breier 2002	?	+	+	+	-	?	-
Bristol Myers 2004a	?	+	?	?	-	-	-
Bristol-Myers 2005b	?	?	-	+	?	-	-
Brook 2000	+	?	-	-	+	-	-
Calver 2015	+	+	+	+	+	?	-
Currier 2004	?	+	-	+	?	-	-
Dorevitch 1999	+	?	?	+	+	?	?
Eli Lilly 2004	?	?	?	?	-	-	-
Fang 2012	+	?	-	-	+	-	-
Fitzgerald 1969	?	?	+	?	?	?	?
Foster 1997	?	?	?	+	+	-	-
Fruensgaard 1977	?	?	+	+	+	+	-
Garza-Trevino 1989	?	?	-	-	+	-	?
Guo 2007	?	?	-	-	+	-	?
Higashima 2004	?	?	-	-	+	-	-
Huf 2007	+	+	-	-	+	+	?

Figure 3. (Continued)

Huf 2007	+	+	-	-	+	+	?
Kewala 1984	?	?	?	?	+	?	-
Kinon 2001d	?	?	?	?	+	-	-
Li 2006	?	?	?	?	+	?	?
Liang 2013	+	?	?	?	-	+	?
Lim 2010	+	?	-	+	+	-	-
Liu 2012	+	?	?	?	+	+	+
Man 1973	-	?	?	?	-	+	-
Nobay 2004	+	+	+	+	+	+	-
Paprocki 1977	?	?	+	?	?	+	-
Reschke 1974	?	?	?	?	+	+	?
Resnick 1984	?	+	+	?	+	+	-
Ritter 1972	?	?	+	?	+	-	?
Salzman 1991	-	?	+	+	-	+	-
Shi 2010	+	?	?	?	+	+	+
Shu 2010	?	?	-	+	?	?	-
Stotsky 1977	?	?	+	+	+	+	-
Taymeeyapradit 2002	?	?	-	?	+	+	?
Tuason 1986	?	?	-	+	-	-	?
Walther 2014	+	+	?	?	-	?	?
Yin 2012	?	?	?	?	+	+	+
Zhou 2014	+	?	?	?	-	+	+

Allocation

All but two of the included studies were said to be randomised. [Man 1973](#) and [Salzman 1991](#) were not explicitly described as randomised, however they were described as double-blind and we felt randomisation was implied. Few of the randomised studies reported information regarding the method of allocation. For the studies that did report further details, methods are described as computer-generated randomisation ([Battaglia 1997](#); [Brook 2000](#); [Huf 2007](#); [Nobay 2004](#)), random number sequence ([Bailine 1987](#); [Dorevitch 1999](#); [Liang 2013](#); [Liu 2012](#); [Shi 2010](#); [Zhou 2014](#)), pre-defined randomisation code ([Lim 2010](#)) and randomisation in blocks ([Stotsky 1977](#); [Walther 2014](#)).

Blinding

Twenty-two studies described that they were 'double-blind' but the variations on how this was carried out were almost as many ([Table](#)

[2](#)); some studies were reported to be 'single-blind' ([Baldacara 2011](#); [Yin 2012](#)), whilst in five studies it was not stated ([Liang 2013](#); [Liu 2012](#); [Shi 2010](#); [Walther 2014](#); [Zhou 2014](#)). Trialists were more or less concerned with issues of performance and detection bias and most 'double-blind' studies are likely not to have been in reality. Open studies ([Brook 2000](#); [Fang 2012](#); [Garza-Trevino 1989](#); [Guo 2007](#); [Higashima 2004](#); [Huf 2007](#)) recognised that blinded giving of drugs added the possibility that findings may be problematic to apply in routine care. These studies also attempted to use outcomes that were less prone to rater bias.

Incomplete outcome data

There was no evidence of attrition bias for approximately two thirds of the studies with all participants being accounted for ([Baldacara 2011](#); [Brook 2000](#); [Calver 2015](#); [Dorevitch 1999](#); [Fang 2012](#); [Foster 1997](#); [Fruensgaard 1977](#); [Garza-Trevino 1989](#); [Guo 2007](#); [Higashima 2004](#); [Huf 2007](#); [Kewala 1984](#); [Kinon 2001d](#); [Li 2006](#); [Lim 2010](#); [Liu](#)

2012; Nobay 2004; Reschke 1974; Resnick 1984; Ritter 1972; Shi 2010; Stotsky 1977; Taymeeyapradit 2002; Yin 2012).

In the [Bristol-Myers 2005b](#), [Currier 2004](#) and [Shu 2010](#) studies, the reasons why participants left early are clearly presented in a table, but the numbers of participants completing different assessments at the same time point vary and reasons are not given. Other studies where attrition is not clearly described are [Bailine 1987](#), [Fitzgerald 1969](#) and [Paprocki 1977](#).

In [Battaglia 1997](#), two people were excluded from the efficacy analysis post randomisation; in [Walther 2014](#) nine people were excluded due to refusal to provide post-hoc consent and in three studies ([Man 1973](#); [Liang 2013](#); [Zhou 2014](#)), although reasons are given for why participants discontinued, results are only presented for those who completed and missing data have not been imputed. Reasons why participants left the study early were not reported in [Battaglia 2002](#), [Breier 2002](#), [Bristol Myers 2004a](#), [Eli Lilly 2004](#), [Salzman 1991](#) and [Tuason 1986](#). Also, attrition approached 50% at 24 hours in [Salzman 1991](#), and was over 50% in [Tuason 1986](#) at 10 days for some outcomes.

Selective reporting

Overall, there was much redundancy in reporting. It is difficult to judge whether this is bias or inadvertent but even-handed poor reproduction of data. For continuous data, reasons for data being rendered unusable where often that the mean was presented but no variance could be extracted. Also, inexact P values were often presented alone and several studies presented their findings in graphs from which we could not estimate numbers.

Even simple issues were not reported well. For example, use of additional medication can indicate how treatments are not effective and need to be supplemented or that adverse effects are prevalent. The amount of additional benzodiazepine in the [Foster 1997](#) study, the amount of anticholinergic and hypnotic medication in [Tuason 1986](#), and the amount of antiparkinson drugs administered in [Lim 2010](#) and [Walther 2014](#) are not reported. Further, there is inconsistent reporting regarding additional medications in [Brook 2000](#) and [Tuason 1986](#).

In general, there was poor reporting with regard to adverse effects. In many studies it is not clear whether all side effects are reported ([Breier 2002](#); [Dorevitch 1999](#); [Fitzgerald 1969](#); [Garza-Trevino 1989](#); [Higashima 2004](#); [Ritter 1972](#)). Further, in other trials adverse effects are only reported where they occurred in a particular percentage of participants ([Battaglia 1997](#); [Bristol Myers 2004a](#); [Bristol-Myers 2005b](#); [Brook 2000](#); [Currier 2004](#); [Fang 2012](#); [Fitzgerald 1969](#); [Kinon 2001d](#); [Shu 2010](#)). This further undermines methodology that is already weak to pick up adverse effect that may not be common, but could well be important.

Other potential sources of bias

That a study is small or sponsored by industry (or both) is not indicative of bias in itself. There are, however, associations between small trials tending to be positive ([Easterbrook 1991](#)), and those sponsored by industry favouring the drug for which they have the pecuniary interest ([Lexchin 2003](#)). Many studies in this review were small ([Bailine 1987](#); [Dorevitch 1999](#); [Foster 1997](#); [Fruensgaard 1977](#); [Higashima 2004](#); [Man 1973](#); [Paprocki 1977](#); [Resnick 1984](#); [Stotsky 1977](#); [Walther 2014](#)), many were sponsored by industry ([Battaglia 1997](#); [Battaglia 2002](#); [Breier 2002](#); [Bristol Myers 2004a](#); [Bristol-Myers](#)

[2005b](#); [Brook 2000](#); [Currier 2004](#); [Eli Lilly 2004](#); [Fang 2012](#); [Kewala 1984](#); [Kinon 2001d](#); [Lim 2010](#); [Salzman 1991](#); [Shu 2010](#); [Stotsky 1977](#)), and one was both ([Stotsky 1977](#)).

Effects of interventions

See: **Summary of findings for the main comparison**
HALOPERIDOL compared to PLACEBO/NIL for psychosis-induced aggression or agitation; **Summary of findings 2** HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: a. ARIPIRAZOLE for psychosis-induced aggression or agitation; **Summary of findings 3** HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: b. CHLORPROMAZINE for psychosis-induced aggression or agitation; **Summary of findings 4** HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: c. DROPERIDOL for psychosis-induced aggression or agitation; **Summary of findings 5** HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: d. LOXAPINE for psychosis-induced aggression or agitation; **Summary of findings 6** HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: e. OLANZAPINE for psychosis induced aggression or agitation; **Summary of findings 7** HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: f. PERPHENAZINE for psychosis-induced aggression or agitation; **Summary of findings 8** HALOPERIDOL compared to OTHER ANTIPSYCHOTICS: g. QUETIAPINE for psychosis-induced aggression or agitation; **Summary of findings 9** HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: h. RISPERIDONE for psychosis-induced aggression or agitation; **Summary of findings 10** HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: i. THIOTHIXENE for psychosis-induced aggression or agitation; **Summary of findings 11** HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE for psychosis-induced aggression or agitation; **Summary of findings 12** HALOPERIDOL compared to OTHER ANTIPSYCHOTIC: k. ZUCLOPENTHIXOL ACETATE for psychosis-induced aggression or agitation; **Summary of findings 13** HALOPERIDOL compared to BENZODIAZEPINE: a. FLUNITRAZEPAM for psychosis-induced aggression or agitation; **Summary of findings 14** HALOPERIDOL compared to BENZODIAZEPINE: b. LORAZEPAM for psychosis-induced aggression or agitation; **Summary of findings 15** HALOPERIDOL compared to BENZODIAZEPINE: c. MIDAZOLAM for psychosis-induced aggression or agitation; **Summary of findings 16** HALOPERIDOL compared to COMBINATIONS: a. HALOPERIDOL + LEVOMEPRIMAZINE for psychosis-induced aggression or agitation; **Summary of findings 17** HALOPERIDOL compared to COMBINATIONS: b. HALOPERIDOL + LORAZEPAM for psychosis-induced aggression or agitation; **Summary of findings 18** HALOPERIDOL compared to COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM for psychosis-induced aggression or agitation; **Summary of findings 19** HALOPERIDOL compared to COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE for psychosis-induced aggression or agitation; **Summary of findings 20** HALOPERIDOL compared to COMBINATIONS: e. HALOPERIDOL + RISPERIDONE for psychosis-induced aggression or agitation; **Summary of findings 21** HALOPERIDOL compared to COMBINATIONS: f. OLANZAPINE + MAGNESIUM VALPROATE for psychosis-induced aggression or agitation; **Summary of findings 22** HALOPERIDOL compared to COMBINATIONS: g. QUETIAPINE + MAGNESIUM VALPROATE for psychosis-induced aggression or agitation; **Summary of findings 23** HALOPERIDOL compared to COMBINATIONS: h. RISPERIDONE + CLONAZEPAM for psychosis induced aggression or agitation; **Summary of findings 24**

HALOPERIDOL compared to COMBINATIONS: i. ZIPRASIDONE + CLONAZEPAM for psychosis-induced aggression or agitation

In the following section, the summary statistic for binary outcome results is reported as risk ratio (RR) and for continuous outcomes, we used mean difference (MD). In all instances 95% confidence intervals (CIs) are reported.

1. COMPARISON 1. HALOPERIDOL versus PLACEBO

Five studies (total $n = 700$) compared haloperidol with placebo (Battaglia 2002; Breier 2002; Bristol Myers 2004a; Bristol-Myers 2005b; Reschke 1974).

1.1 Tranquillisation or asleep - asleep

Significantly more people in the haloperidol group were asleep at two hours compared with those in the placebo group (2 RCTs, $n = 220$, RR 0.88, 95% CI 0.82 to 0.95, $I^2 = 76\%$, Analysis 1.1).

1.2 Repeated need for tranquillisation

More people in the placebo group required repeated rapid tranquillisation compared with those in the haloperidol group (4 RCTs, $n = 660$, RR 0.51, 95% CI 0.42 to 0.62, Analysis 1.2).

1.3 Specific behaviour - agitation

Two studies ($n = 395$) found that significantly more people in the haloperidol group were reported as demonstrating at least a 40% reduction in agitation scores at two hours according to the PANSS-EC response at two hours (RR 1.62, 95% CI 1.28 to 2.07, Analysis 1.3).

Studies used a series of measures at different time periods. By about two hours, participants in the haloperidol group were reported to have experienced a significantly greater reduction in agitation than those in the placebo group according to the Agitated Behavior Scale (ABS) (CABS) rating scale (3 RCTs, $n = 474$, MD -4.48, 95% CI -5.78 to -3.18). In the same way, again at about two hours, people allocated to placebo had significantly smaller average change scores on the Agitation-Calmness Evaluation Scale (ACES) compared with those in the haloperidol group (2 RCTs, $n = 390$, MD 0.83, 95% CI 0.39 to 1.26). One study ($n = 199$) examined the difference in the Positive and Negative Syndrome Scale Excited Component (PANSS-EC) scores between the haloperidol and placebo groups in the schizophrenia subgroup, in non-sedated people based on the ACES and in non-sedated people. People who received haloperidol had significantly improved scores for agitation when compared with placebo in all three of these subgroups (MD -2.57, 95% CI -4.05 to -1.09, MD -2.64, 95% CI -3.95 to -1.33, and MD -2.97, 95% CI -4.27 to -1.67, respectively, Analysis 1.4). Again, according to average change and endpoint combined scores on the PANSS-EC, participants in the haloperidol group had significantly reduced scores for agitation (3 RCTs, $n = 514$, MD -3.67, 95% CI -4.75 to -2.59, Analysis 1.4).

By 24 hours, average change on the ABS (CABS) rating scale was significantly changed in favouring haloperidol (1 RCT, $n = 85$, MD -2.40, 95% CI -4.13 to -0.67). This was not found for the PANSS-EC average change scores (1 RCT, $n = 85$, MD -1.40, 95% CI -2.97 to 0.17, Analysis 1.5).

1.4 Global outcome

Significantly more people in the placebo group exhibited 'not marked improvement' than those who received haloperidol (1 RCT,

$n = 40$, RR 0.61, 95% CI 0.44 to 0.84, Analysis 1.6). Further, people in the placebo group were more likely to be rated as showing 'not any improvement' in global functioning than those receiving haloperidol, however this difference was not statistically significant (1 RCT, $n = 40$, RR 0.28, 95% CI 0.08 to 1.07, Analysis 1.6).

Four studies found that more people in the placebo group required benzodiazepines ($n = 660$, RR 0.44, 95% CI 0.31 to 0.62, Analysis 1.7).

Global rating took place around two hours and 24 hours. By two hours people who received haloperidol were reported to have had significantly more symptom improvement according to their average endpoint scores on the Clinical Global Impression - Improvement (CGI-I), when compared with placebo (2 RCTs, $n = 390$, MD -0.73, 95% CI -0.96 to -0.51), and in the schizophrenia subgroup (1 RCT, $n = 199$, MD -0.64, 95% CI -0.98 to -0.30). When the same outcome was measured using average change score on the Clinical Global Impression - Severity (CGI-S), participants in the haloperidol group were rated as having significantly less severe symptoms than those in the placebo group (2 RCTs, $n = 390$, MD -0.48, 95% CI -0.78 to -0.18, Analysis 1.8).

By 24 hours, average change scores on the CGI-I remained significantly improved (1 RCT, $n = 169$, MD -0.40, 95% CI -0.62 to -0.18). Using the CGI-S scale to rate symptom severity at 24 hours, one trial ($n = 85$) found that although those in the haloperidol group were rated as less severe, the difference was not statistically significant (MD -0.20, 95% CI -0.46 to 0.06, Analysis 1.9).

1.5 Mental state

Participants in the haloperidol group had significantly improved average change scores on the Brief Psychiatric Rating Scale (BPRS) positive sub-scale when rated at two (3 RCTs, $n = 372$, MD -1.04, 95% CI -1.54 to -0.54, Analysis 1.10) and 24 hours (2 RCTs, $n = 264$, MD -1.61, 95% CI -2.34 to -0.89, Analysis 1.11). On the BPRS Total sub-scale, participants in the haloperidol group had significantly improved scores when rated at two (3 RCTs, $n = 371$, MD -5.69, 95% CI -7.38 to -4.00, Analysis 1.10) and 24 hours (2 RCTs, $n = 264$, MD -4.81, 95% CI -6.82 to -2.81, Analysis 1.11).

1.6 Adverse effects

1.6.1 General

Bristol Myers 2004a ($n = 273$) specifically reported that no deaths occurred. Significantly more people in the haloperidol group experienced one or more adverse reactions than those in the placebo group (2 RCTs, $n = 395$, RR 1.64, 95% CI 1.22 to 2.20). One trial ($n = 273$) found that people in the haloperidol group were significantly more likely than those in the placebo group to report overall adverse effects at 72 hours (RR 1.78, 95% CI 1.23 to 2.59) and to experience increased severity of adverse effects after the second injection (RR 3.25, 95% CI 1.88 to 5.63, Analysis 1.12).

Bristol-Myers 2005b reported that of all adverse effects reported, only one was rated as 'serious' - this was related to a participant in the placebo group. This difference was not statistically significant ($n = 122$, RR 0.34, 95% CI 0.01 to 8.29, Analysis 1.13).

1.6.2 Specific

a. Arousal

Insomnia was reported by people in both the haloperidol and placebo groups and the difference between them was not

statistically significant (1 RCT, $n = 273$, RR 1.31, 95% CI 0.61 to 2.82). Somnolence was reported in both the placebo and haloperidol group and the difference between the groups was not statistically significant (4 RCTs, $n = 615$, RR 2.28, 95% CI 0.97 to 5.36, [Analysis 1.14](#)). Significantly more people in the haloperidol group than the placebo group were reported to be "over sedated" (2 RCTs, $n = 313$, RR 3.36, 95% CI 1.42 to 7.99, [Analysis 1.14](#)).

b. Cardiac

We included dizziness in 'cardiac' adverse effects (2 RCTs, $n = 392$, RR 1.33, 95% CI 0.48 to 3.65). More specific cardiac effects also did not suggest an effect for the active drug (QTc abnormality, 1 RCT, $n = 122$, RR 3.10, 95% CI 0.33 to 28.98; sinus tachycardia, 1 RCT, $n = 122$, RR 3.10, 95% CI 0.33 to 28.98; tachycardia, 1 RCT, $n = 122$, RR 1.03, 95% CI 0.07 to 16.15, [Analysis 1.15](#)). Although in favour of placebo, there were no statistically significant differences between the haloperidol and placebo groups with regards to QTc abnormality according to the average change score (2 RCTs, $n = 265$, MD 3.63, 95% CI -2.67 to 9.93), when rated at 24 hours ([Analysis 1.16](#)).

c. Movement disorders

In the haloperidol group, six people reported experiencing akathisia within 24 hours, compared with none in the placebo group. This difference between groups was not statistically significant (1 RCT, $n = 122$, RR 13.43, 95% CI 0.77 to 233.23). Dystonia was reported in the haloperidol group within 24 hours by six people compared with none in the placebo group; the difference between groups was not statistically significant (2 RCTs, $n = 207$, RR 7.49, 95% CI 0.93 to 60.21). Significantly more people in the haloperidol group required antiparkinson drugs due to Extrapyramidal Side Effects (EPS) (1 RCT, $n = 180$, RR 5.57, 95% CI 1.37 to 22.65) with significantly more people experiencing EPS during 24 hours (3 RCTs, $n = 398$, RR 6.79, 95% CI 2.19 to 21.07), [Analysis 1.17](#)).

There was no statistically significant difference between the average change scores on the Barnes Akathisia Scale (BAS) for participants in the haloperidol group when compared with the placebo group (1 RCT, $n = 168$, MD 0.09, 95% CI -0.17 to 0.35, [Analysis 1.18](#)). In one study ($n = 167$) participants in the haloperidol group experienced significantly higher levels of EPS compared with participants in the placebo group according to their average change scores on the Simpson-Angus Scale (SAS) (MD 1.89, 95% CI 0.77 to 3.01, [Analysis 1.18](#)).

d. Miscellaneous

Two studies (total $n = 395$) listed a range of effects including agitation (2 RCTs, RR 0.80, 95% CI 0.29 to 2.19), headache (2 RCTs, RR 1.28, 95% CI 0.55 to 3.00), nausea (2 RCTs, RR 0.69, 95% CI 0.13 to 3.67) and vomiting (1 RCT, $n = 122$, RR 1.03, 95% CI 0.07 to 16.15). Although not statistically significant, two people in the placebo group reported pain at the injection site compared with none in the haloperidol group (1 RCT, $n = 122$, RR 0.21, 95% CI 0.01 to 4.22). This changed to significantly more people in the haloperidol group complaining of pain at the injection site following the second injection, when compared with the placebo group (44% versus 14%, 1 RCT, $n = 273$, RR 3.25, 95% CI 1.88 to 5.63, [Analysis 1.19](#)).

1.7 Leaving the study early

In three studies (total $n = 435$), although more people in the haloperidol group left the study early (for any reason) when

compared with the placebo group, the difference was not statistically significant (RR 1.06, 95% CI 0.53 to 2.11, $I^2 = 83\%$). There were significant differences between groups for number of participants who left early due to lack of efficacy (3 RCTs, $n = 435$, RR 0.12, 95% CI 0.03 to 0.49), but not for reasons of consent (2 RCTs, $n = 395$, RR 6.20, 95% CI 0.77 to 49.98), adverse effects (3 RCTs, $n = 435$, RR 1.78, 95% CI 0.21 to 15.39), being asleep (1 RCT, $n = 40$, RR 1.20, 95% CI 0.05 to 27.44), or 'other' reasons (2 RCTs, $n = 395$, RR 1.81, 95% CI 0.27 to 12.07). In one study ($n = 273$), three people in the haloperidol group were withdrawn from the study compared with none in the placebo, although the difference was not statistically significant (RR 3.35, 95% CI 0.17 to 64.15, [Analysis 1.20](#)).

2. COMPARISON 2. HALOPERIDOL versus OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE

We found two studies comparing haloperidol with aripiprazole (total $n = 599$, [Bristol Myers 2004a](#); [Bristol-Myers 2005b](#)).

2.1 Repeated need for rapid tranquillisation

In both studies, people in the haloperidol group required fewer injections than those in the aripiprazole, the difference between the groups being statistically significant (2 RCTs, $n = 473$, RR 0.78, 95% CI 0.62 to 0.99, [Analysis 2.1](#)).

2.2 Specific behaviours - agitation

In both studies, more people in the aripiprazole group demonstrated at least a 40% reduction in agitation scores (PANSS-EC) at two hours than in the haloperidol group, though the difference between the groups was not statistically significant (2 RCTs, $n = 477$, RR 1.07, 95% CI 0.92 to 1.26, [Analysis 2.2](#)).

At two hours, participants showed similar average change scores according to the Agitation-Calmness Evaluation Scale (ACES) rating scale (2 RCTs, $n = 470$, MD 0.12, 95% CI -0.30 to 0.55). Also, at two hours, participants showed similar average change scores according to the CABS rating scale (2 RCTs, $n = 470$, MD -0.55, 95% CI -2.10 to 1.01). One study found that no differences between the haloperidol and aripiprazole groups in average change scores on the PANSS-EC ($n = 357$, MD -0.48, 95% CI -2.12 to 1.16). This study also examined whether there were differences in various subgroups (schizophrenia subgroup, non-sedated people based on the ACES and non-sedated). There were no significant differences between haloperidol and aripiprazole within any of these subgroups ([Analysis 2.3](#)).

2.3 Global outcomes

Neither those allocated haloperidol or aripiprazole required additional benzodiazepines (2 RCTs, $n = 477$, RR 1.26, 95% CI 0.74 to 2.16, [Analysis 2.4](#)).

At two hours, participants in both groups showed similar levels of severity of symptoms according to their average change scores on the CGI-S rating scale (2 RCTs, $n = 470$, MD 0.07, 95% CI -0.22 to 0.36). On another scale, at the same time, participants showed similar levels of symptom improvement according to their average endpoint scores on the CGI-I rating scale (2 RCTs, $n = 470$, MD -0.02, 95% CI -0.23 to 0.19). Again, at two hours, people in the haloperidol group (schizophrenia subgroup), were not rated as more improved on the CGI-I than those in the aripiprazole group (1 RCT, $n = 257$, MD -0.02, 95% CI -0.30 to 0.26, [Analysis 2.5](#)).

2.4 Mental state

One study (n = 103) found no difference for average change scores for participants on the BPRS (positive sub-scale MD -0.23, 95% CI -1.28 to 0.82 and total score n = 102, MD -2.03, 95% CI -5.76 to 1.70, [Analysis 2.6](#)).

2.5 Adverse effects

2.5.1 General

Studies found no difference between groups for 'adverse effects within 24 hours' (2 RCTs, n = 477, RR 1.18, 95% CI 0.95 to 1.46). However, significantly more people in the haloperidol group suffered with increased severity of adverse effects after the second injection (1 RCT, n = 360, RR 1.34, 95% CI 1.03 to 1.74), and overall adverse effects at 72 hours (1 RCT, n = 360, RR 1.33, 95% CI 1.04 to 1.70, [Analysis 2.7](#)).

There were no significant differences between the groups with respect to 'serious' adverse effects ([Analysis 2.8](#)). In the [Bristol Myers 2004a](#) study, it was reported that no participants in either group died during the study. Four people developed severe adverse effects; three in the aripiprazole group compared with none in the haloperidol group (2 RCTs, n = 477, RR 0.57, 95% CI 0.18 to 1.81). In the [Bristol Myers 2004a](#) study, one person in the haloperidol group developed severe dystonia and in the [Bristol-Myers 2005b](#) study, one person in the aripiprazole group had a tonic-clonic seizure.

2.5.2 Specific

a. Arousal

Significantly more people in the haloperidol group complained of insomnia during the intramuscular (IM) phase (1 RCT, n = 360, RR 2.08, 95% CI 1.01 to 4.27). There were no clear differences between aripiprazole and haloperidol for insomnia during the oral phase (1 RCT, n = 360, RR 0.82, 95% CI 0.47 to 1.44), and for people being recorded as being "over sedated" during the study (1 RCT, n = 360, RR 1.66, 95% CI 0.93 to 2.95). With regards to the various ways of reporting somnolence, again, no difference was apparent (e.g. 1 RCT, n = 360, RR somnolence - between zero to one days 0.71, 95% CI 0.25 to 2.00, [Analysis 2.9](#)).

b. Cardiac

Similar numbers of people in each group experienced dizziness (2 RCTs, n = 477, RR 0.69, 95% CI 0.33 to 1.48), sinus tachycardia (1 RCT, n = 117, RR 6.66, 95% CI 0.35 to 126.06) and tachycardia (1 RCT, n = 117, RR 0.24, 95% CI 0.03 to 2.06, [Analysis 2.10](#)).

c. Gastrointestinal problems

Participants in the haloperidol groups complained of significantly more dyspepsia (1 RCT, n = 360, RR 10.41, 95% CI 1.36 to 79.76), and similar problems with diarrhoea (1 RCT, n = 360, RR 0.95, 95% CI 0.28 to 3.21), nausea (1 RCT, n = 360, RR 0.14, 95% CI 0.02 to 1.09), and vomiting (1 RCT, n = 360, RR 0.38, 95% CI 0.07 to 1.92) between three and seven days ([Analysis 2.11](#)).

d. Movement disorder

These all tended to be more common in the group allocated to haloperidol (e.g. 2 RCTs, n = 477, RR akathisia 2.85, 95% CI 0.94 to 8.61). More people in the haloperidol group were reported to have experienced dystonia (2 RCTs, n = 477, RR 6.63, 95% CI 1.52 to 28.86). Significantly more people in the haloperidol group experienced

EPS during both the IM (1 RCT, n = 360, RR 9.46, 95% CI 1.22 to 73.13) and oral phase (1 RCT, n = 360, RR 7.09, 95% CI 1.65 to 30.58), and required antiparkinson medication during the oral phase (1 RCT, n = 360, RR 4.94, 95% CI 2.50 to 9.78, [Analysis 2.12](#)).

e. Miscellaneous

In both studies, participants in both the haloperidol and aripiprazole groups were reported to have agitation (2 RCTs, n = 477, RR 1.22, 95% CI 0.46 to 3.22), and headache (2 RCTs, n = 477, RR 0.85, 95% CI 0.45 to 1.59) between days zero and one. Although not statistically significant, more people in the aripiprazole group reported pain at the injection site following the first injection (1 RCT, n = 360, RR 0.32, 95% CI 0.06 to 1.54, [Analysis 2.13](#)).

2.6 Leaving the study early

In both studies, more people left the study early in the haloperidol group than in the aripiprazole group, however the difference between groups was not statistically significant (2 RCTs, n = 480, RR 2.07, 95% CI 0.86 to 4.98). When reasons were given these data were also equivocal (e.g. 2 RCTs, n = 480, RR leaving because of adverse effects 0.97, 95% CI 0.17 to 5.59, [Analysis 2.14](#)).

3. COMPARISON 3. HALOPERIDOL versus OTHER ANTIPSYCHOTIC: b. CHLORPROMAZINE

3.1 Tranquillisation or asleep - asleep

One small study (n = 39) reported that more people allocated to chlorpromazine did experience sleep by two hours (RR 1.93, 95% CI 1.04 to 3.60, [Analysis 3.1](#)).

3.2 Repeated need for rapid tranquillisation

In the [Man 1973](#) study, all participants in the haloperidol group and all but one participant in the chlorpromazine study required more than one injection, with several in both groups requiring more than three (1 RCT, n = 30, 1.50, 95% CI 0.53 to 4.26, [Analysis 3.2](#)).

3.3 Global outcome - not marked improvement

For the outcome of "not marked improvement" there were no clear differences between groups (2 RCTs, n = 89, RR 0.79, 95% CI 0.61 to 1.02). More people in the chlorpromazine group were rated as showing "not any improvement" (2 RCTs, n = 89, RR 0.15, 95% CI 0.05 to 0.49, [Analysis 3.3](#)).

3.4 Adverse effects - specific

Reporting of adverse effects was sporadic. One person developed status epilepticus (1 RCT, n = 30, RR 0.33, 95% CI 0.01 to 7.58), and two people had hypotension (3 RCTs, n = 69, RR 0.51, 95% CI 0.10 to 2.60). In the [Man 1973](#) study (n = 30), there were no reported incidences of EPS or irritation at the injection site in either group.

One person in the chlorpromazine group had an elevated SGPT (serum glutamic pyruvic transaminase) level (1 RCT, n = 30, RR 0.33, 95% CI 0.01 to 7.81) and another in the haloperidol group developed mild leucopenia (1 RCT, n = 50, RR 3.00, 95% CI 0.13 to 70.30, [Analysis 3.4](#)).

3.5 Leaving the study early - for any reason

Few people left these studies early (5%) with little, but significant, difference between groups (4 RCTs, n = 153, RR 0.21, 95% CI 0.07 to 0.71, [Analysis 3.5](#)).

4. COMPARISON 4. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: c. DROPERIDOL

We found two relevant studies ($n = 255$) ([Calver 2015](#); [Resnick 1984](#)).

4.1 Tranquillisation or asleep

There was no statistical significant difference in terms of being not asleep at two hours between the compared groups (1 RCT, $n = 228$, RR 1.07, 95% CI 0.44 to 2.60, [Analysis 4.1](#)). People with haloperidol or droperidol experienced no different time to sleep (1 RCT, $n = 114$, MD 5.20, 95% CI -6.05 to 16.45, [Analysis 4.2](#)).

4.2 Repeated need for rapid tranquillisation

More people in the haloperidol group required more than one injection compared with those allocated to droperidol (2 RCTs, $n = 255$, RR 2.38, 95% CI 1.27 to 4.47); however, this was not statistically significant for the outcome of 'required more than three injections' (1 RCT, $n = 27$, RR 2.12, 95% CI 0.09 to 47.68, [Analysis 4.3](#)).

4.3 Global outcome - need for benzodiazepine

There was no statistical significant difference in terms of need for adjunctive benzodiazepine between the two comparison groups (1 RCT, $n = 228$, RR 0.31, 95% CI 0.07 to 1.44, [Analysis 4.4](#)).

4.4 Adverse effects

4.4.1 General

In the haloperidol group there was only one adverse effects up to two hours, compared with seven in the droperidol group (1 RCT, $n = 228$, RR 0.15, 95% CI 0.02 to 1.23), involving one person in the haloperidol group and six in the droperidol group (1 RCT, $n = 228$, [Analysis 4.5](#)).

4.4.2 Specific

a. Arousal

There were few differences between groups as for being over-sedated (1 RCT, $n = 228$, RR 0.36, 95% CI 0.01 to 8.68, [Analysis 4.6](#)).

b. Cardiovascular

One person in the haloperidol group experienced hypotension, compared with four in the droperidol group (1 RCT, $n = 228$, RR 0.27, 95% CI 0.03 to 2.36, [Analysis 4.7](#)).

c. Movement disorder

One person in the haloperidol group experienced dystonia, compared with one in the droperidol group (2 RCTs, $n = 255$, RR 0.86, 95% CI 0.11 to 6.56, [Analysis 4.8](#)).

d. Miscellaneous

There was no difference in terms of desaturation events between the two compared groups (1 RCT, $n = 228$, RR 0.36, 95% CI 0.01 to 8.68, [Analysis 4.9](#)).

5. COMPARISON 5. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: d. LOXAPINE

We found three relevant studies (total $n = 119$, [Fruensgaard 1977](#); [Paprocki 1977](#); [Tuason 1986](#)). [Fruensgaard 1977](#), unlike the other two studies also made sure that both groups of participants were given biperiden, an anticholinergic drug which offsets adverse effects such as movement disorders.

5.1 Tranquillisation or asleep – not asleep at about 12 hours

Loxapine does seem more sedating than haloperidol although the difference was not statistically significant (1 RCT, $n = 54$, RR 4.31, 95% CI 0.54 to 34.48, [Analysis 5.1](#)).

5.2 Specific behaviour

There were very few differences between groups for improvement in levels of agitation (1 RCT, $n = 30$, RR 1.17, 95% CI 0.51 to 2.66) or levels of aggression (1 RCT, $n = 30$, RR 1.10, 95% CI 0.69 to 1.76, [Analysis 5.2](#)).

5.3 Global outcome

Studies found no differences found between groups at any time point in regards to the number of people who were reported as having 'no change in global outcome' (1 RCT, $n = 56$, RR at 48 hours 0.93, 95% CI 0.14 to 6.15; 1 RCT, $n = 54$, RR at 72 hours 0.39, 95% CI 0.04 to 3.49 and 1 RCT, $n = 35$, RR up to 4 weeks 0.40, 95% CI 0.12 to 1.32, [Analysis 5.3](#)).

Although never statistically significant, more people in the loxapine group were sedated at 30 minutes (1 RCT, $n = 30$, RR 1.29, 95% CI 0.65 to 2.54), 60 minutes (1 RCT, $n = 30$, RR 3.50, 95% CI 0.86 to 14.18) and at 120 minutes (1 RCT, $n = 30$, RR 7.00, 95% CI 0.98 to 50.16) than in the haloperidol group ([Analysis 5.4](#)).

5.4 Mental state

One trial found no clear differences between the haloperidol and loxapine in average endpoint scores on the BPRS ($n = 52$, MD 6.10, 95% CI 4.48 to 7.72, [Analysis 5.6](#)).

5.5 Adverse effects

Over 20 different adverse effects are reported including anticholinergic disorders ([Analysis 5.8](#)), arousal levels ([Analysis 5.9](#)), cardiovascular effects ([Analysis 5.10](#)), movement disorders ([Analysis 5.11](#)) and several that were difficult to classify ([Analysis 5.12](#)), but these are probably best summarised by the general impression from [Analysis 5.7](#) where eight people in the haloperidol group and 10 people in the loxapine group experienced one or more drug-related adverse effects (1 RCT, $n = 30$, RR 0.80, 95% CI 0.44 to 1.45).

5.6 Leaving the study early

a. General reasons

At no point at anytime did statically more people leave one group than the other (e.g. 1 RCT, $n = 35$, RR leaving around 72 hours 0.63, 95% CI 0.21 to 1.85). The [Tuason 1986](#) study found no significant differences between the groups for the number of people who discontinued with the study by 10 days ($n = 54$, RR 1.18, 95% CI 0.67 to 2.07, [Analysis 5.13](#)).

b. Specific reasons

In [Paprocki 1977](#) they withdrew one person because of "toxicity" during the IM phase ($n = 35$, RR 2.84, 95% CI 0.12 to 65.34, [Analysis 5.14](#)).

6. COMPARISON 6. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: e. OLANZAPINE

We found six studies (total $n = 720$) that compared haloperidol with olanzapine (Baldacara 2011; Battaglia 2002; Breier 2002; Eli Lilly 2004; Kinon 2001d; Walther 2014).

6.1 Tranquillisation or asleep

By two hours significantly more people in the olanzapine group were asleep compared with those allocated haloperidol (1 RCT, $n = 257$, RR 1.16, 95% CI 1.02 to 1.32, Analysis 6.1).

6.2 Repeated need for tranquillisation

The need for additional injection results were statistically significant between the two groups in favour of olanzapine at 12 hours (1 RCT, $n = 60$, RR 3.29, 95% CI 1.67 to 6.47); however, the two groups results are similar at two hours (1 RCT, $n = 60$, RR 4.00, 95% CI 0.47 to 33.73) and at 24 hours (3 RCTs, $n = 392$, RR 1.06, 95% CI 0.75 to 1.51, Analysis 6.2).

6.3 Specific behaviour

a. Agitation

At two hours, a similar number of participants in both the haloperidol and olanzapine group were rated as having at least a 40% reduction according to the PANSS-EC (1 RCT, $n = 45$, RR 0.96, 95% CI 0.58 to 1.58). Also agitation was rated as an 'adverse effect' (1 RCT, $n = 49$, RR 1.04, 95% CI 0.07 to 15.73), and again reported by 21 days (1 RCT, $n = 100$, RR 1.08, 95% CI 0.33 to 3.51, Analysis 6.3).

By 30 minutes haloperidol and olanzapine showed similar findings at average change scores of ACES (15 minutes: 1 RCT, $n = 46$, MD 0.50, 95% CI -0.35 to 1.35; 30 minutes: 1 RCT, $n = 46$, MD 0.90, 95% CI -0.06 to 1.86) and PANSS-EC (15 minutes: 1 RCT, $n = 46$, MD 0.30, 95% CI -1.64 to 2.24; 30 minutes: 1 RCT, $n = 46$, MD 1.30, 95% CI -1.16 to 3.76, Analysis 6.4).

By about two hours, continuous measures of agitation showed similar findings. The ABS change score did favour olanzapine (1 RCT, $n = 85$, MD 2.70, 95% CI 0.38 to 5.02) and so did the ACES change score at one hour (1 RCT, $n = 46$, MD 1.20, 95% CI 0.06 to 2.34) and the Overt Agitation Severity Scale (OASS) endpoint score at one hour (1 RCT, $n = 60$, MD 2.00, 95% CI 1.18 to 2.82) and two hours (1 RCT, $n = 60$, MD 4.20, 95% CI 3.54 to 4.86). However, the others measurements did not show similar findings (for example, average change and endpoint combined scores PANSS-EC: 3 RCTs, $n = 332$, MD -0.31, 95% CI -1.49 to 0.86, Analysis 6.5).

At four hours, continuous measures of agitation showed similar findings. The average endpoint score on the OASS was statistically significant (1 RCT, $n = 60$, MD 0.20, 95% CI -0.05 to 0.45, Analysis 6.6).

When it came to 24 hours, continuous measures of agitation showed similar findings. For example, the average change score on the ABS (1 RCT, $n = 86$, MD 2.40, 95% CI 0.01 to 4.79) and the average endpoint score on the OASS at six hours (1 RCT, $n = 60$, MD 0.20, 95% CI 0.05 to 0.35) and 12 hours (1 RCT, $n = 60$, MD 0.90, 95% CI 0.82 to 0.98) were statistically significant, but on the PANSS-EC it was not (1 RCT, $n = 86$, MD 1.40, 95% CI -0.55 to 3.35, Analysis 6.7).

b. Aggression

The two groups showed no statistical significant difference on the average endpoint OASS score at one hour (1 RCT, $n = 60$, MD 0.90, 95% CI 0.13 to 1.67) and at two hours (1 RCT, $n = 60$, MD 0.50, 95% CI -0.41 to 1.41, Analysis 6.8).

At four hours, the average endpoint score on the OASS showed similar findings (1 RCT, $n = 60$, MD 0.10, 95% CI -0.11 to 0.31, Analysis 6.9).

When it came to 24 hours, the continuous measures of aggression at six hours showed similar findings at the average endpoint OASS score (1 RCT, $n = 60$, MD -0.10, 95% CI -0.26 to 0.06); however, at 12 hours the average endpoint OASS score resulted in favour of haloperidol (1 RCT, $n = 60$, MD -0.20, 95% CI -0.39 to -0.01, Analysis 6.10).

c. Hostility

This was recorded – again as an adverse effect – but there was not a clear difference between groups (1 RCT, $n = 49$, RR 0.35, 95% CI 0.01 to 8.12, Analysis 6.11)

6.4 Global outcome

a. General

There were no significant differences between the groups with regards to the number of people who required additional benzodiazepines during 24 hours (2 RCTs, $n = 343$, RR 1.05, 95% CI 0.63 to 1.74). This applied up to seven days (1 RCT, $n = 100$, RR 1.00, 95% CI 0.79 to 1.27), and use of additional restraint, seclusion or special observation (2 RCTs, $n = 160$, RR 1.49, 95% CI 0.70 to 3.17, Analysis 6.12). In one study ($n = 100$), there was a significant difference between the haloperidol and olanzapine group for time until the person discontinued the study – it was significantly earlier in the olanzapine group (MD -3.48, 95% CI -6.28 to -0.68). In the same study, the dose of adjunctive lorazepam needed was very similar (MD 0.34, 95% CI -0.14 to 0.82, Analysis 6.13).

People in the haloperidol and olanzapine group were rated as demonstrating similar levels of improvement in global functioning according to average endpoint scores using the CGI-I when rated at two hours (1 RCT, $n = 42$, MD -0.10, 95% CI -0.65 to 0.45, Analysis 6.14), and their average change scores at 24 hours (1 RCT, $n = 243$, MD 0.00, 95% CI -0.20 to 0.20). There was no difference by 24-72 hours according to average change scores using the CGI-S (1 RCT, $n = 86$, MD 0.00, 95% CI -0.24 to 0.24, Analysis 6.15). Kinon 2001d used a non-parametric test and reported that at 21 days, those in the olanzapine group did have a significantly greater change in the CGI-I score ($n = 100$, -0.38 ± 1.31 and -0.2 ± 1.44 , $P = 0.25$, F test 1.71 df). Further, one study demonstrated similar levels of improvement according to average change scores when using the CGI-I by 21 days (1 RCT, $n = 100$, MD 0.36, 95% CI -0.18 to 0.90, Analysis 6.15).

b. Specific

One study repeatedly recorded levels of arousal (alert to asleep) across several periods of time (one to 24 hours). There were no convincing differences between the two drugs (Analysis 6.16).

6.5 Service use

There was no difference between groups for the outcome of average time to discharge (about 13 days) (1 RCT, $n = 100$, MD -0.60, 95%

CI -1.85 to 0.65, [Analysis 6.28](#)). Time was also measured for the average number of patient-hours used. These data were skewed and therefore presented in a table ([Analysis 6.29](#)).

6.6 Mental state

Several adverse effects seem to be better classified in 'mental state'. Two different studies reported anxiety ($n = 49$, RR 0.35, 95% CI 0.01 to 8.12), or anxiety by 21 days ($n = 100$, RR 0.36, 95% CI 0.08 to 1.70). Delusions were also reported as an adverse effect (1 RCT, $n = 49$, RR 0.35, 95% CI 0.01 to 8.12), as was nervousness (1 RCT, $n = 100$, RR 2.17, 95% CI 0.70 to 6.74, [Analysis 6.30](#)). There were no significant differences between olanzapine and haloperidol with respect to improved mental state according to average change scores on the PANSS Total sub-scale (1 RCT, $n = 100$, MD 4.70, 95% CI -0.18 to 9.58, [Analysis 6.34](#)). There was also few differences according to average change scores on positive sub-scale of the BPRS at any time period (1 RCT, $n = 46$) nor on BPRS totals ([Analysis 6.31](#)).

6.7 Adverse effects

6.7.1 General

In three studies significantly more people in the haloperidol group experienced a drug-related adverse event (3 RCTs, $n = 209$, RR 1.38, 95% CI 1.09 to 1.76, [Analysis 6.35](#)).

6.7.2 Serious

There was no clear difference in risk of treatment emergent adverse events between groups (1 RCT, $n = 49$, RR 0.81, 95% CI 0.36 to 1.83). Only a few of these were attributed to the use of the drug. No serious adverse effects nor death were experienced by any participant who received haloperidol or olanzapine ([Analysis 6.36](#)).

6.7.3 Specific

a. Arousal level

Despite being reported in a way that may have underestimated the overall prevalence, complaints of insomnia and/or somnolence were not different between groups ([Analysis 6.38](#)).

b. Cardiovascular

Binary and continuous measures of any cardiovascular change did not show differences between groups (e.g. hypotension 2 RCTs, $n = 146$, RR 0.27, 95% CI 0.03 to 2.38, [Analysis 6.42](#); change in QT interval 1 RCT, $n = 257$, MD 1.80, 95% CI -3.81 to 7.41, [Analysis 6.43](#)).

c. Movement disorders

Studies reported different movement disorders in several different ways. There was the impression that more people who received haloperidol complained of these effects (e.g. EPS at 24 hours, 3 RCTs, $n = 403$, RR 8.35, 95% CI 2.27 to 30.63, [Analysis 6.45](#)).

This finding is supported by the continuous measure of SAS – again, this tends to favour olanzapine at 24 hours (1 RCT, $n = 242$, MD 1.31, 95% CI 0.56 to 2.06, [Analysis 6.46](#)). Other skewed SAS endpoint data are not so favourable, but these are presented only in tabular form ([Analysis 6.47](#)). This also applies to the akathisia endpoint measure (see [Analysis 6.47](#)) and, finally, change scores on the same scale (BAS) at 24 hours favoured olanzapine (1 RCT, $n = 241$, MD 0.28, 95% CI 0.09 to 0.47, [Analysis 6.46](#)).

d. Miscellaneous

Reports recorded several other adverse effects, none of which were different in the two groups. These included anticholinergic effects (increased salivation, 1 RCT, $n = 100$, RR 9.73, 95% CI 0.54 to 176.18, [Analysis 6.37](#)), gastric problems (e.g. abdominal pain 1 RCT, $n = 49$, RR 3.12, 95% CI 0.13 to 73.04, [Analysis 6.44](#)), and nose bleeds (1 RCT, $n = 49$, RR 3.12, 95% CI 0.13 to 73.04, [Analysis 6.49](#)).

No participants in either group were reported to have had a clinically relevant change in laboratory test results.

6.8 Leaving the study early

About 20% of each group left the study early but slightly fewer in the olanzapine group (2 RCTs, $n = 148$, RR 1.66, 95% CI 1.04 to 2.65). Power to investigate specific reasons was low but eight people in the haloperidol group, compared with one in the olanzapine group left the study early due to adverse events (1 RCT, $n = 100$, RR 8.67, 95% CI 1.13 to 66.75, [Analysis 6.50](#)).

7. COMPARISON 7. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: f. PERPHENAZINE

We found one relevant study ($n = 44$, [Fitzgerald 1969](#)).

7.1 Global outcome - no improvement

One person in the haloperidol group and two people in the perphenazine were reported to have shown no global improvement ($n = 44$, RR 0.46, 95% CI 0.04 to 4.68, [Analysis 7.1](#)).

7.2 Adverse effects

7.2.1 General

Ten people in the haloperidol group and seven in the perphenazine group experienced one or more adverse effect, however, this difference is not statistically significant ($n = 44$, RR 1.30, 95% CI 0.61 to 2.80, [Analysis 7.2](#)). No participants in either group developed clinically significant laboratory changes.

7.2.2 Specific

Six people in the haloperidol group and two people in the perphenazine group required antiparkinson medication, but again, this result was not statistically significant ($n = 44$, RR 2.74, 95% CI 0.62 to 12.12). One person in the perphenazine group developed a hypotensive episode compared with none in the haloperidol group ($n = 44$, RR 0.31, 95% CI 0.01 to 7.12).

Two people in the haloperidol group and one person in the perphenazine discontinued with the study because they developed EPS ($n = 44$, RR 1.83, 95% CI 0.18 to 18.70). Also one person was withdrawn from the haloperidol group due to drowsiness/tension and no therapeutic effect ($n = 44$, RR 2.75, 95% CI 0.12 to 64.04, [Analysis 7.3](#)).

7.3 Leaving the study early

We felt that we should add these specific reasons into the data on adverse effects ([Analysis 7.3](#)), but there were no differences between groups ([Analysis 7.4](#)).

8. COMPARISON 8. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: g. QUETIAPINE

We found one relevant study ($n = 80$, [Liang 2013](#)).

8.1 Specific behaviour - agitation

There was no difference between the two groups at the average change score of the PANSS-EC at 24 hours (MD 0.10, 95% CI -0.56 to 0.76, [Analysis 8.1](#)).

Continuous measures over 24 hours showed similar findings, from 72 hours (MD 0.20, 95% CI -0.87 to 1.27) to 28 days (MD 0.00, 95% CI -1.87 to 1.87, [Analysis 8.2](#)).

8.2 Mental state

No statistically significant difference was found at 28 days in average change score of PANSS (MD -0.40, 95% CI -7.04 to 6.24), PANSS positive sub-scale (MD 2.00, 95% CI -0.87 to 4.87), and PANSS negative sub-scale (MD -0.60, 95% CI -2.27 to 1.07, [Analysis 8.3](#)).

8.3 Adverse effects

8.3.1 General

People in the haloperidol group experienced more adverse events in 28 days when comparing with the quetiapine group (RR 1.81, 95% CI 1.19 to 2.77, [Analysis 8.4](#)).

8.3.2 Specific - movement disorder

Average endpoint SAS scores at 28 days were higher in the haloperidol group (MD 1.30, 95% CI 0.57 to 2.03, [Analysis 8.5](#)).

8.4 Leaving the study early

There were no differences between groups (RR 1.25, 95% CI 0.36 to 4.32, [Analysis 8.6](#)).

9. COMPARISON 9. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: h. RISPERIDONE

We found three relevant studies (n = 314), one of which - [Currier 2004](#) - randomised haloperidol versus risperidone but both groups were given lorazepam (2 mg).

9.1 Tranquillisation or asleep

One trial reported on the outcome of 'not asleep' across different time periods (30-120 minutes). In the company of lorazepam, haloperidol caused more people to be asleep than risperidone (e.g. awake at 30 minutes: 1 RCT, n = 162, RR 0.84, 95% CI 0.74 to 0.95, [Analysis 9.1](#)).

9.2 Specific behaviour

About 80% of participants in both groups had at least a 50% reduction in agitation scores according to the PANSS-EC (n = 124, RR 0.96, 95% CI 0.79 to 1.16). By 24 hours, one person in the risperidone group was withdrawn from the study early due to agitation, compared with no one in the haloperidol group (n = 124, RR 0.33, 95% CI 0.01 to 8.03, [Analysis 9.3](#)).

Continuous measures showed similar findings, confirming the absence of a difference between the two groups (1 RCT, n = 28, MD -0.40, 95% CI -5.22 to 4.42, [Analysis 9.4](#)).

9.3 Global outcome - need for benzodiazepine

Oddly, in the study that used lorazepam, the great majority of both groups were given additional benzodiazepine. This did not apply to the other trial where benzodiazepine was not part of the routine treatment. There was no difference between groups ([Analysis 9.7](#)).

9.4 Adverse effects

9.4.1 General

There were no statistically significant differences between the groups with regard to the number of people who reported adverse effects (e.g. one or more adverse effects during 24 hours, 2 RCTs, n = 286, RR 1.01, 95% CI 0.84 to 1.23, [Analysis 9.9](#)).

9.4.2 Specific

Again, levels of arousal suggested that haloperidol was more sedating than risperidone (e.g. sedated at 30 minutes 1 RCT, n = 162, RR 1.90, 95% CI 1.39 to 2.59, [Analysis 9.10](#)). Cardiovascular measures were not different except that risperidone seems to marginally increase pulse and haloperidol decrease it ([Analysis 9.12](#)).

Although not statistically significant, eight people the haloperidol group compared with five allocated risperidone suffered EPS (n = 124, RR 1.60, 95% CI 0.55 to 4.62). There was no difference in the levels of acute dystonia between the old and the new drug (about 2%, 2 RCTs, n = 286, RR 1.44, 95% CI 0.28 to 7.28, [Analysis 9.14](#)). Continuous measures of akathisia and movement disorders did, however, favour risperidone within 24 hours ([Analysis 9.15](#)), finding no difference at 96 hours ([Analysis 9.16](#)).

Other adverse effects were no different between groups ([Analysis 9.17](#)).

9.5 Leaving the study early

Again, few people left these studies early with no clear differences between groups ([Analysis 9.18](#)).

10. COMPARISON 10. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: i. THIOTHIXENE

We found two relevant studies (n = 74, [Kewala 1984](#); [Stotsky 1977](#)).

10.1 Repeated need for rapid tranquillisation

There were no significant differences between haloperidol and thiothixene with respect to the number of participants who required more than one injection (1 RCT, n = 30, RR 1.07, 95% CI 0.89 to 1.28) and more than three (1 RCT, n = 30, RR 2.50, 95% CI 0.57 to 10.93, [Analysis 10.1](#)).

10.2 Specific behaviours

Although agitation was rated as 'adverse effect', we have reported it here as it also reflects how we have done so with other comparisons. The study found no difference between those allocated to haloperidol and those given thiothixene (1 RCT, n = 44, RR 0.28, 95% CI 0.01 to 6.52, [Analysis 10.2](#)).

10.3 Global outcome

In one study six people who received haloperidol compared with two people who received thiothixene exhibited no response to the first injection (n = 44, RR 2.50, 95% CI 0.57 to 11.05). This one trial continued to rate people hourly for five hours and never found a clear difference ([Analysis 10.3](#)).

10.4 Adverse effects

10.4.1 General

There were no differences between the groups in terms of the number of participants who experienced one or more adverse effects (2 RCTs, $n = 74$, $RR\ 1.47$, 95% $CI\ 0.97$ to 2.22 , [Analysis 10.4](#)).

10.4.2 Specific

There were no clear differences between the two drugs in terms of movement disorders (e.g. ataxia, 1 RCT, $n = 44$, $RR\ 0.83$, 95% $CI\ 0.06$ to 12.49 , [Analysis 10.5](#)) and a series of other adverse effect outcomes ([Analysis 10.6](#)).

11. COMPARISON 11. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: j. ZIPRASIDONE

We found five relevant studies (total $n = 829$, [Baldacara 2011](#); [Brook 2000](#); [Li 2006](#); [Shu 2010](#); [Yin 2012](#)).

11.1 Repeated need for tranquillisation

There were no significant differences between haloperidol and ziprasidone in terms of additional drugs for tranquillisation up to two hours (1 RCT, $n = 60$, $RR\ 0.36$, 95% $CI\ 0.13$ to 1.01); however, the same outcome up to 24 hours revealed a significant difference in favour of ziprasidone (1 RCT, $n = 60$, $RR\ 1.64$, 95% $CI\ 1.07$ to 2.53 , [Analysis 11.1](#)).

11.2 Specific behaviour

11.2.1 Agitation

In one study ($n = 231$), there were no significant differences between groups in their average endpoint scores on the PANSS-EC across any time period from two hours (MD 0.06 , 95% $CI\ -1.13$ to 1.25), right on to 72 hours (MD 0.62 , 95% $CI\ -0.45$ to 1.69 , [Analysis 11.2](#)).

However, in another study ($n = 60$), there were significant differences between groups in their average endpoint scores on the OASS: in favour of haloperidol at one hour (MD -7.70 , 95% $CI\ -9.41$ to -5.99), and at two hours (MD -3.90 , 95% $CI\ -6.38$ to -1.42 , [Analysis 11.2](#)), in favour of ziprasidone at six hours (MD 0.20 , 95% $CI\ 0.05$ to 0.35), and at 12 hours (MD 0.90 , 95% $CI\ 0.82$ to 0.98 , [Analysis 11.4](#)).

11.2.2 Aggression

There were no significant differences between groups in their average endpoint scores on the OAS until six hours, when there was a significant difference in favour of ziprasidone (1 RCT, $n = 60$, MD 0.70 , 95% $CI\ 0.60$ to 0.80), confirmed at 12 hours (MD 0.20 , 95% $CI\ 0.07$ to 0.33 , [Analysis 11.8](#)).

11.3 Global outcome

In one study ($n = 132$), there were no significant differences between the groups with regard to the need for anxiolytic drugs ($RR\ 1.11$, 95% $CI\ 0.84$ to 1.48) or hypnotic medications ($RR\ 0.71$, 95% $CI\ 0.20$ to 2.50) up to seven days.

Also, there were no significant differences between groups in terms of additional restraint (1 RCT, $n = 60$, $RR\ 0.60$, 95% $CI\ 0.25$ to 1.44 , [Analysis 11.9](#)).

Using the CGI-S, this same study found people in the ziprasidone group had significantly improved average change scores for symptom severity when rated at three days ($n = 132$, MD 0.34 , 95%

$CI\ 0.13$ to 0.55) and seven days ($n = 132$, MD 0.51 , 95% $CI\ 0.07$ to 0.95 , [Analysis 11.10](#)).

11.4 Mental state

In one trial ($n = 60$), people were rated with BPRS at two hours (MD -4.25 , 95% $CI\ -10.16$ to 1.66 , [Analysis 11.11](#)), four hours (MD -4.60 , 95% $CI\ -9.60$ to 0.40 , [Analysis 11.12](#)), 24 hours (MD -4.00 , 95% $CI\ -9.09$ to 1.09 , [Analysis 11.13](#)), and 48 hours (MD -2.05 , 95% $CI\ -6.45$ to 2.35 , [Analysis 11.14](#)); the BPRS was also rated at 72 hours (3 RCTs, $n = 511$, MD -0.43 , 95% $CI\ -1.93$ to 1.07), and at seven days (1 RCT, $n = 132$, MD 2.93 , 95% $CI\ -0.81$ to 6.67 , [Analysis 11.14](#)).

Another trial ($n = 231$) rated people at 72 hours on the PANSS. There were no differences between the groups on the positive sub-scale (MD 1.11 , 95% $CI\ -0.45$ to 2.67), general pathology score (MD 0.00 , 95% $CI\ -2.27$ to 2.27) or total score (MD 2.45 , 95% $CI\ -2.19$ to 7.09). However, there was a significant difference in favour of haloperidol between groups on the negative sub-scale (MD -4.19 , 95% $CI\ -5.71$ to -2.67 , [Analysis 11.14](#)).

11.5 Adverse effects

For several of the studies reporting of adverse effects only occurred when $> 2\%$ or 5% or even 10% of people suffered the effect. This results in under-reporting.

11.5.1 General

No significant differences were found between the groups in terms of experiencing one or more adverse effects at 12 hours (1 RCT, $n = 60$, $RR\ 1.38$, 95% $CI\ 0.64$ to 2.93 , [Analysis 11.15](#)). At 72 hours, significantly more people in the haloperidol group had experienced one or more adverse effects (4 RCTs, $n = 799$, $RR\ 1.74$, 95% $CI\ 1.47$ to 2.06). However, by seven days, the difference between the groups was not statistically significant (1 RCT, $n = 132$, $RR\ 1.31$, 95% $CI\ 0.93$ to 1.83 , [Analysis 11.16](#)).

No one experienced an adverse effect that was thought to be 'severe' ([Analysis 11.17](#)).

11.5.2 Specific

a. Anticholinergic

More people with haloperidol experienced episodes of blurred vision (1 RCT, $n = 231$, $RR\ 3.97$, 95% $CI\ 1.15$ to 13.68), dry mouth (1 RCT, $n = 231$, $RR\ 2.97$, 95% $CI\ 1.22$ to 7.22) and hypersalivation (1 RCT, $n = 231$, $RR\ 2.55$, 95% $CI\ 1.11$ to 5.87); no significant differences were found concerning constipation (1 RCT, $n = 376$, $RR\ 0.40$, 95% $CI\ 0.08$ to 2.06 , [Analysis 11.18](#)).

b. Arousal

There were no differences between haloperidol and ziprasidone using both dichotomous assessments of excessive sedation (1 RCT, $n = 60$, $RR\ 1.00$, 95% $CI\ 0.22$ to 4.56), insomnia (1 RCT, $n = 376$, $RR\ 0.51$, 95% $CI\ 0.05$ to 5.53), lethargy (1 RCT, $n = 231$, $RR\ 1.26$, 95% $CI\ 0.67$ to 2.35) and somnolence (1 RCT, $n = 376$, $RR\ 1.16$, 95% $CI\ 0.43$ to 3.12 , [Analysis 11.19](#)), and continuous rating scales ([Analysis 11.20](#), [Analysis 11.21](#), [Analysis 11.22](#)).

b. Cardiovascular

Cardiac adverse effects seemed more common in the ziprasidone group (e.g. ECG abnormal by 72 hours 2 RCTs, $n = 607$, $RR\ 0.38$, 95%

CI 0.17 to 0.84; tachycardia by 72 hours 3 RCTs, $n = 739$, RR 0.49, 95% CI 0.23 to 1.04, [Analysis 11.23](#)).

c. Gastrointestinal

This also seems to apply to gastric upset – although not to a level that is statistically significant (e.g. vomiting between zero and three days 1 RCT, $n = 132$, RR 0.30, 95% CI 0.02 to 5.72; between zero and seven days 1 RCT, $n = 132$, RR 0.16, 95% CI 0.01 to 2.82, [Analysis 11.24](#)).

d. Haematological

Abnormal laboratory results were recorded for participants in both groups, but there were no significant differences across a range of measures (e.g. abnormal lab tests – by three days, 2 RCTs, $n = 508$, RR 0.90, 95% CI 0.69 to 1.17, [Analysis 11.25](#)).

e. Movement disorders

Over a wide range of measures, ziprasidone caused fewer movement disorders than haloperidol (e.g. akathisia by 72 hours 3 RCTs, $n = 739$, RR 2.32, 95% CI 1.34 to 4.01; acute dystonia by 72 hours 2 RCTs, $n = 508$, RR 10.26, 95% CI 1.67 to 63.17). Overall, more people in the haloperidol group were reported to suffer EPS during 12 hours (1 RCT, $n = 60$, RR 11.00, 95% CI 0.64 to 190.53) and at a significant level during 72 hours (2 RCTs, $n = 508$, RR 19.13, 95% CI 7.59 to 48.21) and at seven days (1 RCT, $n = 132$, RR 34.29, 95% CI 4.70 to 250.02, [Analysis 11.26](#)).

Continuous measures of movement disorders partially supported these findings ([Analysis 11.27](#)): a higher incidence of EPS and akathisia in the haloperidol group compared with the ziprasidone group according to the average score on the SAS at three days (2 RCTs, $n = 191$, MD 3.59, 95% CI 2.15 to 5.03), average change scores at seven days ($n = 131$, MD 6.10, 95% CI 3.91 to 8.29), and the average change score on the BAS at three days ($n = 131$, MD 0.47, 95% CI 0.18 to 0.76) and seven days ($n = 131$, MD 0.90, 95% CI 0.51 to 1.29).

11.6 Leaving the study early

There were no significant differences with regard to the number of participants who left the study early for any reasons during 72 hours (1 RCT, $n = 376$, RR 1.77, 95% CI 0.53 to 5.94) or within seven days (1 RCT, $n = 132$, RR 2.14, 95% CI 0.86 to 5.32, [Analysis 11.29](#)).

There were also no differences between the groups with regards to the number of participants who left studies early because of adverse effects (by 72 hours, 2 RCTs, $n = 508$, RR 2.40, 95% CI 0.45 to 12.75; seven days 1 RCT, $n = 132$, RR 0.54, 95% CI 0.06 to 4.65). There were no differences between number of people who withdrew consent (1 RCT, $n = 376$, RR 5.05, 95% CI 0.24 to 104.55) or left for any other reason (1 RCT, $n = 376$, RR 0.20, 95% CI 0.01 to 4.18) by 72 hours ([Analysis 11.30](#)).

12. COMPARISON 12. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: k. ZUCLOPENTHIXOL ACETATE

We found one study ($n = 70$, [Taymeeyapradit 2002](#)).

12.1 Repeated need for tranquillisation

More people in the haloperidol group required more than three injections ($n = 70$, RR 2.54, 95% CI 1.19 to 5.46, [Analysis 12.1](#)).

12.2 Adverse effects

One person in the haloperidol group suffered a reaction at the injection site, whereas no one allocated zuclopenthixol experienced this effect ($n = 70$, RR 3.55, 95% CI 0.15 to 84.14, [Analysis 12.3](#)). Seven people allocated to haloperidol and two in the zuclopenthixol acetate group developed a tremor during seven days ($n = 70$, RR 4.16, 95% CI 0.93 to 18.62, [Analysis 12.2](#)).

12.3 Leaving the study early

No participants in either group left the study early ([Analysis 12.4](#)).

13. HALOPERIDOL versus BENZODIAZEPINE: a. FLUNITRAZEPAM

We found one relevant study ($n = 28$, [Dorevitch 1999](#)).

13.1 Specific behaviours - aggression

The [Dorevitch 1999](#) study found no differences in the number of participants in the haloperidol and flunitrazepam groups who had a $\geq 50\%$ reduction on the OAS by 90 minutes ($n = 28$, RR 1.15, 95% CI 0.86 to 1.55, [Analysis 13.1](#)).

13.2 Global outcome - need for seclusion or restraint

No participants in either the haloperidol or flunitrazepam group required further seclusion or restraint ([Analysis 13.2](#)).

13.3 Adverse effects - EPS

No participants in either the haloperidol or flunitrazepam group have experienced extrapyramidal symptoms ([Analysis 13.3](#)).

14. COMPARISON 14. HALOPERIDOL versus BENZODIAZEPINE: b. LORAZEPAM

We found four relevant studies ($n = 207$, [Battaglia 1997](#); [Foster 1997](#); [Garza-Trevino 1989](#); [Salzman 1991](#)).

14.1 Tranquillisation or asleep - not asleep

There were no significant differences between the haloperidol and lorazepam groups with regard to the number of participants asleep at one hour (1 RCT, $n = 60$, RR 1.05, 95% CI 0.76 to 1.44). However, by three hours, significantly more people were asleep in the lorazepam group compared with the haloperidol group (1 RCT, $n = 66$, RR 1.93, 95% CI 1.14 to 3.27, [Analysis 14.1](#)).

14.2 Repeated need for rapid tranquillisation

There were no differences in numbers requiring more than one and more than three injections (1 RCT, $n = 66$, RR 1.14, 95% CI 0.91 to 1.43 and RR 1.11, 95% CI 0.50 to 2.45, respectively, [Analysis 14.2](#)).

14.3 Global outcome - no overall improvement

One trial ($n = 44$) found no difference between groups for the outcome of 'showing no improvement at 30 minutes' (RR 1.10, 95% CI 0.70 to 1.71). By 60 minutes this had not changed (1 RCT, $n = 44$, RR 1.64, 95% CI 0.54 to 5.03). All participants in both groups had reportedly exhibited global improvement beyond 60 minutes ([Analysis 14.3](#)).

14.4 Mental state - average endpoint score

No differences were found between the haloperidol and lorazepam groups for improvement in mental state as measured by BPRS at

one hour (1 RCT, $n = 37$, MD 3.26, 95% CI -4.13 to 10.65), and several times up to four hours ([Analysis 14.4](#)).

14.5 Adverse effects

14.5.1 General

Approximately a third of the participants in both groups experienced one or more drug-related adverse effect (1 RCT, $n = 66$, RR 1.13, 95% CI 0.60 to 2.10). No participants in either group were reported to have experienced serious adverse effects ([Analysis 14.6](#)).

14.5.2 Specific

There were no differences between those allocated haloperidol and lorazepam in terms of dry mouth (1 RCT, $n = 66$, RR 0.53, 95% CI 0.14 to 2.04, [Analysis 14.8](#)).

In terms of movement disorders and the emergence of ataxia (1 RCT, $n = 66$, RR 0.44, 95% CI 0.04 to 4.65), dystonia (1 RCT, $n = 66$, RR 3.54, 95% CI 0.42 to 30.03), speech disorder (1 RCT, $n = 66$, RR 1.77, 95% CI 0.35 to 9.01), rigidity (1 RCT, $n = 66$, RR 6.22, 95% CI 0.33 to 115.91), tremor (1 RCT, $n = 66$, RR 1.77, 95% CI 0.17 to 18.60) and the need for benztropine (1 RCT, $n = 66$, RR 1.99, 95% CI 0.68 to 5.83). One study ($n = 60$) reported that significantly more participants experienced EPS in the haloperidol group compared with the lorazepam group (RR 15.00, 95% CI 2.11 to 106.49, [Analysis 14.11](#)).

Some other adverse effects are reported. One study ($n = 60$) found no differences between the groups in the number of participants asleep at one hour (RR 0.89, 95% CI 0.40 to 1.99). Clonidine was used for hypertension but not significantly more in one group or the other (1 RCT, $n = 66$, RR 2.67, 95% CI 0.11 to 63.17). Dizziness was not more prevalent in either group (1 RCT, $n = 66$, RR 0.89, 95% CI 0.19 to 4.07, [Analysis 14.9](#)).

15. HALOPERIDOL versus BENZODIAZEPINE: c. MIDAZOLAM

We found one relevant study ($n = 84$, [Nobay 2004](#)).

15.1 Tranquillisation or asleep - average time

Significant differences favouring midazolam over haloperidol were reported for the average time to sedation, however these data were skewed and are presented in tabular form only ([Analysis 15.1](#)).

15.2 Global outcome

There were no clear differences between the groups for the number of people requiring a rescue drug ($n = 84$, RR 1.14, 95% CI 0.46 to 2.87, [Analysis 15.2](#)).

15.3 Adverse effects

Two participants were reported to have experienced adverse effects. These were both in the haloperidol group with one person experiencing hypotension ($n = 84$, RR 3.00, 95% CI 0.13 to 71.61) and the other, apnoea ($n = 84$, RR 3.00, 95% CI 0.13 to 71.61, [Analysis 15.3](#)).

16. HALOPERIDOL versus COMBINATIONS: a. HALOPERIDOL + LEVOMEPRMAZINE

We found one study ($n = 19$, [Higashima 2004](#)).

16.1. Global outcome

There was no clear difference for the outcome of 'not improved' (1 RCT, $n = 19$, RR 8.18, 95% CI 0.50 to 133.66, [Analysis 16.1](#)).

17. HALOPERIDOL versus COMBINATIONS: b. HALOPERIDOL + LORAZEPAM

We found two studies that compared haloperidol alone with a combination of haloperidol and lorazepam ($n = 113$, [Battaglia 1997](#); [Garza-Trevino 1989](#)).

17.1 Tranquillisation or asleep

Significantly more people in the combination group were asleep at three hours than those in the haloperidol alone group (1 RCT, $n = 67$, RR 1.83, 95% CI 1.11 to 3.02, [Analysis 17.1](#)).

17.2 Repeated need for rapid tranquillisation

Fewer people in the combination group required more than one (1 RCT, $n = 67$, RR 1.05, 95% CI 0.87 to 1.27) and more than three additional injections within 24 hours than in the haloperidol alone group (1 RCT, $n = 67$, RR 3.05, 95% CI 0.92 to 10.10, [Analysis 17.2](#)).

17.3 Global outcome - no overall improvement

Significantly fewer people in the combination group showed no overall improvement by 30 minutes (1 RCT, $n = 45$, RR 2.67, 95% CI 1.25 to 5.68). By 60 minutes, although six people in the haloperidol alone group continued to show no overall improvement compared with none in the combination group; the difference between the groups was not statistically significant (1 RCT, $n = 45$, RR 14.77, 95% CI 0.88 to 247.54). No participants in either group showed 'no overall improvement' after 60 minutes. All participants in both groups demonstrated overall improvement ([Analysis 17.3](#)).

17.4 Adverse effects

17.4.1 General

One study ($n = 67$) found no difference for numbers experiencing one or more adverse effects (RR 1.16, 95% CI 0.62 to 2.18, [Analysis 17.4](#)).

17.4.2 Specific

In terms of a series of movement disorders, one study ($n = 67$) found no statistically significant differences between the groups (e.g. dystonia RR 8.25, 95% CI 0.46 to 147.45; required antiparkinson medication RR 2.74, 95% CI 0.81 to 9.25, [Analysis 17.7](#)).

The anticholinergic effect of dry mouth was not more common in the haloperidol alone group (1 RCT, $n = 67$, RR 0.91, 95% CI 0.20 to 4.21, [Analysis 17.5](#)). This also applies to cardiovascular effects such as hypertension (1 RCT, $n = 67$, RR 0.91, 95% CI 0.06 to 14.02) and dizziness (1 RCT, $n = 67$, RR 1.37, 95% CI 0.24 to 7.69, [Analysis 17.6](#)).

18. HALOPERIDOL versus COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM

We found one study ($n = 60$) comparing haloperidol versus haloperidol plus midazolam ([Baldacara 2011](#)).

18.1 Repeated need for tranquillisation

There were significant differences between groups in terms of need for additional drugs for tranquillisation at two hours in favour of

haloperidol (RR 0.27, 95% CI 0.10 to 0.71); however, this was not confirmed at 12 hours (RR 0.88, 95% CI 0.69 to 1.13, [Analysis 18.1](#)).

18.2 Specific behaviour

18.2.1 Agitation

Average endpoint scores on OASS showed a significant difference in favour of haloperidol at one hour (MD -8.50, 95% CI -9.93 to -7.07), at two hours (MD -6.70, 95% CI -7.46 to -5.94, [Analysis 18.2](#)), at four hours (MD -5.50, 95% CI -6.48 to -4.52, [Analysis 18.3](#)), at six hours (MD -3.70, 95% CI -4.56 to -2.84), and at 12 hours (MD -11.20, 95% CI -12.24 to -10.16, [Analysis 18.4](#)).

18.2.2 Aggression

Continuous measurement on OAS at different rating time points showed a significant advantage of haloperidol over the combination group ([Analysis 18.5](#)), with the exception of the rating at four hours (MD -0.10, 95% CI -0.68 to 0.48, [Analysis 18.6](#)) and at six hours (MD 0.40, 95% CI 0.30 to 0.50, [Analysis 18.7](#)).

18.3 Global outcomes

Six people in the haloperidol group needed additional restraint, seclusion or special observation compared with 21 people in the combination group (1 RCT, n = 60, RR 0.29, 95% CI 0.13 to 0.61, [Analysis 18.8](#)).

18.4 Adverse effects

18.4.1 General

There were no significant differences between the compared groups in terms of number of people experiencing one or more adverse effect (RR 0.73, 95% CI 0.41 to 1.32, [Analysis 18.9](#)).

18.4.2 Specific

a. Arousal

More people in the combination group experienced excessive sedation by 12 hours (RR 0.25, 95% CI 0.08 to 0.80, [Analysis 18.11](#)). Continuous measurement on the Ramsay Sedation Scale (RSS) at various time points however failed to show differences between the compared groups ([Analysis 18.12](#); [Analysis 18.13](#), [Analysis 18.14](#)).

b. Cardiovascular

Five people in the combination group experienced hypotension episodes compared to no people in the haloperidol group (RR 0.09, 95% CI 0.01 to 1.57, [Analysis 18.15](#)).

c. Movement disorders

There were no significant differences in terms of EPS by 12 hours (RR 1.67, 95% CI 0.44 to 6.36, [Analysis 18.16](#)).

19. HALOPERIDOL versus COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE

We found two relevant studies (n = 376) ([Baldacara 2011](#); [Huf 2007](#)).

19.1 Tranquillisation or asleep

More people in the haloperidol group compared with those in the combination group were not tranquil or asleep by 20 minutes (n = 316, RR 1.60, 95% CI 1.18 to 2.16), 40 minutes (n = 316, RR 1.26, 95% CI 0.83 to 1.91), 60 minutes (n = 316, RR 1.42, 95% CI 0.85 to 2.37) and 120 minutes (n = 316, RR 1.32, 95% CI 0.68 to 2.56). The difference

between the groups was only statistically significant at 20 minutes ([Analysis 19.1](#); [Analysis 19.2](#)).

19.2 Repeated need for tranquillisation

No more people in the haloperidol alone group required repeat rapid tranquillisation by two hours (2 RCTs, n = 376, RR 0.78, 95% CI 0.43 to 1.41) nor by 12 hours (1 RCT, n = 60, RR 1.10, 95% CI 0.81 to 1.49, [Analysis 19.3](#)).

19.3 Specific behaviours

19.3.1 Agitation

When rated on OASS, there was a significant advantage of haloperidol at one hour (1 RCT, n = 60, MD -24.50, 95% CI -27.32 to -21.68) and two hours (1 RCT, n = 60, MD -9.40, 95% CI -10.39 to -8.41, [Analysis 19.4](#)), and of combination at four hours (1 RCT, n = 60, MD 3.80, 95% CI 3.27 to 4.33, [Analysis 19.5](#)), at six hours (1 RCT, n = 60, MD 2.60, 95% CI 2.13 to 3.07) and at 12 hours (1 RCT, n = 60, MD 0.80, 95% CI 0.55 to 1.05, [Analysis 19.6](#)).

19.3.2 Aggression

While at continuous rating on OAS there was a significant advantage of haloperidol at one hour (1 RCT, n = 60, MD -4.50, 95% CI -6.28 to -2.72, [Analysis 19.7](#)), subsequent rating resulted significantly in favour of combination at four hours (1 RCT, n = 60, MD 0.60, 95% CI 0.49 to 0.71, [Analysis 19.8](#)), at six hours (1 RCT, n = 60, MD 1.10, 95% CI 0.91 to 1.29), and at 12 hours (1 RCT, n = 60, MD 1.80, 95% CI 1.67 to 1.93, [Analysis 19.9](#)).

No more people in the haloperidol alone group exhibited a further aggressive outburst within 24 hours (n = 316, RR 1.06, 95% CI 0.68 to 1.65, [Analysis 19.10](#)).

19.4 Global outcomes

More people in the haloperidol alone group had to be restrained by 120 minutes, however the difference between groups was not statistically significant (n = 316, RR 1.21, 95% CI 0.84 to 1.76, [Analysis 19.11](#)). The number of people who refused oral medication was roughly the same in both the haloperidol alone and combination group (n = 316, RR 0.99, 95% CI 0.61 to 1.60). Significantly more people in the haloperidol alone group required extra visits from the doctor (n = 316, RR 1.50, 95% CI 1.05 to 2.14, [Analysis 19.12](#)).

19.5 Adverse effects

19.5.1 General

Significantly more people in the haloperidol alone group than the combination group experienced one or more adverse effects (2 RCTs, n = 376, RR 2.01, 95% CI 1.07 to 3.80, [Analysis 19.13](#)). There were no differences between the compared group in terms of serious adverse events ([Analysis 19.14](#)).

19.5.2 Specific

a. Arousal

No significant differences were found between the compared groups ([Analysis 19.15](#); [Analysis 19.16](#); [Analysis 19.17](#); [Analysis 19.18](#)).

b. Cardiovascular

Three people in the combination group experienced hypotension episodes compared to no people in the haloperidol group (1 RCT, $n = 60$, RR 0.14, 95% CI 0.01 to 2.65, [Analysis 19.19](#)).

c. Movement disorders

Nine people in the haloperidol alone group experienced acute dystonia compared with no one in the combination group ($n = 316$, RR 19.48, 95% CI 1.14 to 331.92); there were no differences in terms of number of EPS episodes by 12 hours between the compared groups (1 RCT, $n = 60$, RR 1.00, 95% CI 0.32 to 3.10, [Analysis 19.20](#)).

19.6 Service outcomes

Less people in the haloperidol alone group were not discharged by 14 days, but this was not a statistically significant difference between groups ($n = 316$, RR 0.87, 95% CI 0.72 to 1.05, [Analysis 19.21](#)).

19.7 Leaving the study early

Seventeen people from each group left the study early, therefore there were no statistical differences between groups ([Analysis 19.22](#)). When specific reasons were cited, again, no differences were clearly shown (e.g. withdrew consent $n = 316$, RR 1.03, 95% CI 0.06 to 16.25, [Analysis 19.22](#)).

20. HALOPERIDOL versus COMBINATIONS: e. HALOPERIDOL + RISPERIDONE

We found one study ($n = 100$) ([Liu 2012](#)).

20.1 Specific behaviour - agitation

There was a statistically significant difference between the compared groups on the average scores of PANSS-EC at 24 hours in favour of the comparison group (MD 3.82, 95% CI 1.35 to 6.29, [Analysis 20.1](#)); rating at three days, five days, one week and two weeks failed to show a significant difference ([Analysis 20.2](#)).

20.2 Global outcomes

More people in the combination group experienced no improvement at 72 hours (RR 0.38, 95% CI 0.11 to 1.33), at five days (RR 0.25, 95% CI 0.03 to 2.16), at one week (RR 0.67, 95% CI 0.12 to 3.82), at two weeks (RR 0.33, 95% CI 0.01 to 7.99) compared to people in the haloperidol group; however, differences were not significant ([Analysis 20.3](#)).

20.3 Adverse effects

a. Arousal

There were no significant differences in terms of insomnia (RR 1.20, 95% CI 0.39 to 3.68) and somnolence (RR 0.92, 95% CI 0.45 to 1.88, [Analysis 20.4](#)).

b. Movement disorders

There were no significant differences between the compared groups in terms of tremor (RR 0.94, 95% CI 0.54 to 1.65) and akathisia (RR 0.88, 95% CI 0.48 to 1.60) during 14 days ([Analysis 20.5](#)).

c. Miscellaneous

No significant differences were found when comparing the two groups concerning dry mouth (RR 0.89, 95% CI 0.37 to 2.12), constipation (RR 1.50, 95% CI 0.45 to 4.99), and increased salivation (RR 2.00, 95% CI 0.19 to 21.36) during 14 days ([Analysis 20.6](#)).

21. HALOPERIDOL versus COMBINATIONS: f. OLANZAPINE + MAGNESIUM VALPROATE

We found one study ($n = 64$) that compared haloperidol versus olanzapine plus magnesium valproate ([Shi 2010](#)).

21.1 Specific behaviour - agitation

There were no significant differences on average endpoint scores on PANSS-EC at 72 hours (MD -0.04, 95% CI -1.33 to 1.25); however, rating on PANSS-EC showed significant differences at seven days (MD 1.61, 95% CI 0.15 to 3.07) and 14 days (MD 1.82, 95% CI 0.61 to 3.03) in favour of combination group ([Analysis 21.1](#)).

21.2 Global outcomes - no improvement

The same number of people experienced no improvement (RR 1.06, 95% CI 0.38 to 2.95, [Analysis 21.2](#)).

21.3 Mental state

There were no significant differences on average endpoint PANSS scores at 72 hours (MD -1.58, 95% CI -7.79 to 4.63), at seven days (MD 1.94, 95% CI -2.22 to 6.10), and at 14 days (MD 2.42, 95% CI -0.86 to 5.70, [Analysis 21.3](#)).

21.4 Adverse effects

Few people on the combination group experienced one or more adverse effects (RR 3.06, 95% CI 1.62 to 5.79) compared to haloperidol alone group ([Analysis 21.4](#)).

22. HALOPERIDOL versus COMBINATIONS: g. QUETIAPINE + MAGNESIUM VALPROATE

We found one study that compared haloperidol with a combination of quetiapine + magnesium valproate ($n = 60$, [Guo 2007](#)).

22.1 Specific behaviour - agitation

Regarding the binary measure of agitation (less than 25% decrease in PANSS-EC score) there was no difference between groups ($n = 60$, RR 1.17, 95% CI 0.44 to 3.06, [Analysis 22.1](#)).

When endpoint scores for agitation were recorded at three, seven and 14 days no differences were apparent ($n = 60$, MD 14 days 1.24, 95% CI -0.49 to 2.97, [Analysis 22.2](#)).

23. HALOPERIDOL versus COMBINATIONS: h. RISPERIDONE + CLONAZEPAM

We found two studies ($n = 305$) that compared haloperidol versus risperidone plus clonazepam ([Fang 2012](#); [Liu 2012](#)).

23.1 Tranquillisation or asleep

There were no significant differences in the number of people not asleep at two hours (1 RCT, $n = 205$, RR 1.07, 95% CI 0.82 to 1.39) and at four hours (1 RCT, $n = 205$, RR 0.82, 95% CI 0.66 to 1.03, [Analysis 23.1](#)).

23.2 Specific behaviour - agitation

Continuous measurement on average PANSS-EC scores showed no significant differences between the compared groups at two hours (1 RCT, $n = 108$, MD -0.50, 95% CI -3.80 to 2.80, [Analysis 23.2](#)), at four hours (1 RCT, $n = 126$, MD -2.40, 95% CI -5.20 to 0.40, [Analysis 23.3](#)), at 24 hours (2 RCTs, $n = 305$, MD 0.71, 95% CI -0.56 to 1.98, [Analysis 23.4](#)), at three days (2 RCTs, $n = 305$, MD -0.41, 95% CI -1.66 to 0.83), at five days (2 RCTs, $n = 305$, MD 0.14, 95% CI -1.24 to 1.52), and at two weeks (1 RCT, $n = 100$, MD -0.88, 95% CI -2.34 to 0.58); the only exception was at week one in favour of haloperidol (1 RCT, $n = 100$, MD -2.56, 95% CI -4.57 to -0.55, [Analysis 23.5](#)).

23.3 Global outcomes

More people on the combination group experienced no improvement at 72 hours (RR 0.60, 95% CI 0.15 to 2.38), at five days (RR 0.25, 95% CI 0.03 to 2.16), at one week (RR 0.20, 95% CI 0.01 to 4.06), and at two weeks (RR 0.50, 95% CI 0.10 to 2.61) compared with haloperidol group ([Analysis 23.6](#)).

23.4 Mental state

There were no significant differences on average change score on PANSS at five days (1 RCT, $n = 205$, MD -3.50, 95% CI -7.07 to 0.07, [Analysis 23.7](#)).

23.5 Adverse effects

23.5.1 General

In the combination group there were fewer adverse events at a significant level (1 RCT, $n = 205$, RR 1.72, 95% CI 1.29 to 2.29, [Analysis 23.8](#)).

23.5.2 Specific

a. Arousal

There were no significant differences concerning arousal adverse events (e.g. insomnia during five days 1 RCT, $n = 205$, RR 1.54, 95% CI 0.45 to 5.31; insomnia during 14 days 1 RCT, $n = 100$, RR 0.50, 95% CI 0.20 to 1.23; somnolence during 14 days 1 RCT, $n = 100$, RR 2.75, 95% CI 0.94 to 8.06) between the compared groups ([Analysis 23.9](#)).

b. Cardiovascular

There were no significant differences between the groups concerning tachycardia during five days (1 RCT, $n = 205$, RR 1.29, 95% CI 0.36 to 4.66, [Analysis 23.10](#)).

c. Movement disorders

People in the haloperidol group experienced more movement disorder-related adverse events (e.g. akathisia during five days 1 RCT, $n = 205$, RR 1.63, 95% CI 1.12 to 2.38; EPS during five days 1 RCT, $n = 205$, RR 2.22, 95% CI 1.52 to 3.23; tremor during 14 days 1 RCT, $n = 100$, RR 3.20, 95% CI 1.27 to 8.07, [Analysis 23.11](#)).

d. Miscellaneous

There were no significant differences in terms of anticholinergic adverse events ([Analysis 23.12](#)).

24. HALOPERIDOL versus COMBINATIONS: i. ZIPRASIDONE + CLONAZEPAM

We found 1 study ($n = 76$) that compared haloperidol versus ziprasidone plus clonazepam ([Zhou 2014](#)).

24.1 Specific behaviour - agitation

There were no significant differences on the average endpoint PANSS-EC scores at 12 hours (MD -0.28, 95% CI -0.80 to 1.24, [Analysis 24.1](#)), at 48 hours (MD -1.54, 95% CI -3.26 to 0.18), at three days (MD -0.42, 95% CI 1.63 to 0.79), at five days (MD -0.22, 95% CI -1.78 to 1.34), and at seven days (MD -0.19, 95% CI -1.57 to 1.19, [Analysis 24.2](#)).

24.2 Global outcomes

There were no significant differences on average endpoint CGI-S scores at seven days (MD 0.00, 95% CI -0.26 to 0.26, [Analysis 24.3](#)).

24.3 Mental state

There were no significant differences on average endpoint PANSS total scores at seven days (MD 1.46, 95% CI -4.58 to 7.50) and sub-scales ([Analysis 24.4](#)).

24.4 Leaving the study early

Three people in the haloperidol group left the study early compared to two people in the combination group (RR 1.50, 95% CI 0.27 to 8.48, [Analysis 24.5](#)).

DISCUSSION

Summary of main results

1. COMPARISON 1. HALOPERIDOL versus PLACEBO

It can be seen from the [Summary of findings for the main comparison](#) that, despite five trials providing data (total $n = 700$), the overall quality of evidence was not high. No study discussed economic outcomes.

1.1 Tranquillisation or asleep, repeated need for tranquillisation, agitation

When compared with placebo, significantly more people in the haloperidol group were asleep at two hours and demonstrated at least a 40% reduction in Positive and Negative Syndrome Scale Excited Component (PANSS-EC) agitation score ([Bristol Myers 2004a](#); [Bristol-Myers 2005b](#)). With respect to evidence for rapid tranquillisation, the evidence was downgraded to 'very low' for 'tranquil or asleep' because there was no measure at 30 minutes, therefore, two hours was the closest time point. We think this is quite distant from the emergency situation and most data for this outcome came from [Battaglia 2002](#) study, in which there is a serious risk of bias because the method of randomisation was not described, allocation concealment was not stated, there were incomplete outcome data and the trial was sponsored by the drug company, which could possibly impact on publication bias (please see [Summary of findings for the main comparison](#)).

On the continuous measures of agitation, haloperidol seems to demonstrate a discernable effect. For example, people in the haloperidol group were reported to have experienced a significantly greater reduction in agitation than those in the placebo group at two hours (3 RCTs, $n = 474$, MD -4.48, 95% CI -5.78 to -3.18, [Analysis 1.4](#)). A four-point reduction may be of clinical meaning, but we could not find this clearly highlighted.

1.2 Global outcome, mental state

Fewer people in the placebo group exhibited 'marked improvement' than those who received haloperidol ([Analysis 1.6](#)), and four studies found that more people in the placebo group required benzodiazepines ([Analysis 1.7](#)). All data point to the real effect of use of haloperidol in these difficult and dangerous situations. Mental state data support this impression.

1.3 Adverse effects

More people in the haloperidol group experienced one or more adverse reactions than those in the placebo group ([Analysis 1.12](#)). [Bristol-Myers 2005b](#) reported that of all adverse effects reported, only one was rated as 'serious' - this was related to a participant in the placebo group.

More people in the haloperidol group than the placebo group were reported to be "over sedated". Haloperidol did seem to not cause more cardiac problems than placebo but, unsurprisingly, movement disorders were more prevalent. Often, however, and on a range of outcomes, the differences were not statistically significant. However, significantly more people in the haloperidol group did require antiparkinson drugs and experienced Extrapyramidal Side Effects (EPS) ([Analysis 1.17](#)).

1.4 Leaving the study early

It is problematic how to interpret data on leaving the study as, in some of these trials, people may not have been fully free to leave. Only 27 of 335 people left early - with no differences between groups.

2. COMPARISON 2. HALOPERIDOL versus OTHER ANTIPSYCHOTIC: a. ARIPIRAZOLE

This comparison contains the larger trials (two RCTs, total n = 599), but, as can be seen from the [Summary of findings 2](#), the selected outcomes are absent or of low or very low quality.

2.1 Repeated need for rapid tranquillisation

Considering how difficult repeated injections are, that use of haloperidol necessitates less is important. The quality of these data is, however, low. We do not have any direct data on tranquillisation, but there is the implication from these data that haloperidol either sedates or tranquillises more effectively than aripiprazole.

2.2 Specific behaviours - agitation

There was no difference between groups at about two hours with about half of each group having a considerable reduction in agitation scores. We graded these data as of very low quality ([Summary of findings 2](#)). Two hours, is, however, a considerable time after an acute aggressive or agitated event and data at an earlier state would be helpful.

2.3 Global outcomes

All global outcomes – again all around two hours – show no difference between the two drugs (requiring additional benzodiazepines; average change scores on the Clinical Global Impression - Severity (CGI-S) rating scale; average endpoint scores on the Clinical Global Impression - Improvement (CGI-I) rating scale; average change scores on the CGI-I rating scale in the schizophrenia subgroup). This is reassuring that aripiprazole does have some

effect but its value in the very acute stage of management is not supported by these data. The few mental state data concur.

2.4 Adverse effects

For some adverse effects haloperidol seems to come off worse than aripiprazole although there was no difference between groups for 'adverse effects within 24 hours'. One person in the haloperidol group developed severe dystonia and one in the aripiprazole group had a tonic-clonic seizure.

Evidence was not convincing that there were any differences when it came to cardiac or gastrointestinal effects. Movement disorders, however, all tended to be more common in the group allocated to haloperidol and risk of use of antiparkinson medication was increased if a person was allocated to haloperidol. Although this is not often the problem that is most paramount within an emergency, these effects result in an already difficult situation being even more unpleasant. Akathisia (RR nearly 3), for example, can be mistaken for the original agitation and result in administration of yet more haloperidol. These common effects (~1/3 needed additional antiparkinson medication) are a considerable limitation for the value of haloperidol.

2.5 Leaving the study early

There were no significant differences in the number of people leaving early and, again, the meaning of these data is unclear as we are not sure how at liberty people were to leave the studies by the end of the trial.

Economic outcomes were not measured. Aripiprazole is considerably more expensive than haloperidol.

3. COMPARISON 3. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: b. CHLORPROMAZINE

Although we found four relevant studies, all were small (total n = 153) and we could not rate overall findings more than of very low grade ([Summary of findings 3](#)), and no important outcome had all trials contributing.

3.1 Tranquillisation or asleep, repeated need for rapid tranquillisation, not marked improvement

One small study (n = 39) of poor quality reported that more people allocated to chlorpromazine did experience sleep by two hours ([Analysis 3.1](#)). This does seem to fit with clinical experience.

Another study reported the number of people requiring more than one injection and, despite the suggestion of more sedation in the chlorpromazine group, there was no clear indication that either group avoided more or fewer additional injections – but, again the trial was small and of very limited quality ([Analysis 3.2](#)).

Although there was no clear difference for the outcome of "not marked improvement", more people in the chlorpromazine group were rated as showing "not any improvement" ([Analysis 3.3](#)). We are unclear if this has meaning or is a chance finding.

3.2 Adverse effects

Reporting of adverse effects was sporadic and therefore, not very helpful. From these data it is not clear that either drug has clear advantage over the other for allergic, cardiovascular, or movement disorders ([Analysis 3.4](#)).

3.3 Leaving the study early - for any reason

This was the only outcome where all trials contributed. Few people left these studies early (5%). This could be a function of the coercive aspect of the treatment. Perhaps people were not fully free to leave; significantly fewer people left the haloperidol group ([Analysis 3.5](#)).

4. COMPARISON 4. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: c. DROPERIDOL

Droperidol has been taken off the market because of “commercial reasons” ([Khokhar 2016](#)). However, there remains interest in its use ([Isbister 2010](#)). We found two relevant studies ($n = 255$) and the outcomes were of moderate grade (higher than others in this review) ([Summary of findings 4](#)).

4.1 Tranquillisation or asleep, repeated need for rapid tranquillisation

The same amount of people in the haloperidol and droperidol groups did experience sleep by two hours ([Analysis 4.1](#)), with no significant differences concerning the time to get asleep ([Analysis 4.2](#)).

Droperidol did seem more effective - more people in the haloperidol group required more than one injection, and this held true for the outcome of ‘required more than three injections’ ([Analysis 4.3](#)).

4.2 Global outcomes

To note, more people in the droperidol group required more benzodiazepines up to two hours ([Analysis 4.4](#)).

4.3 Adverse effects - specific

Droperidol could be equally or more toxic than haloperidol in terms of total adverse events ([Analysis 4.5](#)) and specific adverse events ([Analysis 4.6](#); [Analysis 4.7](#); [Analysis 4.8](#); [Analysis 4.9](#)).

Further studies are needed; much better evaluation is justified but in times of austerity in health care, it could be a clinically useful option.

5. COMPARISON 5. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: d. LOXAPINE

We found three relevant studies (total $n = 119$), one of which offset adverse effects using biperiden, but that we could only rate the outcomes as being of low or very low grade ([Summary of findings 5](#)). Loxapine is widely available in several European countries, Australasia and the USA. It is a shame that we do not have better data on this interesting compound that some consider to be inexpensive, but with ‘atypical’ properties ([Glazer 1999](#)).

5.1 Tranquillisation or asleep, agitation, aggression

Loxapine did seem more sedating than haloperidol although the one study lacked power ($n = 54$, [Analysis 5.1](#)). There were no differences between groups for improvement in levels of agitation or levels of aggression but, again, these outcomes were reported by only one very small study ($n = 30$, [Analysis 5.2](#)).

5.2 Global outcomes, mental state

There are several ways these trials reported global outcomes and measured mental state, but never any real suggestion of an effect favouring one or other drug.

5.3 Adverse effects

Even with the use of biperiden in one study, it might have been expected to see some differences between drugs. None were apparent in these trials.

6. COMPARISON 6. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: e. OLANZAPINE

We found six studies (total $n = 720$) but no pre-selected outcome of importance contained data from more than 400 people and all important outcomes we had to rate as being of very low grade ([Summary of findings table 6](#)).

6.1 Tranquillisation or asleep, need for more tranquillisation, specific behaviours

Olanzapine does seem more sedating than haloperidol ([Analysis 6.1](#)). In terms of requiring additional injections, a slight advantage in favours of olanzapine was demonstrated ([Analysis 6.2](#)).

When analysing agitation rating scales, a slight tendency in favour of olanzapine was demonstrated ([Analysis 6.3](#); [Analysis 6.4](#), [Analysis 6.5](#); [Analysis 6.6](#); [Analysis 6.7](#)).

In terms of both aggression and hostility, haloperidol and olanzapine showed no difference ([Analysis 6.8](#); [Analysis 6.9](#); [Analysis 6.10](#); [Analysis 6.11](#)).

6.2 Global outcome, mental state

Most global outcomes demonstrated no clear difference between the compounds. For example, use of additional restraint, seclusion or special observation was comparable ([Analysis 6.12](#)). Olanzapine seems as effective as the older drug. This also seems to apply to the measures of mental state ([Analysis 6.30](#)).

6.3 Adverse effects

It was a concern that these industry-sponsored trials were designed to under-report adverse effects. It had been stipulated in the design of a trial that adverse effects would be reported only if, for example, more than 10% of people in a group experienced them ([Kinon 2001d](#)). Therefore, should 9% of people experienced worrying skin reactions in [Kinon 2001d](#), this could have been omitted from the report.

Nevertheless, it does seem clear that haloperidol does cause more movement disorders than olanzapine. This must be seen as an advantage for the newer drug when being compared with haloperidol used alone.

6.4 Economic

There was no measure for economic outcomes. In older trials this is understandable, but not for comparisons of expensive drugs and the older compounds. It is likely that the expensive olanzapine could save money in the short term - but, disappointingly, this is supposition.

7. COMPARISON 7. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: f. PERPHENAZINE

Considering the recent resurgence of interest in perphenazine (Lieberman 2005), this is an important comparison. The one relevant study ($n = 44$) at least proves that comparative testing of these two treatments is possible, although outcomes are limited and the grade of evidence largely poor (Summary of findings 7). Now that data suggest that perphenazine is a real option for people wanting a more gentle effective antipsychotic drug (Lieberman 2005), there is the possibility that it also would be of value in the emergency situation.

7.1 Global outcome - no improvement

Limited data suggest global impression of improvement is not different between groups (Analysis 7.1).

7.2 Adverse effects

Considerably more people allocated to haloperidol experienced one or more adverse effects (Analysis 7.2) and required antiparkinson medication (Analysis 7.3), but these were not statistically significant in this small trial. Nevertheless, there is a suggestion that perphenazine may have advantages in terms of adverse effects.

This must be an area for future research. However, without the push of industry it is possible that perphenazine will continue to be under evaluated.

8. COMPARISON 8. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: g. QUETIAPINE

We found only one study ($n = 80$) with the most of the outcomes rated at 28 days and of low grade (Summary of findings 8).

8.1 Specific behaviour - agitation

No differences were found between haloperidol and quetiapine across when using rating scales (Analysis 8.1; Analysis 8.2).

8.2 Mental state

Haloperidol and quetiapine showed similar results (Analysis 8.3); however, the Positive and Negative Syndrome Scale (PANSS) and sub-scales were rated at 28 days, inappropriately with the aim of this review.

8.3 Adverse effects

More people allocated to haloperidol experienced one or more adverse effects (Analysis 8.4); this held true when taking into account movement disorders rating scales (Analysis 8.5).

9. COMPARISON 9. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: h. RISPERIDONE

We found three relevant studies ($n = 314$), the outcomes of which we had to grade as of moderate, low or very low quality (Summary of findings 9).

9.1 Tranquillisation or asleep, agitation, need for more benzodiazepine

We thought the best graded evidence was that, if given haloperidol as well as lorazepam, more people were asleep compared with risperidone – also plus lorazepam (Analysis 9.1). Tranquillisation is

often now considered desirable rather than full sedation to sleep, but in certain circumstances the latter may be preferable. The key issue is whether agitation is decreased and, according to PANNS-EC scores, both medications are effective (Analysis 9.3).

In the trial randomising lorazepam as part of the package of care, more subsequent benzodiazepines were used. There was no difference between groups (Analysis 9.7), just the indication that if benzodiazepines are part of the package of care they will be used again and again. This could be problematic for the cumulative adverse effects.

9.2 Adverse effects

The two drugs did not have many differences as regards adverse effects; one seems as good or as bad as the other (Analysis 9.9). For binary measures of movement disorder no differences were clearly apparent (Analysis 9.14). Continuous measures of akathisia and movement disorders did, however, favour risperidone (Analysis 9.15). These fine-grain measures may pick up differences not clearly shown by the more blunt binary outcomes. Akathisia is unpleasant and can lead to clinicians misinterpreting that the person is agitated again. Even subtle decreases of akathisia could be valuable in management of agitation and aggression.

9.3 Economic

We found no data on economic outcomes of this comparison. These trials were undertaken at a time when risperidone was not an inexpensive option and omission of this type of data does not seem random.

10. COMPARISON 10. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: i. THIOTHIXINE

The two small studies provided low level data (total $n = 74$, Summary of findings 10).

10.1 Repeated need for rapid tranquillisation, agitation, global outcome

In the two studies that compared haloperidol with thiothixene, there was no outcome measure for tranquillisation or asleep. There were no significant differences between haloperidol and thiothixene with respect to the number of participants who required more than one injection (Analysis 10.1), agitation (Analysis 10.2), and 'no response to the first injection' (Analysis 10.3), but the study reporting these outcomes was very small indeed.

10.2 Adverse effects

There were no differences between the groups in terms of the number of participants who experienced one or more adverse effects (2 RCTs, $n = 74$, RR 1.47, 95% CI 0.97 to 2.22, Analysis 10.4).

The quality of these outcomes was, however, rated as 'low' or 'very low' because of risk of several biases (Summary of findings 10).

11. COMPARISON 11. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: j. ZIPRASIDONE

Ziprasidone is a relatively new drug when compared with others in this set of comparisons. It is clear that the pharmaceutical industry has recognised the paucity of data in this area and that profits are to be made if an expensive drug can be shown to be as effective as the older drug and have fewer adverse effects. We found five

relevant studies (total $n = 829$) but, again we had to grade the quality of outcome evidence as low or very low ([Summary of findings 11](#)). We have reconsidered as to whether we have been too harsh in judging the quality of evidence but, for the reasons outlined in the [Summary of findings 11](#), we feel we have to continue to grade these data as poor. For example, these large studies did not report the outcome of 'Tranquillisation or asleep' or even 'Repeated need for tranquillisation'. In larger trials focused on the coercive treatment of aggressive people we do feel that this is an odd set of omissions.

11.1 Repeated need for tranquillisation, specific behaviours, global outcomes and mental state

All measures are not convincingly better for either drug. The CGI-S measures are better for the ziprasidone group as are some sub-scores of mental state measures.

11.2 Adverse effects

The technique of reporting adverse effects only if more than 2%/5% or even 10% of people suffered the effect results in under-reporting. Haloperidol can be confidently expected to provide frequent adverse effects ([Adams 2013](#)), so this device may assist the new drug seem less toxic.

At 72 hours, significantly more people in the haloperidol group had experienced one or more adverse effects but, by seven days, the difference between the groups was not statistically significant ([Analysis 11.15](#)).

Ziprasidone does seem to cause fewer movement disorders than haloperidol ([Analysis 11.26](#)) and, perhaps less anticholinergic problems. However, even these trials, generated by the company with a pecuniary interest in the outcomes, did nevertheless suggest, that cardiac adverse effects seemed more common in the ziprasidone group ([Analysis 11.28](#)). This must be seen as an important concern.

11.3 Economic

In these trials, so obviously comparing an inexpensive with expensive drug we find no economic analysis. This bias by omission casts a further cloud on reporting of all other data.

12. COMPARISON 12. HALOPERIDOL versus OTHER ANTIPSYCHOTICS: k. ZUCLOPENTHIXOL ACETATE

Although we understand that zuclopenthixol acetate may be a choice when rather longer-term sedation is required that would be expected with haloperidol ([Jayakody 2012](#)), we did expect to find more relevant trials. Whether to use haloperidol or zuclopenthixol acetate is a prevalent clinical dilemma in some countries. We found one small study ($n = 70$), the outcomes of which we had to grade as 'very low' ([Summary of findings 12](#)).

This study reported few outcome with some expected results. For example, in this trial of a little longer duration, more people in the haloperidol group required more than three injections ([Analysis 12.1](#)) and there was the suggestion that, at seven days, haloperidol caused more tremor ([Analysis 12.2](#)).

This pioneering study is helpful in showing that fair testing could be undertaken for this important comparison. At present, however, best evidence from trials does not support or refute the choice one of these drugs over another in the emergency situation.

13. COMPARISON 13. HALOPERIDOL versus BENZODIAZEPINE: a. FLUNITRAZEPAM

We found one very small relevant study ($n = 28$) the outcomes of which were graded as being of very low quality ([Summary of findings 13](#)).

Unsurprisingly with such a small study there were no differences for any outcomes. That even adverse effects were similar highlights how lacking in power this trial was to show real differences. Flunitrazepam may be effective and a real alternative, but this trial does not show this. It does, however, illustrate that trials with this comparison are possible.

14. COMPARISON 14. HALOPERIDOL versus BENZODIAZEPINE: b. LORAZEPAM

Although we found four relevant studies with more than the average number of total participants ($n = 207$), few outcomes contain data from more than 70 people. The use of haloperidol alone in preference to lorazepam or vice versa is based on such limited evidence of poor quality that this could be considered an embarrassment to a subspecialty that recommends coercive use of these interventions ([NICE 2005](#); [NICE 2015](#)).

14.1 Tranquillisation or asleep

Evidence is limited and we had to grade these data as of 'very poor' quality ([Summary of findings 14](#)). There was a suggestion that lorazepam is significantly more sedating after a few hours ([Analysis 14.1](#)). This may not be desirable as tranquillisation is frequently the aim without necessarily sedating.

14.2 Other global outcome

There were no differences in numbers requiring more than one and more than three injections and no difference between groups for the outcome of 'showing no improvement' at 30 and 60 minutes ([Analysis 14.3](#)). Mental state scores on 37 people were the same.

14.3 Adverse effects

These underpowered studies did not highlight many expected differences. However, significantly more participants experienced EPS in the haloperidol group compared with the lorazepam group ([Analysis 14.11](#)). Apnoea was not reported.

15. COMPARISON 15. HALOPERIDOL versus BENZODIAZEPINE: c. MIDAZOLAM

We found one small relevant study with data we could use ($n = 84$), but the limited outcomes we could use for the [Summary of findings 15](#) we had to grade as being of low quality.

15.1 Tranquillisation or asleep: average time

Midazolam is a highly effective hypnotic ([Olkkola 2008](#)) and the skewed data on tranquillisation did seem to support that it was significantly faster than haloperidol in this clinical situation.

15.2 Adverse effects

Midazolam is known to cause apnoea but in this trial it was a person in the haloperidol group that experienced this ([Analysis 15.3](#)). Nevertheless, data from this small study should not be considered conclusive. Other data from better trials ([TREC CG](#); [TREC 2003](#)) and other sources suggest that use of midazolam could be accompanied

with full resource for control of airways and intravenous reversal of the apneic effects with flumazenil (Olkola 2008).

16. COMPARISON 16. HALOPERIDOL versus COMBINATIONS: a. HALOPERIDOL + LEVOMEPRIMAZINE

We found one very small study ($n = 19$), the outcomes of which we had to grade as being of very low quality (Summary of findings 16).

Clinicians find themselves in an unusual position because of the prevalence of haloperidol and its use for psychosis-induced aggression but in the recognition of its toxicity. We think it understandable that researchers investigate whether combinations could offset some adverse effects and levomepromazine may do this, but the included trial is too small to illustrate anything except that such a study is possible.

17. COMPARISON 17. HALOPERIDOL versus COMBINATIONS: b. HALOPERIDOL + LORAZEPAM

We were rather surprised to find that there are only two trials of this comparison. We think that this combination is in common use but the two trials have a total of only just over 100 people and we had to grade outcomes all of very low quality (Summary of findings 17).

17.1 Tranquillisation or asleep

More people in the combination group were asleep at three hours than those in the haloperidol alone group but these were limited very low-grade data. Also, three hours is a very long time when a person is acutely disturbed and not an outcome of great clinical utility in regard to the focus of this review.

17.2 Repeated need for rapid tranquillisation

There was some suggestion that the addition of lorazepam did assist the tranquillisation process but no data were conclusive and all were of very poor grade. Significantly fewer people in the combination group exhibited "no overall improvement by 30 minutes".

17.3 Adverse effects

Overall, there was no difference between the groups. Best evidence suggests that addition of lorazepam does not offset adverse effects. In these small poorly reported and probably poorly conducted trials dystonia was common and could be considered to be at unacceptable levels. People undergoing coercive rapid tranquillisation have enough difficult experiences without the addition of needless distressing adverse effects. We have no data that this will cause further mistrust of healthcare professionals, but it does seem a reasonable assumption.

18. COMPARISON 18. HALOPERIDOL versus COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM

We found one relevant study ($n = 60$) with outcomes of very low grade only (Summary of findings 18).

18.1 Repeated need for tranquillisation, specific behaviours, global outcomes

An overall advantage in favour of haloperidol was shown in terms of agitation and aggression rating scales, and need for seclusion or restraint (Analysis 18.1; Analysis 18.2; Analysis 18.3; Analysis 18.4; Analysis 18.5; Analysis 18.6; Analysis 18.7).

18.2 Adverse effects

Haloperidol and combination data resulted in similar but, again, data from this small study should not be considered conclusive.

19. HALOPERIDOL versus COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE

We found two relevant study ($n = 376$) and outcomes were graded of moderate quality (higher than others in this review) or low quality (Summary of findings 19).

19.1 Tranquillisation or asleep, repeated need for tranquillisation

Haloperidol used on its own clearly is quickly tranquillising, but the combination with promethazine makes is significantly more swift. The difference between being tranquil at 20 and 40 minutes can be of great importance in the clinical front line (Analysis 19.1; Analysis 19.2).

There were no significant differences in terms of requiring additional drugs for tranquillisation (Analysis 19.3).

19.2 Specific behaviours - agitation, aggression

When using agitation and aggression rating scales, an initial advantage for haloperidol was demonstrated (Analysis 19.4; Analysis 19.7); however, this should be replicated by other studies as it appears to come from one trial only of low-moderate quality (Baldacara 2011).

On other measures of aggression there was not much difference (requiring repeat rapid tranquillisation, exhibiting further aggressive outburst within 24 hours. More people in the haloperidol alone group had to be restrained by 120 minutes, but the difference between groups was not statistically significant. However, significantly more people in the haloperidol alone group required extra visits from the doctor, which is an indication of the clinical concern of the healthcare professionals. Such visits can be for further management of aggression or disturbance, or to address adverse effects.

19.3 Adverse effects

Haloperidol is a potent cause of adverse effects. Significantly more people in the haloperidol alone group than the combination group experienced one or more adverse effects. Acute dystonia – a most distressing effect – was common ($n = 316$, RR 19.48, 95% CI 1.14 to 331.92, Analysis 19.20), and frequency of occurrence resulted in this trial being terminated prematurely. Use of promethazine offsets the dystonia. The report specifically states that the Steering Group of this trial felt that the guidance from APA in the USA and NICE in the UK was unethical to continue.

20. COMPARISON 20. HALOPERIDOL versus COMBINATIONS: e. HALOPERIDOL + RISPERIDONE

We found one study ($n = 100$) with outcomes of low-grade quality (Summary of findings 20).

20.1 Specific behaviour, global outcomes

An initial advantage - at 24 hours - for the combination group was demonstrated (Analysis 20.1), even though it was not confirmed by subsequent time points (Analysis 20.2); further studies should be

carried out, in particularly focused on the first hours in order to better understand the effect of the combination versus haloperidol.

More people allocated to the haloperidol group showed improvement when compared to the haloperidol + risperidone group, even if this difference was not significant (Analysis 20.3).

20.2 Adverse effects

The two compared groups showed comparable results (Analysis 20.4, Analysis 20.5, Analysis 20.6).

21. COMPARISON 21. HALOPERIDOL versus COMBINATIONS: f. OLANZAPINE + MAGNESIUM VALPROATE

We found one small study (n = 64) with outcomes of very low-grade quality (Summary of findings 21).

21.1 Specific behaviour - agitation

People allocated in the combination group showed a lower level of agitation, rated using the PANSS-EC, at seven and 14 days; unfortunately, the first time point available in this trial was at 72 hours and showed no differences between the compared groups (Analysis 21.1). These timings are not adequate in order to fulfil the aim of this study, and even more highlight the need for further studies focused on the early effects.

21.2 Global outcomes, mental state

No significant differences were found as an equal number of people showed no improvement in both groups (Analysis 21.2); the PANSS total scores showed a trend over the weeks in advantage for the combination group, even if not significant (Analysis 21.3).

21.3 Adverse effects

The number of people experiencing one or more adverse effects was almost three times in the haloperidol group (Analysis 21.4).

22. COMPARISON 22. HALOPERIDOL versus COMBINATIONS: g. QUETIAPINE + MAGNESIUM VALPROATE

The one study that compared haloperidol with a combination of quetiapine + magnesium valproate was small and outcomes were graded as very low quality (n = 60, Summary of findings 22).

Although this one trial was probably underpowered, there did not seem to be a suggestion that the quetiapine-magnesium valproate combination had any affect over and above that seen with haloperidol alone.

Global effect had to be indirectly inferred from no effect on the PANSS-EC, and further aggressive behaviour had to be surmised from scale-derived data for agitation at 72 hours. Risk of bias was rated as serious because the method of randomisation was not reported, allocation concealment was not stated and it was an open-label trial. There was no measure for economic outcomes. Any use of this quetiapine-valproate combination must be considered experimental.

23. COMPARISON 23. HALOPERIDOL versus COMBINATIONS: h. RISPERIDONE + CLONAZEPAM

We found two studies (n = 305) and outcomes were graded as of low and very low grade (Summary of findings 23).

23.1 Tranquillisation or asleep, specific behaviour

There were no differences when comparing the number of people not asleep in the first four hours (Analysis 23.1) and the PANSS-EC rating scores (Analysis 23.2; Analysis 23.3; Analysis 23.4; Analysis 23.5).

23.2 Global outcomes, mental state

More people allocated to the haloperidol group experienced an improvement across the explored time points, even though the difference was not significant (Analysis 23.6). At five days when looking at the PANSS total scores, the two groups showed similar results (Analysis 23.7).

23.3 Adverse effects

More adverse events were recorded in the haloperidol group (Analysis 23.8), with a significant predominance of movement disorders-related adverse effects (Analysis 23.11).

24. COMPARISON 24. HALOPERIDOL versus COMBINATIONS: i. ZIPRASIDONE + CLONAZEPAM

We found one study (n = 76) and outcomes were graded as of very low quality (Summary of findings 24).

There did not seem to be a suggestion of superiority of ziprasidone + clonazepam than haloperidol alone; no significant differences were found between the compared groups.

Overall completeness and applicability of evidence

1. Completeness

Despite the great number of comparisons in this review, there are no data on several important questions. For example, although guidelines recommend use of haloperidol intravenously in exceptional circumstances, not one randomised controlled trial was found that investigated the efficacy and safety of haloperidol via this route of administration. For other routes of administration, even where some data do exist, evidence is graded as being of largely poor quality. For example, there is no strong evidence of the clear advantage of olanzapine or risperidone over the older haloperidol. The evidence for perphenazine is not good either, but it could be seen as an inexpensive option to haloperidol. In addition, not one of the studies evaluated satisfaction with care, acceptance of treatment, quality of life or economic costs. Unfortunately, a large proportion of the efficacy and safety data produced by the trials was left redundant due to poor and/or selective reporting. This is clearly an area of evaluation in which researchers have invested heavily and this leaves those compiling guidance having to use judgements that are based less on good evidence than they should be.

2. Applicability

Many of the included trials were more of the explanatory type of study rather than practical/pragmatic and easily applicable to real-world practice (Thorpe 2009). Few seem to have been designed with clear clinical needs as a priority. This does not mean that applicability is impossible. It just necessitates a level of conjecture that could have been lessened should more pragmatic trials have been available. Two examples of how trials are near or far from real-world practice and needs are outlined below.

2.1 Participants seen in real-world practice

Many of these trials seem to have been undertaken in circumstances very far from real clinical practice. For example, the UK's NICE guidelines recommend that rapid tranquillisation should only be considered "once de-escalation and other strategies have failed to calm the service user" (NICE 2005, p14) and that "p.r.n. medication can be used as part of a de-escalation strategy but p.r.n. medication used alone is not de-escalation." (NICE 2015, p15). However, in a number of studies, participants (in some cases their legal representatives) were required to provide written informed consent (Battaglia 2002; Breier 2002; Bristol Myers 2004a; Bristol-Myers 2005b; Brook 2000; Eli Lilly 2004; Kinon 2001d; Currier 2004; Higashima 2004; Lim 2010). Further, participants in many studies all had to undergo a screening period ranging from two to 96 hours (Battaglia 2002; Breier 2002; Bristol Myers 2004a; Bristol-Myers 2005b; Brook 2000; Currier 2004; Eli Lilly 2004; Fruensgaard 1977; Kinon 2001d; Lim 2010; Shu 2010). Participants in these studies have intervention delayed for screening and are deemed competent to give informed consent - a situation that is not common in the heat of an emergency and makes applicability problematic - a matter of conjecture.

Other studies, however, recognised that conduct of the study had to involve deferred or waived consent - and in this way making it more like clinical practice. In Battaglia 1997, Huf 2007, Nobay 2004 and Walther 2014, informed consent was obtained wherever possible. It was waived in Bailine 1987, Dorevitch 1999 and Foster 1997 because it was acknowledged that if people were that violent or psychologically disorganised to require rapid tranquillisation they would not be likely to be able to give informed consent. In Tuason 1986, informed consent was obtained as soon as participants condition permitted and in Resnick 1984 it was deemed that participants were unable to give informed consent because they were involuntary patients. This last group of trials is likely to be involving people closer to real-world circumstances.

2.2 Outcomes of value in real-world practice

Where people are a risk to themselves or others, rapid tranquillisation is often used to prevent the prolonged physical restraint of people, which can result in injury and death. It was therefore surprising that few studies measured tranquillisation, asleep, or global functioning outcomes at time points that one would associate with a rapid intervention. There were only three studies that measured tranquil or asleep on or around 30 minutes (Huf 2007; Currier 2004; Fruensgaard 1977), with a further two measuring at 60 minutes (Kinon 2001d; Salzman 1991), with global improvement being measured at 30 minutes in the Garza-Trevino 1989 and at 60 minutes in the Kewala 1984 study. Also, not all studies documented simple and important outcomes such as repeated need for rapid tranquillisation. We are therefore left inferring the clinical implications of outcomes that are not recorded in routine care. This is an entirely avoidable situation if those designing trials listened to the needs of clinicians and people who are subject to this coercive care.

Quality of the evidence

1. General

Twenty-nine of the 41 included studies were published after the first CONSORT guidance (1996). Few studies complied with this internationally agreed standard. Disappointingly, a large

proportion of data was rendered unusable, which could have been avoided had the trials been better reported.

2. Specific

Although this review includes 41 studies and approximately 3500 participants, there were 24 different comparisons. Trials were largely very small and meta-analysis could be undertaken for a minority of comparisons and, even then, for only some outcomes. Overall, the majority of the evidence had to be graded as 'very low', usually because of considerable - and often avoidable - limits in research design, selective reporting and imprecision.

The best available evidence came from the Huf 2007 trial which compared haloperidol with haloperidol plus promethazine, from the Calver 2015 trial which compared haloperidol with droperidol, and from the Nobay 2004 trial that compared haloperidol with midazolam. Nevertheless, these studies also had limitations to their research design which led to the evidence being graded as 'moderate'.

Potential biases in the review process

There are a number of ways in which bias could have been introduced into this review.

i. Double-data extraction

In some systematic reviews, two people independently inspect all citations obtained from the search and extracted the data to minimise bias. However, this was not the case with this review, with only 20% of the screening of citations and only 10% of the data extraction being performed by two review authors.

ii. Potential conflict of interest

Another source of potential bias is that one of the authors of this review (CEA) was an investigator in the Huf 2007 trial.

Agreements and disagreements with other studies or reviews

The review authors are not aware of any other studies or reviews that have focused solely on use of haloperidol alone for people who are aggressive secondary to serious mental illnesses. Pratt 2008 conducted a systematic review (not meta-analysis), focusing on the practice of rapid tranquillisation in the UK and concluded that there was a lack of high-quality evidence.

AUTHORS' CONCLUSIONS

Implications for practice

1. For people agitation/aggression due to psychosis

Some clinical guidelines recommend the use of haloperidol, either intramuscularly combined with benzodiazepine or even, in the extreme situation, given intravenously. There is no good-quality evidence for the latter. These treatments are administered under coercion and people having experienced this very unpleasant situation, when recovered, may wish to enquire just why such a treatment was administered. They could suggest that an advanced directive be stipulated that only treatments given either from a firm evidence base or within the context of a randomised trial - should they fall so ill again. All evidence reported in this review is limited and flawed. Best evidence is probably for the use of

haloperidol combined with promethazine; the last update of UK's NICE guidelines supports this evidence (NICE 2015). Evidence of the benefit of haloperidol combined with lorazepam is limited to a few very small trials.

People do not like experiencing dystonia. This is common with haloperidol used with nothing to offset this problem. One larger and independent trial was stopped early because it was felt that the NICE/APA recommended use of haloperidol on its own was unethical (Huf 2007). People with schizophrenia may feel that coercion is enough without use of treatments of dubious ethicality.

2. For clinicians

Use of haloperidol clearly calms difficult situations. Clinicians (as well as people with psychosis and aggression), however, are left vulnerable when evidence is of such limited quality and so thinly spread over many comparisons. In some situations, where a person is clearly both mentally unwell and aggressive, haloperidol may be the only choice - and in this case, a reasonable one. However, where resources are more available, haloperidol used completely on its own does not seem defensible. Haloperidol may be made more sedating by lorazepam but evidence is poor. Adverse effects may be offset by anticholinergics but data are lacking and this is unlikely to happen with lorazepam. Best evidence exists for addition of the sedating antihistamine promethazine. If researchers continue to let down the clinical community by avoiding evaluative studies in this difficult area clinicians should request such investigations formally. Not to have good evidence in this area is difficult to justify.

3. For policy makers

Even in the situation of very limited high-grade data, guidance has to happen. There is enough evidence to suggest that haloperidol given intramuscularly may be helpful, but probably giving it alone should be avoided. Theoretically anticholinergics are indicated at the same time and this has been best demonstrated by the use of the anticholinergic-antihistamine promethazine - at least within the context of use of restraints. There is not compelling evidence that newer drugs are better enough to justify their routine use and no economic data whatsoever. More independent trials are indicated.

Implications for research

1. General

All studies should now comply with the Consolidated Standards of Reporting Trials (CONSORT, Moher 2001). More transparency in the reporting of randomised controlled trials, would enable readers to understand the design, conduct, analysis and interpretation, and to assess the validity of results. Although binary data are easier to interpret, where continuous data are used, some measure of

variance should be provided. Data presented in graphs should be accompanied by exact numbers and standard deviations in the text.

The review authors strongly highlight that, despite the recommendations provided by CONSORT reporting guidelines and the first edition of this review, out of nine studies included in this update, only one study (Calver 2015) contributed to a moderate-grade evidence, whilst the remaining eight studies provided low- or very low-grade evidence only.

2. Specific

This review is an indictment of six decades of research and routine coercive use. Large pragmatic well-designed randomised trials led by independent researchers, which measure simple and meaningful outcomes such as 'tranquil', 'asleep', 'serious adverse effect', 'needing additional medication', 'further aggressive episodes' are still required. These studies do not need large funds. They just need modest financial support and firm help from ethics committees who understand the need to produce good evidence for practice in this difficult edge of health care. See Table 3 for a suggested design for a study.

ACKNOWLEDGEMENTS

We would like to thank Gene Amason, Dr Anand, Dr Baldaçara, Dr Dubin, Dr El-Sayeh, Dr Herrera, Dr Isbister, Dr Kinon, Dr Lerner, Dr Mueller, Dr Resnick, Dr Simon and Dr Spivak for answering our requests for further information regarding studies.

In addition, we would like to thank Sam Roberts at the Cochrane Schizophrenia Group Editorial Base in Nottingham for designing the search strategy, Jun Xia of Systematic Review Solutions (<http://www.srs-org.com/>) for data extraction of Chinese papers and correspondence and Claire Irving for her help and patience. Also, Ben Gray for writing the Plain language summary and Hannah Jones for her contributions for the previous version of this review.

The Cochrane Schizophrenia Group Editorial Base in Nottingham produces and maintains standard text for use in the Methods section of their reviews. We have used this text as the basis of what appears here and adapted it as required. Further, this is one of a series of reviews (Table 1). The authors of this review also acknowledge the help and guidance of the structure of the Chlorpromazine for psychosis-induced aggression or agitation (Ahmed 2010).

This project was supported by the National Institute for Health Research, via Cochrane Infrastructure funding to the Schizophrenia Group. The views and opinions expressed therein are those of the authors and do not necessarily reflect those of the Systematic Reviews Programme, NIHR, NHS or the Department of Health.

REFERENCES

References to studies included in this review

Bailine 1987 {published data only}

Bailine SH, Lesser MS, Krubit G, Ravasz TJ, Davis RA, Kane JM. Comparison of IM haloperidol and IM chlorpromazine in the treatment of acutely psychotic patients. *Psychiatric Hospital* 1987;**18**(3):127-9.

Baldacara 2011 {published data only}

Baldacara L, Sanches M, Cordeiro DC, Jackowski AP. Rapid tranquilization for agitated patients in emergency psychiatric rooms: A randomized trial of olanzapine, ziprasidone, haloperidol plus promethazine, haloperidol plus midazolam and haloperidol alone. *Revista Brasileira de Psiquiatria* 2011; Vol. 33, issue 1:30-9.

Battaglia 1997 {published data only}

Battaglia J, Moss S, Rush J, Kang J, Mendoza R, Leedom L, et al. Haloperidol, lorazepam, or both for psychotic agitation? A multicenter, prospective, double-blind, emergency department study. *American Journal of Emergency Medicine* 1997;**15**(4):335-40. [MEDLINE: 9217519]

Battaglia 2002 {published data only}

* Battaglia J, David SR, Alaka K, Meehan K, Wright P. Calming versus sedative effects of IM olanzapine in agitated patients. *Schizophrenia Research* 2002;**53**(3 Suppl 1):183.

Battaglia J, Houston JP, Ahl J, Meyers AL, Kaiser CJ. A post hoc analysis of transitioning to oral treatment with olanzapine or haloperidol after 24-hour intramuscular treatment in acutely agitated adult patients with schizophrenia. *Clinical Therapeutics* 2005;**27**(10):1612-8. [MEDLINE: 16330297]

Battaglia J, Lindborg SR, Alaka K, Meehan K, Wright P. Calming versus sedative effects of intramuscular olanzapine in agitated patients. *American Journal of Emergency Medicine* 2003;**21**(3):192-8. [MEDLINE: 12811711]

David SR, Battaglia J, Alaka K, Meehan K, Wright P. Calming versus sedative effects of IM olanzapine in agitated patients. *Proceedings of the 11th Congress of the Association of European Psychiatrists*; 2002 May 4-8; Stockholm, Sweden 2002;**17**(Suppl 1):104s.

David SR, Beasley CM Jr, Alaka K. Analysis of the QTC interval in acutely agitated patients with schizophrenia, bipolar mania, or dementia treated with intramuscular (IM) olanzapine vs. IM placebo or IM haloperidol. *European Neuropsychopharmacology* 2001;**11**(3):276.

David SR, Beasley CM Jr, Alaka K. QTC intervals during treatment with IM olanzapine in acutely agitated patients. *Schizophrenia Research* 2002;**53**(3 Suppl 1):164.

David SR, Meehan K, Birkett MA, Wright P, Ferchland I, Alaka K, et al. Intramuscular olanzapine versus intramuscular haloperidol and intramuscular placebo: an international double-blind study in acutely agitated patients with schizophrenia. *Schizophrenia Research* 2001;**49**(1-2):224.

Houston JP, Kaiser CJ, Ahl J. Oral olanzapine transition dose following intramuscular olanzapine treatment. International Congress on Schizophrenia Research; 2005 Apr 2-5, Savannah, Georgia USA. 2005.

King KG, Wright P, Meehan K, Birkett M, David SR, Taylor CC. Transition from intramuscular to oral olanzapine. Proceedings of the 12th World Congress of Psychiatry; 2002 Aug 24-29; Yokohama, Japan. 2002.

Lindborg SR, Beasley CM, Alaka K, Taylor CC. Effects of intramuscular olanzapine vs haloperidol and placebo on QTc intervals in acutely agitated patients. *Psychiatry Research* 2003;**119**(1-2):113-23. [MEDLINE: 12860365]

Meehan K, Alaka KJ, Birkett M, Breier A, David S, Ferchland I, et al. Short-acting intramuscular olanzapine in acutely agitated schizophrenic patients: A randomized double blind trial vs. placebo and haloperidol. *Psychosomatics* 2001;**42**(2):197-8.

Meehan K, Wright P, David SR, Alaka K, Taylor CC, Breier A, et al. Agitation as a treatable symptom common to schizophrenia, acute mania and dementia. Proceedings of the 57th Annual Meeting of the Society of Biological Psychiatry; 2002 May 16-18; Philadelphia, Pennsylvania, USA. 2002.

Meehan K, Wright P, David SR, Alaka K, Taylor CC, Breier A, et al. Agitation as a treatable symptom common to schizophrenia, acute mania, and dementia. *Schizophrenia Research* 2002;**53**(3 Suppl 1):63.

Wright P, Birkett M, David SR, Meehan K, Ferchland I, Alaka KJ, et al. Double-blind, placebo-controlled comparison of intramuscular olanzapine and intramuscular haloperidol in the treatment of acute agitation in schizophrenia. *American Journal of Psychiatry* 2001;**158**(7):1149-51. [MEDLINE: 11431240]

Wright P, Birkett M, Ferchland I, David S, Alaka K, Pullen P, et al. A double-blind study of intramuscular olanzapine, haloperidol and placebo in acutely agitated schizophrenic patients. *European Neuropsychopharmacology* 2000;**10**(Suppl 3):S304.

Wright P, Birkett M, Meehan K, Ferchland I, David S, Alaka K, et al. A double-blind study of intramuscular olanzapine, haloperidol and placebo in acutely agitated schizophrenic patients. *International Journal of Neuropsychopharmacology* 2000;**3**(Suppl 1):S139.

Wright P, Lindborg SR, Birkett M, Meehan K, Jones B, Alaka K, et al. Intramuscular olanzapine and intramuscular haloperidol in acute schizophrenia: antipsychotic efficacy and extrapyramidal safety during the first 24 hours of treatment. *Canadian Journal of Psychiatry [Revue Canadienne de Psychiatrie]* 2003;**48**(11):716-21. [MEDLINE: 14733451]

Wright P, Meehan K, Birkett M, Lindborg SR, Taylor CC, Morris PBA. A comparison of the efficacy and safety of olanzapine versus haloperidol during transition from intramuscular to oral therapy. *Clinical Therapeutics* 2003;**25**(5):1420-8. [MEDLINE: 12867218]

Breier 2002 {published data only}

* Breier A, Meehan K, Birkett M, David S, Ferchland I, Sutton V, et al. A double-blind, placebo-controlled dose-response comparison of intramuscular olanzapine and haloperidol in the treatment of acute agitation in schizophrenia. *Archives of General Psychiatry* 2002;**59**(5):441-8.

Breier A, Wright P, Birkett M, David S, Brook S, Meehan K. A double-blind dose response study comparing intramuscular (IM) olanzapine, haloperidol, and placebo in acutely agitated schizophrenic patients. *Biological Psychiatry* 2001;**49**(8):121S.

Breier A, Wright P, Birkett M, Meehan K, David S, Brook S. A double-blind dose response study comparing intramuscular olanzapine, haloperidol and placebo in acutely agitated schizophrenic patients. Proceedings of the 39th Annual Meeting of the American College of Neuropsychopharmacology; 2000 Dec 10-14; San Juan, Puerto Rico. 2000.

Breier AF, Wright P, Birkett M, Meehan K, David SR, Brook S. Intramuscular olanzapine: dose-related improvement in acutely agitated patients with schizophrenia. Proceedings of the 154th Annual Meeting of the American Psychiatric Association; 2001 May 5-10; New Orleans, Louisiana, USA. Marathon Multimedia, 2001.

Smith RC. 10 mg intramuscular olanzapine reduces acute agitation in schizophrenia more effectively than lower doses [Commentary]. *Evidence-Based Mental Health* 2003;**6**(1):27. [MEDLINE: 12588833]

Wright P, Birkett MA, Meehan K, David SR, Brook S, Breier A. A double-blind dose response study comparing intramuscular olanzapine, haloperidol and placebo in acutely agitated schizophrenic patients. *Schizophrenia Research* 2001;**49**(1-2):250-1.

Bristol Myers 2004a {published data only}

Andreuzina R, Josiassen RC, Marcus RN, Oren DA, Manos G, Stock E, et al. Intramuscular aripiprazole for the treatment of acute agitation in patients with schizophrenia or schizoaffective disorder: a double-blind, placebo-controlled comparison with intramuscular haloperidol. *Psychopharmacology* 2006;**188**(3):281-92. [MEDLINE: 16953381]

Andreuzina R, Marcus RN, Oren DA, Manos G, Stock E, Carson WH, et al. Intramuscular aripiprazole or haloperidol and transition to oral therapy in patients with agitation associated with schizophrenia: sub-analysis of a double-blind study. *Current Medical Research and Opinion* 2006;**22**(11):2209-19. [MEDLINE: 17076982]

* Bristol-Myers S. A randomized, double-blind comparison of the efficacy and safety of aripiprazole intramuscular formula, haloperidol, or placebo in the treatment of acutely agitated patients with a diagnosis of schizophrenia or schizoaffective disorder. <http://www.clinicalstudyresults.org/> 2004.

Citrome LL, Vester-Blokland E, Archibald D, McQuade R, Kostic D, Pikalov A, et al. Benefits of a second dose of intramuscular (IM) aripiprazole to control agitation in patients with schizophrenia or bipolar I disorder. Proceedings of the

159th Annual Meeting of the American Psychiatric Association; 2006 May 20-25, Toronto, Canada. 2006.

Currier GW, Crandall D, Archibald D, Nyilas M, Kostic D, Pikalov A, et al. Intramuscular (IM) aripiprazole controls agitation in patients with schizophrenia or bipolar disorder without excessive sedation. Proceedings of the 159th Annual Meeting of the American Psychiatric Association; 2006 May 20-25; Toronto, Canada. 2006.

Daniel DG, Currier GW, Zimbardo DL, Allen MH, Oren D, Manos G, et al. Efficacy and safety of oral aripiprazole compared with haloperidol in patients transitioning from acute treatment with intramuscular formulations. *Journal of Psychiatric Practice* 2007;**13**(3):170-7. [MEDLINE: 17522560]

Daniel DG, Stock EG, Wilber CH, Marcus RN, Carson WH Jr, Manos G, et al. Intramuscular aripiprazole in acutely agitated psychotic patients. Proceedings of the 157th Annual Meeting of the American Psychiatric Association; 2004 May 1-6; New York, USA. 2004.

Dillenschneider A, Oren D, Marcus R, Kostic D, McQuade R, Iwamoto T, et al. Comparison of intramuscular aripiprazole and haloperidol with placebo in acutely agitated schizophrenia patients. *European Neuropsychopharmacology* 2005;**15**(Suppl 3):S509.

Dillenschneider A, Oren D, Marcus R, Kostic D, McQuade R, Iwamoto T, et al. Intramuscular aripiprazole in acute schizophrenia: a pivotal phase III study. Proceedings of the 18th European College of Neuropsychopharmacology Congress; 2005 Oct 22-26; Amsterdam, Netherlands 2005.

McQuade R, Auby P, Oren D, Marcus R, Kostic D, Iwamoto T, et al. Intramuscular aripiprazole in acute schizophrenia: a double-blind, placebo-controlled comparison with IM haloperidol. Proceedings of the 13th Biennial Winter Workshop on Schizophrenia Research; 2006 Feb 4-10; Davos, Switzerland. 2006.

Oren D, Marcus R, Kostic D, McQuade R, Iwamoto T, Archibald D. Efficacy and safety of intramuscular aripiprazole, haloperidol or PLACEBO in acutely agitated patients with schizophrenia or schizoaffective disorder. *Schizophrenia Bulletin* 2005;**31**:499.

Oren D, Marcus R, Kostic D, McQuade R, Iwamoto T, Archibald D. Efficacy and safety of intramuscular aripiprazole, haloperidol or placebo in acutely agitated patients with schizophrenia or schizoaffective disorder. Proceedings of the 158th Annual Meeting of the American Psychiatric Association; 2005 May 21-26; Atlanta, Georgia, USA 2005.

Oren DA, Marcus RN, Manos G, Carson WH, McQuade D. Non-inferiority of intramuscular aripiprazole versus intramuscular haloperidol for the treatment of agitation associated with schizophrenia/schizoaffective disorder. *Schizophrenia Bulletin* 2007;**33**(2):451.

Yocca F, Marcus R, Oren D, Manos G, Carson W, Iwamoto T, et al. Intramuscular aripiprazole in acute schizophrenia: a pivotal phase III study. Proceedings of the 158th Annual Meeting of the American Psychiatric Association; 2005 May 21-26, Atlanta, Georgia, USA 2005.

Bristol-Myers 2005b {published data only}

* Bristol-Myers S. Randomized, double-blind, dose-ranging study of intramuscular aripiprazole in the treatment of acute agitation in patients with a diagnosis of schizophrenia, schizoaffective, or schizophreniform disorder. <http://www.clinicalstudyresults.org/2005>.

Currier GW, Crandall D, Archibald D, Nyilas M, Kostic D, Pikalov A, et al. Intramuscular (IM) aripiprazole controls agitation in patients with schizophrenia or bipolar disorder without excessive sedation. Proceedings of the 159th Annual Meeting of the American Psychiatric Association; 2006 May 20-25; Toronto, Canada. 2006.

Gismondi R, Daniel D, Stock E, Wilber R, Marcus R, Carson W, et al. Intramuscular aripiprazole treatment for acute agitation in patients with psychosis. *European Neuropsychopharmacology* 2004;**14**(Suppl 3):S261.

Kungel M, Czaniera R, Ebrecht M, Werner C, Oren D, McQuade R, et al. Intramuscular aripiprazole for the treatment of acute agitation accompanied with schizophrenia: Subanalysis of a double-blind controlled study to find the dosage. Proceedings of the 25th Symposium of the Arbeitsgemeinschaft Neuropsychopharmakologie und Pharmakopsychiatrie; 2007 Oct 3-6; Munich, Germany. 2007.

Kungel M, Daniel D, Stock E, Wilber R, Marcus R, Carson W, et al. Efficacy and safety of intramuscular aripiprazole in acutely agitated patients with psychosis. Proceedings of the Thematic Conference of the World Psychiatric Association on "Treatments in Psychiatry: An Update"; 2004 Nov 10-13; Florence, Italy. 2004.

Oren DA, Manos G, Markovic O, McQuade RD. Intramuscular aripiprazole for the treatment of acute agitation associated with schizophrenia: Sub-analysis of a double-blind, controlled, dose-ranging study. *European Psychiatry* 2007;**22**:S124-S.

Tran-Johnson TK, Sack DA, Marcus RN, Auby P, McQuade RD, Oren DA. Efficacy and safety of intramuscular aripiprazole in patients with acute agitation: a randomized, double-blind, placebo-controlled trial. *Journal of Clinical Psychiatry* 2007;**68**(1):111-9. [MEDLINE: 17284138]

Brook 2000 {published data only}

Brook S, Krams M, Gunn K. Intramuscular (IM) ziprasidone vs. IM haloperidol in patients with acute, non-organic psychosis. Proceedings of the 9th Congress of the Association of European Psychiatrists; 1998 Sep 20-24; Copenhagen, Denmark. 1998.

Brook S, Krams M, Gunn KP. Intramuscular ziprasidone versus intramuscular haloperidol. Proceedings of the 21st Collegium Internationale Neuro-Psychopharmacologicum Congress; 1998 Jul 12-16; Glasgow, UK 1998.

Brook S, Krams M, Gunn KP, and the Ziprasidone IM Study Group. The efficacy and tolerability of intramuscular (IM) ziprasidone versus IM haloperidol in patients with acute non-organic psychosis. Proceedings of the 151st Annual Meeting of the American Psychiatric Association; 1998 May 30 - Jun 4; Toronto, Ontario, Canada. 1998.

* Brook S, Lucey JV, Gunn KP. Intramuscular ziprasidone compared with intramuscular haloperidol in the treatment of acute psychosis. *Journal of Clinical Psychological Medicine* [####] 2000;**61**(12):933-41.

Daniel DG, Brook S, Reeves KR. An overview of the efficacy and safety of rapid acting intramuscular ziprasidone. *European Neuropsychopharmacology* 2000;**10**(Suppl 3):S239.

Zimbroff DL, Daniel DG, Brook S, Reeves KR. An overview of the efficacy and safety of rapid-acting intramuscular ziprasidone. Proceedings of the 153rd Annual Meeting of the American Psychiatric Association; 2000 May 13-18; Chicago, Illinois, USA. 2000.

Calver 2015 {published data only}

Calver L, Drinkwater V, Gupta R, Page CB, Isbister GK. Droperidol v. haloperidol for sedation of aggressive behaviour in acute mental health: Randomized controlled trial. *British Journal of Psychiatry* 2015; Vol. 206, issue 3:223-8.

Currier 2004 {published data only}

Currier GW, Chou JC, Feifel D, Bossie CA, Turkoz I, Mahmoud R, et al. Acute treatment of psychotic agitation: a randomized comparison of oral treatment with risperidone and lorazepam versus intramuscular treatment with haloperidol and lorazepam. *Journal of Clinical Psychiatry* 2004;**65**(3):386-94. [MEDLINE: 15096079]

Dorevitch 1999 {published data only}

Dorevitch A, Katz N, Zemishlany Z, Aizenberg D, Weizman A. Intramuscular flunitrazepam versus intramuscular haloperidol in the emergency treatment of aggressive psychotic behavior. *American Journal of Psychiatry* 1999;**156**(1):142-4.

Eli Lilly 2004 {published data only}

Chan HY, Ree SC, Su LW, Chen JJ, Chou SY, Chen CK, et al. A double-blind, randomized comparison study of efficacy and safety of intramuscular olanzapine and intramuscular haloperidol in patients with schizophrenia and acute agitated behavior. *Journal of Clinical Psychopharmacology* 2014; Vol. 34, issue 3:355-8.

Eli Lilly. Comparison of intramuscular olanzapine and intramuscular haloperidol in patients with schizophrenia. Eli Lilly and Company Clinical Trial Registry 2004.

* NCT00485901. A double-blind randomized comparison of the efficacy and safety of intramuscular olanzapine and intramuscular haloperidol in acutely agitated patients with schizophrenia. <http://www.clinicaltrials.gov> 2007.

Fang 2012 {published data only}

Fang M, Chen H, Li LH, Wu R, Li Y, Liu L, et al. Comparison of risperidone oral solution and intramuscular haloperidol with the latter shifting to oral therapy for the treatment of acute agitation in patients with schizophrenia. *International Clinical Psychopharmacology* 2012; Vol. 27, issue 2:107-13.

Fitzgerald 1969 {published data only}

Fitzgerald CH. A double-blind comparison of haloperidol with perphenazine in acute psychiatric episodes. *Current Therapeutic Research* 1969;**11**(8):515-9.

Foster 1997 {published data only}

Foster S, Kessel J, Berman ME, Simpson GM. Efficacy of lorazepam and haloperidol for rapid tranquilization in a psychiatric emergency room setting. *International Clinical Psychopharmacology* 1997;**12**(3):175-9. [MEDLINE: 9248875]

Fruensgaard 1977 {published data only}

Fruensgaard K, Korsgaard S, Jorgensen H, Jensen K. Loxapine versus haloperidol parenterally in acute psychosis with agitation. A double-blind study. *Acta Psychiatrica Scandinavica* 1977;**56**(4):256-64. [MEDLINE: 335787]

Garza-Trevino 1989 {published data only}

Garza-Trevino ES, Hollister LE, Overall JE, Alexander WF. Efficacy of combinations of intramuscular antipsychotics and sedative-hypnotics for control of psychotic agitation. *American Journal of Psychiatry* 1989;**146**(12):1598-601.

Guo 2007 {published data only}

Guo C-R. Study of quetiapine combined with magnesium valproate release tablets in treatment of schizophrenia with symptoms of elation and agitation [奎硫平合并丙戊酸镁缓释片治疗精神分裂症兴奋激越研究]. *Linchuang Jingshen Yixue Zazhi* [#####] 2007;**17**(3):183-4.

Higashima 2004 {published data only}

Higashima M, Takeda T, Nagasawa T, Hirao N, Oka T, Nakamura M, et al. Combined therapy with low-potency neuroleptic levomepromazine as an adjunct to haloperidol for agitated patients with acute exacerbation of schizophrenia. *European Psychiatry* 2004;**19**(6):380-1. [MEDLINE: 15363480]

Huf 2007 {published data only}

Barbui C. Intramuscular haloperidol plus promethazine is more effective and safer than haloperidol alone for rapid tranquillisation of agitated mentally ill patients. *Evidence-Based Mental Health* 2008;**11**(3):86-7.

Huf G. Haloperidol plus promethazine versus haloperidol for psychosis induced aggression. Data on file 2004.

* Huf G, Coutinho ESF, Adams CE. Rapid tranquillisation in psychiatric emergency settings in Brazil: pragmatic randomised controlled trial of intramuscular haloperidol versus intramuscular haloperidol plus promethazine. *BMJ* 2007;**335**(7625):869-72. [MEDLINE: 17954515]

Huf G, Coutinho ESF, Adams CE. The pharmacological management of agitated patients in emergency psychiatric hospitals in Rio de Janeiro, Brazil: the results of two pragmatic randomized clinical trials. Proceedings of the 5th European Congress on Violence in Clinical Psychiatry; 2007 Oct 25-27; Amsterdam, The Netherlands. 2007.

ISRCTN83261243. TREC2 - Rapid tranquillisation for agitated patients in emergency psychiatric rooms in Rio de Janeiro. A randomised trial of intramuscular haloperidol

versus intramuscular haloperidol + promethazine. <http://www.controlled-trials.com> 2005.

Migon MN, Coutinho ES, Huf G, Adams CE, Cunha GM, Allen MH. Factors associated with the use of physical restraints for agitated patients in psychiatric emergency rooms. *General Hospital Psychiatry* 2008;**30**(3):263-8. [MEDLINE: 18433659]

Kewala 1984 {published data only}

Kewala S, Ban TA, Berney SA, Wilson WH. Rapid tranquilization: A comparative study of thiothixene and haloperidol. *Progress in Neuro-Psychopharmacology and Biological Psychiatry* 1984;**8**:77-83.

Kinon 2001d {published data only}

Kinon B, Rotelli MD, Gilmore JA. Efficacy of olanzapine and adjunctive lorazepam to control agitation in schizophrenia. *International Journal of Neuropsychopharmacology* 2002;**5**(Suppl 1):S128.

Kinon BJ, Ahl J, Rotelli MD, McMullen E. Efficacy of accelerated dose titration of olanzapine with adjunctive lorazepam to treat acute agitation in schizophrenia. *American Journal of Emergency Medicine* 2004;**22**(3):181-6. [MEDLINE: 15138953]

* Kinon BJ, Wang L, Rotelli MD, Gilmore JA. The efficacy of olanzapine plus adjunctive lorazepam to control acute agitation in schizophrenia. *European Neuropsychopharmacology* 2001;**11**(3):278.

Li 2006 {published data only}

Li L-H, Zhao J-P, Xu X-F. A comparative study of intramuscular ziprasidone and haloperidol in treating acute agitation in schizophrenia [国产齐拉西酮与氟哌啶醇注射液治疗精神分裂症急性期兴奋激越症状的对照研究]. *Chinese Journal of Psychiatry* 2006;**39**(4):216-9.

Liang 2013 {published data only}

Liang Y, Su Y, Huang J, Tang M, Li K, Yang F, et al. Effect of quetiapine and haloperidol on QTc interval in treating schizophrenic patients with excitement and agitation [喹硫平和氟哌啶醇治疗精神分裂症急性期兴奋激越对QTc间期的影响]. *Chinese New Drugs Journal* [Zhongguo Xin Yao Za Zhi; 中国新药杂志] 2013; Vol. 22, issue 16:1886-9+1894.

* Liang Y, Su Y, Huang J, Tang M, Li K, Yang P, et al. A randomized controlled study of effect and safety of quetiapine and haloperidol in treatment of schizophrenic patients with excitement and agitation [喹硫平和氟哌啶醇治疗精神分裂症急性期兴奋激越的多中心随机对照研究]. *Chinese Mental Health Journal* [Zhongguo Xin Li Wei Sheng Za Zhi; 中国心理卫生杂志] 2013; Vol. 27, issue 4:262-7.

Lim 2010 {published data only}

Kim JJ, Lim HK, Lee CU, Lee C, Pae CU, Paik IH. Risperidone orodispersible tablet and intramuscular haloperidol in treatment of acute psychotic agitation. *European Neuropsychopharmacology* 2007;**17**(Suppl 4):S467.

* Lim HK, Kim JJ, Pae CU, Lee CU, Lee C, Paik IH. Comparison of risperidone orodispersible tablet and intramuscular haloperidol in the treatment of acute psychotic agitation: A randomized

open, prospective study. *Neuropsychobiology* 2010;**62**(2):81-6. [ISI: 000280037400001]

Liu 2012 {published data only}

刘锐, 刘云, 宁南义, 孙建中, 孙亚军, 孙同勋, et al. Consolidation of short-term risperidone intramuscular haloperidol in the treatment of acute agitation in schizophrenia randomized controlled study of the efficacy of behavior [Google Translate] [利培酮合并短期肌注氟哌啶醇治疗精神分裂症急性期激越行为疗效的随机对照研究]. *中国医药导报* 2012, issue 35:104-6.

Man 1973 {published data only}

Man-Pang L, Chen CH. Rapid tranquilization of acutely psychotic patients with intramuscular haloperidol and chlorpromazine. *Psychosomatics* 1973;**14**(1):59-63. [MEDLINE: 4605009]

Nobay 2004 {published data only}

Nobay F, Simon BC, Levitt MA, Dresden GM. A prospective, double-blind, randomized trial of midazolam versus haloperidol versus lorazepam in the chemical restraint of violent and severely agitated patients. *Academic Emergency Medicine* 2004;**11**(7):744-9. [MEDLINE: 15231461]

Paprocki 1977 {published data only}

Paprocki J, Versiani M. A double-blind comparison between loxapine and haloperidol by parenteral route in acute schizophrenia. *Current Therapeutic Research* 1977;**21**(1):80-100.

Reschke 1974 {published data only}

Reschke RW. Parenteral haloperidol for rapid control of severe, disruptive symptoms of acute schizophrenia. *Diseases of the Nervous System* 1974;**35**:112-5.

Resnick 1984 {published data only}

Resnick M, Burton BT. Droperidol vs. haloperidol in the initial management of acutely agitated patients. *Journal of Clinical Psychological Medicine* [#####] 1984;**45**(7):298-9.

Ritter 1972 {published data only}

Ritter RM, Davidson DE, Robinson TA. Comparison of injectable haloperidol and chlorpromazine. *American Journal of Psychiatry* 1972;**129**(1):78-81. [MEDLINE: 4556089]

Salzman 1991 {published data only}

Salzman C, Solomon D, Miyawaki E, Glassman R, Rood L. Parenteral lorazepam versus parenteral haloperidol for the control of psychotic disruptive behavior. *Journal of Clinical Psychiatry* 1991;**52**(4):177-80. [MEDLINE: 1673123]

Shi 2010 {published data only}

Shi Y, Dai T, Yi P. olanzapine combined magnesium valproate sustained-release tablets in the treatment of schizophrenia, acute agitation excited [Google Translate] [奥氮平合并丙戊酸镁缓释片治疗精神分裂症急性期兴奋激越研究]. *Modern Practical Medicine* 2010; Vol. 22, issue 4:398-9.

Shu 2010 {published data only}

NCT00723606. A randomized, open-label, multi-center study to evaluate the efficacy and safety of intramuscular ziprasidone in patients with agitation. <http://www.clinicaltrials.gov> 2008.

Pfizer. A randomized, open label, rater blind, flexible dose multi-center study comparing the efficacy and safety of intramuscular ziprasidone with haloperidol for three days in patients with agitation of schizophrenia. http://www.clinicalstudyresults.org/documents/company-study_10505_0.pdf (accessed 13 June 2011).

* Shu L, Zhang H, Wang G, Zhao J, Xie S, Xu X, et al. Intramuscular ziprasidone has improved tolerability over haloperidol and comparable efficacy for control of agitation in schizophrenia in Chinese patients. *Schizophrenia Research* 2010;**117**(2-3):260.

Zhang H, Wang G, Zhao J, Xie S, Xu X, Shi J, et al. Intramuscular ziprasidone versus haloperidol for managing agitation in Chinese patients with schizophrenia. *Journal of Clinical Psychopharmacology* 2013; Vol. 33, issue 2:178-85.

Stotsky 1977 {published data only}

Stotsky BA. Relative efficacy of parenteral haloperidol and thiothixene for the emergency treatment of acutely excited and agitated patients. *Diseases of the Nervous System* 1977;**38**(12):967-73. [MEDLINE: 338270]

Taymeeyapradit 2002 {published data only}

Taymeeyapradit U, Kuasirikul S. Comparative study of the effectiveness of zuclopenthixol acetate and haloperidol in acutely disturbed psychotic patients. *Journal of the Medical Association of Thailand* 2002;**85**(12):1301-8. [MEDLINE: 12678168]

Tuason 1986 {published data only}

Tuason VB. A comparison of parenteral loxapine and haloperidol in hostile and aggressive acutely schizophrenic patients. *Journal of Clinical Psychiatry* 1986;**47**(3):126-9. [MEDLINE: 3512535]

Walther 2014 {published data only}

* Walther S, Moggi F, Horn H, Moskvitin K, Abderhalden C, Maier N, et al. Rapid tranquilization of severely agitated patients with schizophrenia spectrum disorders: a naturalistic, rater-blinded, randomized, controlled study with oral haloperidol, risperidone, and olanzapine. *Journal of Clinical Psychopharmacology* 2014; Vol. 34, issue 1:124-8.

Walther S, Moggi F, Horn H, Moskvitin K, Maier N, Abderhalden C, et al. Rapid tranquilization of severely agitated patients with schizophrenia spectrum disorders: a naturalistic, rater-blinded, randomized controlled study with oral haloperidol, risperidone, and olanzapine. *PharmacoPsychiatry* 2013; Vol. 46, issue 6:A77.

Yin 2012 {published data only}

尹析凡, 王刚. 齐拉西酮注射液治疗精神分裂症激越症状的研究. *中国当代医药* 2012; Vol. 30:9-11.

Zhou 2014 {published data only}

周升宝, 孙晓丹, 庞琦, 李志梅, 赵继舒, 张峰. 齐拉西酮联合氯硝西泮治疗精神分裂症急性期兴奋激越症状对照研究. *临床心身疾病杂志* 2014; Vol. 20, issue 2:17-20.

References to studies excluded from this review

Addington 1996 {published data only}

* Addington DE, Arvanitis LA. 'Seroquel' (quetiapine): an atypical antipsychotic - results from a multiple fixed dose, placebo-controlled study. Proceedings of the 4th International Conference on Schizophrenia - 1996: Breaking down the Barriers; 1996 Oct 6-9; Vancouver, Canada. 1996.

Arvanitis LA, Sweitzer DE, Goldstein JM, Yeung PP. Serequel reduces aggression in patients with acute exacerbation of schizophrenia. Proceedings of the 11th European College of Neuropsychopharmacology Congress; 1998 Oct 31 - Nov 4; Paris, France. 1998.

Borison RL, Arvanitist LA, Miller BG. A comparison of five fixed doses 'seroquel' (ICI 204,636) with haloperidol and placebo in patients with schizophrenia. *Schizophrenia Research* 1996;**18**(2-3):132.

Cantillon M. Quetiapine fumarate reduces aggression. Proceedings of the 11th Annual Meeting of the American Association for Geriatric Psychiatry; 1998 Mar 8-11; San Diego, California, USA. 1998.

Cantillon M, Arvanitis LA, Miller BG, Kowalczyk. 'Seroquel' (quetiapine fumarate) reduces hostility and aggression in patients with acute schizophrenia. Proceedings of the 36th Annual Meeting of the American College of Neuropsychopharmacology; 1997 Dec 8-12; Waikoloa, Hawaii, USA. 1997.

Cantillon M, Goldstein JM. Efficacy of quetiapine fumarate in affective symptoms of schizophrenia. Proceedings of the 151st Annual Meeting of the American Psychiatric Association; 1998 May 30 - Jun 4; Toronto, Ontario, Canada 1998.

Cantillon M, Goldstein JM. Quetiapine fumarate reduces aggression and hostility in patients with schizophrenia. Proceedings of the 151st Annual Meeting of the American Psychiatric Association; 1998 May 30 - Jun 4; Toronto, Ontario, Canada 1998.

Chengappa KNR, Goldstein JM, Greenwood M, John V, Levine J. A post hoc analysis of the impact on hostility and agitation of quetiapine and haloperidol among patients with schizophrenia. *Clinical Therapeutics* 2003;**25**(2):530-41. [MEDLINE: 12749512]

Hellewell JSE, Cameron-Hands D, Cantillon M. Seroquel: evidence for efficacy in the treatment of hostility and aggression. *Schizophrenia Research* 1998;**29**(1-2):154.

Hellewell JSE, Goldstein J. Seroquel (quetiapine): efficacy in improving mood, aggression and hostility. Proceedings of the 9th Congress of the Association of European Psychiatrists; 1998 Sep 20-24; Copenhagen, Denmark. 1998.

Hong W, Arvanitis L. Reduction of the positive symptoms of schizophrenia by 'ICI 204,636': results from phase II and III clinical trials. Proceedings of the 9th European College of Neuropsychopharmacology Congress; 1996 Sep 21-25; Amsterdam, Netherlands. 1996.

Alexander 2004 {published data only}

Alexander J. Lorazepam versus a combination of haloperidol and promethazine in the acute management of agitation and aggression - a randomized controlled trial [thesis]. Guindy, Chennai: Tamilnadu Dr. MGR Medical University, 2003.

Alexander J, John T, Tharyan P, Adams CE. TREC-India. A second arm of TREC. *Schizophrenia Research* 2002;**53**(3 Suppl 1):236.

* Alexander J, Tharyan P, Adams C, John T, Mol C, Philip J. Rapid tranquillisation of violent or agitated patients in a psychiatric emergency setting. Pragmatic randomised trial of intramuscular lorazepam v haloperidol plus promethazine. *British Journal of Psychiatry* 2004;**185**:63-9.

NCT00455234. Rapid tranquilization of violent or agitated people in psychiatric emergency settings- a pragmatic randomized controlled trial of intramuscular olanzapine vs intramuscular haloperidol + promethazine. <http://www.clinicaltrials.gov> 2007.

Raveendran NS. Rapid tranquillization of acutely agitated patients: intramuscular olanzapine vs haloperidol + promethazine - pragmatic randomized control trial. Proceedings of the 5th European Congress on Violence in Clinical Psychiatry; 2007 Oct 25-27; Amsterdam, The Netherlands. 2007.

Raveendran NS, Tharyan P, Alexander J, Adams CE. Rapid tranquillisation in psychiatric emergency setting in India: pragmatic randomised controlled trial of intramuscular olanzapine versus intramuscular haloperidol plus promethazine. *BMJ* 2007;**335**(7625):865-9. [MEDLINE: 17954514]

Tharyan P. A randomised controlled trial of intra-muscular lorazepam versus intra-muscular haloperidol+promethazine in the management of psychotic agitations and aggression. <http://www.controlled-trials.com> 2002.

Allen 2007 {published data only}

Allen M, Talbott S, Eudicone J, McQuade R, Pikalov A. Similar rates of agitation, anxiety and insomnia in sedating and non-sedating antipsychotics: evaluating clinical trial results with aripiprazole, haloperidol and olanzapine. *Schizophrenia Bulletin* 2007;**33**(2):418.

Asadollahi 2015 {published data only}

Asadollahi S, Heidari K, Hatamabadi H, Vafaee R, Yunesian S, Azadbakht A, et al. Efficacy and safety of valproic acid versus haloperidol in patients with acute agitation: results of a randomized, double-blind, parallel-group trial. *International Clinical Psychopharmacology* 2015; Vol. 30, issue 3:142-50.

Beasley 1998 {published data only}

Beasley CM, Sayler ME, Keisler GM, Potvin JH, Sanger TM, Tollefson GD. The influence of pharmacotherapy on self-directed and externally-directed aggression in schizophrenia. *Schizophrenia Research* 1998;**29**(1-2):28.

Bieniek 1998 {published data only}

Bieniek SA, Ownby RL, Penalver A, Dominguez RA. A double-blind study of lorazepam versus the combination of haloperidol

and lorazepam in managing agitation. *Pharmacotherapy* 1998;**18**(1):57-62.

Brook 1998 {published data only}

Brook S, Berk M, Selemani S, Kolloori J, Nzo I. A randomized controlled double blind study of zuclopenthixol acetate compared to haloperidol in acute psychosis. *Human Psychopharmacology* 1998;**13**(1):17-20.

Campbell 1982 {published data only}

Campbell M, Cohen IL, Small AM. Drugs in aggressive behavior. *Journal of the American Academy of Child Psychiatry* 1982;**21**(2):107-17. [MEDLINE: 7040529]

Chen 2004 {published data only}

Chen M-F, Liu D-X, Lin J-Z. The comparison study on curing agitation by haloperidol and chlorpromazine venoclysis [氟哌啶醇与氯丙嗪静滴治疗急性激越的对照研究]. *Medical Journal of Chinese Civil Administration* 2004;**16**(10):599.

Chen 2008 {published data only}

Chen J, Yang X, Song A, Gu X, Ma X. Intramuscular ziprasidone versus haloperidol in the treatment of acute agitation in schizophrenia [齐拉西酮注射液与氟哌啶醇注射液治疗精神分裂症急性激越症状的对比研究]. *Shanghai Archives of Psychiatry* [#####] [#####] 2008;**20**(6):363-6.

Chouinard 1991 {published data only}

Chouinard G, Annable L, Turnier L, Holobow N, Szkrumelak N. A double-blind randomized clinical trial of rapid tranquilization with IM clonazepam and IM haloperidol in agitated psychotic patients with manic symptoms. *Canadian Journal of Psychiatry [Revue Canadienne de Psychiatrie]* 1993;**38**(Suppl 4):S114-21. [MEDLINE: 8306241]

* Chouinard G, Safadi G, Beauclair L. The management of acutely schizophrenic patients newly admitted from the emergency room: A double-blind clinical trial comparing zuclopenthixol acetate and liquid haloperidol. Proceedings of the 5th World Congress of Biological Psychiatry; 1991 Jun 9-14; Florence, Italy. Amsterdam, Netherlands: Excerpta Medica, 1991:35-43.

Citrome 2001 {published data only}

* Citrome L, Volavka J, Czobor P, Sheitman B, Lindenmayer J-P, McEvoy J, et al. Effects of clozapine, olanzapine, risperidone, and haloperidol on hostility among patients with schizophrenia. *Psychiatric Services* 2001;**52**(11):1510-4. [MEDLINE: 11684748]

Citrome LL, Volavka J, Czobor P, Nolan K, Lieberman JA, Lindenmayer J-P, et al. Overt aggression and psychotic symptoms in patients with schizophrenia treated with clozapine, olanzapine, risperidone, or haloperidol. Proceedings of the 156th Annual Meeting of the American Psychiatric Association; 2003 May 17-22; San Francisco, California, USA. 2003.

Citrome LL, Volavka J, Czobor P, Sheitman BB, Lindenmayer J-P, McEvoy JP, et al. Atypical antipsychotics and hostility in schizophrenia: A double-blind study. Proceedings of the 154th Annual Meeting of the American Psychiatric Association; 2001 May 5-10; New Orleans, Louisiana, USA. 2001.

Citrome LL, Volavka J, Czobor P, Sheitman BB, Lindenmayer J-P, McEvoy JP, et al. Atypical antipsychotics and hostility in schizophrenia: A double-blind study. Proceedings of the 155th Annual Meeting of the American Psychiatric Association; 2002 May 18-23; Philadelphia, Pennsylvania, USA. 2002.

Hoptman MJ, Volavka J, Czobor P, Gerig G, Chakos M, Blocher J, et al. Aggression and quantitative MRI measures of caudate in patients with chronic schizophrenia or schizoaffective disorder. *Journal of Neuropsychiatry and Clinical Neurosciences* 2006;**18**(4):509-15.

Volavka J, Czobor P, Nolan K, Sheitman B, Lindenmayer JP, Citrome LMJP, et al. Overt aggression and psychotic symptoms in patients with schizophrenia treated with clozapine, olanzapine, risperidone, or haloperidol. *Journal of Clinical Psychopharmacology* 2004;**24**(2):225-8. [MEDLINE: 15206671]

Colonna 1998 {published data only}

Colonna L, Martin P, Gerard D. Long-term study of the effectiveness and tolerance of amisulpride versus haloperidol [Etude à long terme de l'efficacité et de la tolérance à l'amisulpride versus haloperidol]. Proceedings of the 96th Congress de Psychiatrie et de Neurologie de Langue Française; 1998 May 10-15; Saint Paul, France. 1998.

Conde 2011 {published data only}

Conde BL, Sobreveg EE, Sionzon MP. A randomized trial of oral risperidone versus intramuscular haloperidol in the emergency treatment of acute psychotic agitation. *ASEAN Journal of Psychiatry* 2011; Vol. 12, issue 1:29-36.

Crandall 2005 {published data only}

Crandall D, Pikalov A, Kostic D, Kaplita S, Berman R, McQuade R, et al. Is sedation needed for effective reduction of acute schizophrenia symptoms?. Proceedings of the 158th Annual Meeting of the American Psychiatric Association; 2005 May 21-26; Atlanta, Georgia, USA. 2005.

Currier 2000 {published data only}

Currier GW, Simpson GA. Risperidone oral solution versus intramuscular haloperidol for control of psychotic agitation. *European Neuropsychopharmacology* 2000;**10**(Suppl 3):S296.

Daniel 2000 {published data only}

Daniel DG, Brook S, Reeves KR. An overview of the efficacy and safety of rapid acting intramuscular ziprasidone. *European Neuropsychopharmacology* 2000;**10**(Suppl 3):S289.

Daniel 2003 {published data only}

Daniel DG, Brook S, Benattia I. Efficacy of intramuscular ziprasidone without adjunctive benzodiazepines. *European Neuropsychopharmacology* 2003;**13**(4):S299.

Daniel 2004 {published data only}

* Daniel DG, Zimbroff DL, Swift RH, Harrigan EP. The tolerability of intramuscular ziprasidone and haloperidol treatment and the transition to oral therapy. *International Clinical Psychopharmacology* 2004;**19**(1):9-15.

Lucey JV, Brook S, Daniel DG, Reeves KR, Harrigan EP. Intramuscular (IM) ziprasidone: a novel treatment for the short

term management of agitated psychotic patients. *Schizophrenia Research* 2000;**41**(1):208.

Modell S, Daniel D, Stock E, Wilber R, Marcus R, Carson W, et al. Intramuscular aripiprazole treatment for acute agitation in patients with psychosis. Proceedings of the 24th Collegium Internationale Neuro-Psychopharmacologicum Congress; 2004 June 20–24; Paris, France 2004.

Swift RH, Harrigan EP, Cappelleri JC, Kramer D, Chandler LP. Validation of the behavioural activity rating scale (BARS): a novel measure of activity in agitated patients. *Journal of Psychiatric Research* 2002;**36**(2):87–95. [MEDLINE: 11777497]

Swift RH, Harrigan EP, Kammen DP. A comparison of intramuscular (IM) ziprasidone with IM haloperidol. Proceedings of the 9th Congress of the Association of European Psychiatrists; 1998 Sep 20–24; Copenhagen, Denmark. 1998.

Swift RH, Harrigan EP, Kammen DP. A comparison of intramuscular ziprasidone with intramuscular haloperidol. Proceedings of the 151st Annual Meeting of the American Psychiatric Association; 1998 May 30 - Jun 4; Toronto, Ontario, Canada 1998.

Swift RH, Harrigan EP, Van Kammen D, Casey DE. A comparison of fixed dose, intramuscular (IM) ziprasidone with flexible dose, IM haloperidol. Proceedings of the 11th European College of Neuropsychopharmacology Congress; 1998 Oct 31 - Nov 4; Paris, France. 1998.

Swift RH, Harrigan EP, Van-Kammen D. A comparison of fixed-dose intramuscular (IM) ziprasidone with flexible-dose IM haloperidol. Proceedings of the 21st Collegium Internationale Neuro-Psychopharmacologicum Congress; 1998 Jul 12–16; Glasgow, UK. 1998.

Yocca F, Daniel D, Stock E, Oren D, Marcus R, Carson W, et al. Efficacy and safety of intramuscular aripiprazole treatment for acute agitation in patients with psychosis. Proceedings of the 43rd Annual Meeting of the American College of Neuro-Psychopharmacology; 2004 Dec 12–16; San Juan, Puerto Rico. 2004.

Fan 2012 {published data only}

樊凌姿, 曹立新. 氯普噻吨注射液与氟哌啶醇注射液治疗精神分裂症兴奋激越对照研究. 中外健康文摘 2012; Vol. 40:31–2.

Faretra 1970 {published data only}

Faretra G, Dooher L, Dowling J. Comparison of haloperidol and fluphenazine in disturbed children. *American Journal of Psychiatry* 1970;**126**:1670–3. [MEDLINE: 5443653]

Freeman 2009 {published data only}

Freeman DJ, DiPaula BA, Love RC. Intramuscular haloperidol versus intramuscular olanzapine for treatment of acute agitation: A cost-minimization study. *Pharmacotherapy* 2009;**29**(8):930–6.

Goldstein 1966 {published data only}

Goldstein BJ, Clyde DJ. Haloperidol in controlling the symptoms of acute psychoses. Part II: A double blind evaluation of

haloperidol and trifluoperazine. *Current Therapeutic Research* 1966;**8**:236–40.

Harvey 2004 {published data only}

Harvey AT, Flockhart D, Gorski JC, Greenblatt DJ, Burke M, Werder S, et al. Intramuscular haloperidol or lorazepam and QT intervals in schizophrenia. *Journal of Clinical Pharmacology* 2004;**44**(10):1173–84. [MEDLINE: 15342619]

Huang 2004 {published data only}

Huang W-S, Di X-L, Yang F-D. A comparative study of the effect of modern electroconvulsive therapy on the agitated behavior of psychotic patients. *Journal of Clinical Psychological Medicine* [#####] 2004;**14**(5):276–7.

Jiang 2009 {published data only}

蒋合萍, 罗毅平. Olanzapine and haloperidol in the treatment of schizophrenia comparative study of excited agitation [Google Translate] [奥氮平与氟哌啶醇治疗精神分裂症兴奋激越症状的对照研究]. *Sichuan Mental Health* [四川精神卫生] 2009; Vol. 22, issue 1:43–4.

Jones 2001 {published data only}

Jones B, Taylor CC, Meehan K. The efficacy of a rapid-acting intramuscular formulation of olanzapine for positive symptoms. *Journal of Clinical Psychological Medicine* [#####] 2001;**62**(2):22–4. [MEDLINE: 11232747]

Kane 2000 {published data only}

Kane J, Ingenito G, Ali M. Efficacy of aripiprazole in psychotic disorders: comparison with haloperidol and placebo. *Schizophrenia Research* 2000;**41**(1):39.

Kinon 2001a {published data only}

Eli Lilly. Beasley 1996a - olz vs placebo vs hpl (N America) (as supplied 2004). Data on file 2001.

Eli Lilly. Beasley 1997 - olz vs hpl (international) (as supplied 2004). Data on file 2001.

Eli Lilly. Tollefson 1997 - olz vs hpl (as supplied 2004). Data on file 2001.

Kinon BJ, Basson MS, Tollefson GD. Gender-specific prolactin response to treatment with olanzapine versus haloperidol in schizophrenia. Proceedings of the 9th Biennial Winter Workshop on Schizophrenia; 1998 Feb 7–13; Davos, Switzerland 1998.

Kinon BJ, Milton DR, Hill AL, Williamson DJ. Effective resolution of acute presentation of behavioral agitation and positive psychotic symptoms in schizophrenia with olanzapine. *Journal of Psychopharmacology* 2000;**14**(3 Suppl):A60.

* Kinon BJ, Roychowdhury SM, Milton DR, Hill AL. Effective resolution with olanzapine of acute presentation of behavioral agitation and positive psychotic symptoms in schizophrenia. *Journal of Clinical Psychiatry* 2001;**62**(Suppl 2):17–21. [MEDLINE: 11232746]

Kinon BJ, Wang L, Stauffer VL. A comparison of prolactin concentrations in the elderly during treatment with olanzapine, haloperidol, and risperidone. Proceedings of the 14th Annual

Meeting of the American Association for Geriatric Psychiatry; 2001 Feb 23-26; San Francisco, California, USA. 2001.

Kinon 2003 {published data only}

* Kinon BJ, Stauffer VL, McGuire HC, Kaiser CJ, Dickson RA, Kennedy JS. The effects of antipsychotic drug treatment on prolactin concentrations in elderly patients. *Journal of the American Medical Directors Association* 2003;**4**(4):189-94. [PUBMED: 12837139]

Kinon BJ, Wang L, Stauffer VL. A comparison of prolactin concentrations in the elderly during treatment with olanzapine, haloperidol, and risperidone. Proceedings of the 14th Annual Meeting of the American Association for Geriatric Psychiatry; 2001 Feb 23-26; San Francisco, California, USA. 2001.

Krakowski 2006 {published data only}

Krakowski M I, Czobor P. Depression and impulsivity as pathways to violence: implications for antiaggressive treatment. *Schizophrenia Bulletin* 2014; Vol. 40, issue 4:886-94.

Krakowski M, Czobor P. Cholesterol and cognition in schizophrenia: A double-blind study of patients randomized to clozapine, olanzapine and haloperidol. *Schizophrenia Research* 2011; Vol. 130, issue 1-3:27-33.

Krakowski M, Czobor P, Citrome L. Weight gain, metabolic parameters, and the impact of race in aggressive inpatients randomized to double-blind clozapine, olanzapine or haloperidol. *Schizophrenia Research* 2009;**110**(1-3):95-102. [MEDLINE: 19269139]

Krakowski MI. Clozapine and olanzapine in violent schizophrenics. CRISP database (<https://www-commons.cit.nih.gov/crisp/index.html>) (accessed 19th February 2001).

Krakowski MI, Czobor P. A prospective longitudinal study of cholesterol and aggression in patients randomized to clozapine, olanzapine, and haloperidol. *Journal of Clinical Psychopharmacology* 2010;**30**(2):198-200.

Krakowski MI, Czobor P. Executive function predicts response to antiaggression treatment in schizophrenia: A randomized controlled trial. *Journal of Clinical Psychiatry* 2012; Vol. 73, issue 1:74-80.

Krakowski MI, Czobor P. Neurocognitive impairment limits the response to treatment of aggression with antipsychotic agents. *Schizophrenia Bulletin* 2011; Vol. 37:311-2.

* Krakowski MI, Czobor P, Citrome L, Bark N, Cooper TB. Atypical antipsychotic agents in the treatment of violent patients with schizophrenia and schizoaffective disorder. *Archives of General Psychiatry* 2006;**63**(6):622-9. [MEDLINE: 16754835]

NCT01123408. Clozapine and Olanzapine Treatment of Aggression. <http://ClinicalTrials.gov/ct2/show/NCT01123408> 2010.

Krakowski 2008 {published data only}

Krakowski M, Czobor P. A prospective longitudinal study of cholesterol and aggression in patients randomized to clozapine, olanzapine and haloperidol. Proceedings of the 15th Biennial Winter Workshop in Psychoses; 2009 Nov 15-18; Barcelona, Spain. 2009.

* Krakowski MI, Czobor P, Nolan KA. Atypical antipsychotics, neurocognitive deficits, and aggression in schizophrenic patients. *Journal of Clinical Psychopharmacology* 2008;**28**(5):485-93.

Nolan KA, Volavka J, Czobor P, Sheitman B, Lindenmayer J-P, Citrome LL, et al. Aggression and psychopathology in treatment-resistant inpatients with schizophrenia and schizoaffective disorder. *Journal of Psychiatric Research* 2005;**39**(1):109-15. [MEDLINE: 15504429]

Lewis 2006 {published data only}

* Lewis CF. Haloperidol versus risperidone in the treatment of aggressive psychotic male inmates. Proceedings of the 159th Annual Meeting of the American Psychiatric Association; 2006 May 20-25; Toronto, Canada. 2006.

NCT00203775. Efficacy of risperidone versus haloperidol in the treatment of aggression and hostility in psychotic inmates. <http://www.clinicaltrials.gov> 2005.

Li 2005 {published data only}

Li GL. Tiapride intramuscular haloperidol in the treatment of acute schizophrenia excited state control study [硫必利肌注与氟哌啶醇治疗精神分裂症急性期兴奋状态的对照研究]. *Nervous Diseases and Mental Health* [神经疾病与精神卫生] 2005; Vol. 5, issue 6:456-7.

Li 2007 {published data only}

Li C, Zhu S, Wang H, Chen H, Yu Y, Liu D, et al. Safety and efficacy of clonazepam, haloperidol and haloperidol combined with clonazepam in the treatment of schizophrenia with excitement and agitation [氯硝西洋、氟哌啶醇及其联合治疗精神分裂症激越症状疗效的比较]. *Shanghai Archives of Psychiatry* [#####] [#####] 2007;**19**(3):150-2.

Liang 2003 {published data only}

梁芝国, 罗维肖. Clonazepam and haloperidol on patients with acute agitation control study [氯硝西洋和氟哌啶醇联用治疗急性激越的对照研究]. *Medical Journal of Chinese Civil Administration* 2003; Vol. 15, issue 5:293-4.

Liu 2012b {published data only}

刘庆梅. 无抽搐电休克治疗精神分裂症伴激越行为疗效观察及护理干预. *临床心身疾病杂志* 2012; Vol. 18, issue 5:437-9.

Mandel 2008 {published data only}

Mandel F, Allen MH, Lombardo I. Early response to intramuscular ziprasidone as a predictor of end point response to oral ziprasidone. Proceedings of the 161st Annual Meeting of the American Psychiatric Association; 2008 May 3-8; Washington DC, USA. 2008.

Mantua 2006 {published data only}

Mantua V, Travis MJ, Atakan Z, Isaac MB, Isaac MT, Smith S, et al. Treatment of agitation caused by severe mental illness: data from the South London and Maudsley intensive care units trial evaluation (SLAMICUTE) study. *European Neuropsychopharmacology* 2006;**16**(Suppl 4):S420.

McEvoy 1994 {published data only}

McEvoy JP. Efficacy of risperidone on positive features of schizophrenia. *Journal of Clinical Psychiatry* 1994;**55**(5 Suppl):18-21. [MEDLINE: 7520904]

Mendelowitz 2004 {published data only}

Mendelowitz AJ. The utility of intramuscular ziprasidone in the management of acute psychotic agitation. *Annals of Clinical Psychiatry* 2004;**16**(3):145-54. [MEDLINE: 15517847]

NCT00189995 2005 {published data only}

NCT00189995. Intramuscular clozapine in the management of aggression in schizophrenic patients. <http://www.clinicaltrials.gov> 2005.

NCT00631722 2008 {published data only}

NCT00631722. Multicenter, open-label, randomised, haloperidol-controlled study to evaluate seroquel as mono-therapy in the treatment of agitated symptoms in the patients with acute episode of schizophrenia. <http://www.clinicaltrials.gov> 2008.

NCT00797277 2008 {published data only}

NCT00797277. IM olanzapine versus IM haloperidol plus lorazepam for acute agitation in schizophrenia. <http://www.clinicaltrials.gov> 2008.

Pan 2005 {published data only}

Pan N, Yang X, Mei Q. Large dose of olanzapine in the ictus treatment of agitated symptoms of schizophrenia. *Journal of Clinical Psychosomatic Diseases* [#####] 2005;**11**(2):113-5.

Pathiraja 1995 {published data only}

Pathiraja AP, Diamond BI, Borison RL, Meibech RC, Anand R. Relationship between creatine phosphokinase, psychotic symptoms and novel antipsychotic drugs. *Schizophrenia Research* 1995;**15**(1-2):160-1.

Pedros 2004 {published data only}

* Pedros Rosello A, Tenias JM. Neuroleptics election and clinical outcomes in acute psychosis: A comparative study of risperidone versus other neuroleptics [Eleccion de neuroleptico y respuesta clinica en psicosis aguda: Un estudio comparativo de risperidona frente a otros neurolepticos]. *Anales de Psiquiatria* 2004;**20**(4):167-71.

Pei 2009 {published data only}

裴树景, 杨红卫. Oral risperidone treatment of schizophrenia clinical observation of excited agitation [Google Translate] [利培酮口服液治疗精神分裂症兴奋激越临床观察]. *Linchuang Jingshen Yixue Zazhi* [临床精神医学杂志] 2009; Vol. 19, issue 6:405-6.

Puech 1998 {published data only}

Muller MJ, Wetzel H, Eich FX, Rein W, Puech A, Benkert O, Amisulpride Study Group. Dose-related effects of amisulpride on five dimensions of psychopathology in patients with acute exacerbation of schizophrenia. *Journal of Clinical Psychopharmacology* 2002;**22**(6):554-60.

* Puech A, Fleurot O, Rein W, Amisulpride Study Group. Amisulpride, an atypical antipsychotic, in the treatment of acute episodes of schizophrenia: a dose-ranging study vs. haloperidol. *Acta Psychiatrica Scandinavica* 1998;**98**(1):65-72. [MEDLINE: 9696517]

Romeo 2009 {published data only}

* Romeo R, Knapp M, Tyrer P, Crawford M, Oliver-Africano P. The treatment of challenging behaviour in intellectual disabilities: cost-effectiveness analysis. *Journal of Intellectual Disability Research* 2009;**53**:633-43.

Tyrer P. Neuroleptics in adults with aggressive challenging behaviour and intellectual disability. <http://www.controlled-trials.com> 2003.

Simpson 2010 {published data only}

Simpson G, Mohammad AR, Fauzia S, Campaneau M, Laska E, Ninah R, et al. A comparison of rapid titration quetiapine and haloperidol in agitated adults in an emergency setting. Proceedings of the 163rd Annual Meeting of the American Psychiatric Association; 2010 May 22-26; New Orleans, LA 2010.

Singh 1981 {published data only}

Singh MM, Kay SR, Pitman RK. Aggression control and structuring of social relations among recently admitted schizophrenics. *Psychiatry Research* 1981;**5**(2):157-69. [MEDLINE: 6945611]

Smith 1974 {published data only}

Smith GR, Taylor CW, Linkous P. Haloperidol versus thioridazine for the treatment of psychogeriatric patients: A double-blind clinical trial. *Psychosomatics* 1974;**15**(3):134-8.

Srinath 2010 {published data only}

Srinath G, Shailaja B, Sai PG, Reddy KA. A comparative study of injection haloperidol and promethazine vs. injection lorazepam in controlling acute psychotic agitation. *Indian Journal of Psychiatry* 2010; Vol. 52:S21.

Teja 1975 {published data only}

Teja JS, Grey WH, Clum JM, Warren C. Tranquilizers or anti-depressants for chronic schizophrenics: A long term study. *Australian and New Zealand Journal of Psychiatry* 1975;**9**(4):241-7.

Thomas 1992 {published data only}

Thomas HJ, Schwartz E, Petrilli R. Droperidol versus haloperidol for chemical restraint of agitated and combative patients. *Annals of Emergency Medicine* 1992;**21**(4):407-13. [MEDLINE: 1554179]

TREC CG {published data only}

Coutinho ES, Huf G, Allen MH, Adams CE. Physical restraints for agitated patients in psychiatric emergency hospitals in Rio de Janeiro, Brazil: a predictive model. *Schizophrenia Bulletin* 2005;**31**:220.

Huf G. Rapid safe tranquillisation for acutely disturbed people attending public psychiatric emergency clinics in Rio de Janeiro. <http://www.controlled-trials.com> 2002.

Huf G, Coutinho ESF, Adams CE. TREC I. Background. *Schizophrenia Research* 2002;**53**(3 Suppl 1):187.

Huf G, Coutinho ESF, Adams CE. TREC III. The protocol and progress of TREC. *Schizophrenia Research* 2002;**53**(3 Suppl 1):187.

Huf G, Coutinho ESF, Adams CE. TREC-Rio trial: A randomised controlled trial for rapid tranquillisation for agitated patients in emergency psychiatric rooms. *BMC Psychiatry* 2002;**2**(11):1-12.

Huf G, Coutinho ESF, Fagundes HM Jr, Carvalho AL, Ramos FA, Keusen AL, Oliveira ES, et al. TREC II. Current practices in managing acutely disturbed patients at three hospitals in Rio de Janeiro, Brazil. *Schizophrenia Research* 2002;**53**(3):236-7.

* TREC CG. Rapid tranquillisation for agitated patients in emergency psychiatric rooms: A randomised trial of midazolam versus haloperidol plus promethazine. *BMJ* 2003;**327**(7417):708-13.

Ungvari 1982 {published data only}

Ungvari G, Petho B. High-dose haloperidol therapy: Its effectiveness and a comparison with electroconvulsive treatment. *Journal of Psychiatric Treatment and Evaluation* 1982;**4**:279-83.

Vaisanen 1981 {published data only}

Vaisanen K, Viukari M, Rimon R, Raisanen P. Haloperidol, thioridazine and placebo in mentally subnormal patients-serum levels and clinical effects. *Acta Psychiatrica Scandinavica* 1981;**63**(3):262-71. [MEDLINE: 7015792]

Veser 2006 {published data only}

Veser F, Zealburg J, Veser B, Zhu Y, Gharabawi G. Oral risperidone in the management of agitated behavior in emergency settings. *European Neuropsychopharmacology* 2002;**12**(Suppl 3):S313.

* Veser FH, Veser BD, McMullan JT, Zealberg J, Currier GW. Risperidone versus haloperidol, in combination with lorazepam, in the treatment of acute agitation and psychosis: a pilot, randomized, double-blind, placebo-controlled trial. *Journal of Psychiatric Practice* 2006;**12**(2):103-8. [MEDLINE: 16728906]

Vives 2015 {published data only}

Vives Vilagut R, Duno R, Planet N, Garcia G, Payes M, Marinosa M, et al. Testing the intranasal route for administration of haloperidol in emergency room: Generation of efficacy data to support clinical practice. *Clinical Therapeutics* 2015; Vol. 1:e152.

Wan 2005 {published data only}

Wan Z, Zhong Z. Curative effect of risperidone with BDZs in the treatment of excitement and agitate with schizophrenia in acute phase. *Heath Psychology Journal* 2005;**13**(1):23-4.

Wang 2004 {published data only}

Wang G, Cai Z-J, Wang L-F. A multicenter study of risperidone treatment for acute agitation in patients with schizophrenia. *Chinese Journal of Psychiatry* 2004;**37**(2):88-91.

Wang 2005 {published data only}

Wang H, Guo X, Shen T. Safety and efficacy of quetiapine and haloperidol in the treatment of schizophrenia with excitement and agitation. *Shanghai Archives of Psychiatry* [#####] [#####] 2005;**17**(5):271-4.

Wyant 1990 {published data only}

Wyant M, Diamond BI, O'Neal E, Sloan A, Borison RL. The use of midazolam in acutely agitated psychiatric patients. *Psychopharmacology Bulletin* 1990;**26**(1):126-9.

Zapletalek 1986 {published data only}

Zapletalek M, Tuma I, Libiger J. Zetidoline compared with haloperidol in a double blind clinical study. *Activitas Nervosa Superior* 1986;**28**(1):31-2.

References to studies awaiting assessment
Daniel 2006 {published data only}

Daniel DG, Crandall D, Manos G, McQuade RD, Gutierrez-Esteinou R, Pikalov AA, et al. Transitioning from intramuscular (IM) to oral aripiprazole in patients with schizophrenia. Proceedings of the 159th Annual Meeting of the American Psychiatric Association; 2006 May 20-25; Toronto, Canada. 2006.

Davis 2008 {published data only}

Davis JM, Wang B, He Y, Jin H, Hu Q, Zhang M. The emergency treatment of acutely agitated psychotic schizophrenia patient. *International Journal of Neuropsychopharmacology* 2008;**11**(Suppl 1):163.

Dubin 1985 {published data only}

Dubin WR, Waxman HM, Weiss KJ, Ramchandani D, Tavani-Petrone C. Rapid tranquilization: the efficacy of oral concentrate. *Journal of Clinical Psychiatry* 1985;**46**(11):475-8. [MEDLINE: 2865251]

Esmailian 2015 {published data only}

Esmailian M, Ahmadi O, Taheri M, Zamani M. Comparison of haloperidol and midazolam in restless management of patients referred to the Emergency Department: PDF: A double-blinded, randomized clinical trial. *Journal of Research in Medical Sciences* 2015; Vol. 20, issue 9:844-9.

Herrera 2005 {published data only}

Herrera M. Double-blind study with risperidone vs haloperidol in schizophrenic patients with agitation and/or aggression. Proceedings of the 8th World Congress of Psychiatry; 2005 Sep 10-15; Cairo, Egypt. 2005.

Hsu 2010 {published data only}

Hsu WY, Huang SS, Lee BS, Chiu NY. Comparison of intramuscular olanzapine, orally disintegrating olanzapine tablets, oral risperidone solution, and intramuscular haloperidol in the management of acute agitation in an acute care psychiatric ward in Taiwan. *Journal of Clinical Psychopharmacology* 2010;**30**(3):230-4.

Istikoglou 2010 {published data only}

Istikoglou C, Vlavianou A, Foutsitzis D, Theodorakopoulos G, Polonifis C, Kanellos P, et al. Intramuscular aripiprazole versus injectable haloperidol in treatment of acute psychotic agitation. *European Neuropsychopharmacology* 2010;**20**(3):S469-70.

Kong 2009 {published data only}

Kong BG, Shim JC, Lee SK. Comparison of the effects of intramuscular antipsychotics injection in the treatment of acute agitation in psychosis. *European Neuropsychopharmacology* 2009; Vol. 19:S525-S6.

Lasic 2006 {published data only}

Lasic D, Rubesa G, Dodig G, Katavic Z, Glavina T, Zuljan-Cvitanovic M. Risperidone oral solution, therapeutic possibility in treatment of acute and agitated psychotic patients. *European Neuropsychopharmacology* 2006;**16**(Suppl 4):S445.

Li 2007a {published data only}

Li C, Zhu S, Wang H, Chen H, Yu Y, Liu D, et al. Safety and efficacy of clonazepam, haloperidol and haloperidol combined with clonazepam in the [氯硝西泮、氟哌啶醇及其联合治疗对精神分裂症急性激越症状疗效的比较]. *Shanghai Archives of Psychiatry [上海精神医学]* 2007; Vol. 19, issue 3:150-2.

Lu 2006 {published data only}

Lu D, Ning J, Zhong X. Clinical evaluation on treatment of acute-agitation in schizophrenia with ziprasidone injection [甲磺酸齐拉西酮注射液治疗精神分裂症急性激越症状的临床评价]. *China Pharmaceuticals* 2006;**15**(4):48-9.

NCT00859872 2009 {published data only}

NCT00859872. Efficacy and safety of risperidone oral solution combination clonazepam versus haloperidol intramuscular (IM) injection for treatment of acute psychotic agitation in schizophrenia. <http://www.clinicaltrials.gov> 2009.

NCT00866645 2009 {published data only}

NCT00866645. A study to evaluate the efficacy and safety of intramuscular levosulpiride in patients with agitation of schizophrenia. <http://www.clinicaltrials.gov> 2009.

Shim 2009 {published data only}

Shim JC, Kong BG, Lee SG, Kim YH. The effect of intramuscular olanzapine in the treatment of acute agitation in patient with psychotic symptoms. *European Neuropsychopharmacology* 2009; Vol. 19:S512-S3.

Si 2009 {published data only}

Si T, Zhong HJ, Yun'Ai S, Qin TM, Qing LK, De YP, et al. A multi-site, open-label, randomised, haloperidol-controlled study to evaluate quetiapine as monotherapy in the treatment of agitated acute schizophrenia. *Proceedings of the 162nd Annual*

Meeting of the American Psychiatric Association; 2009 May 16-21; San Francisco, CA 2009.

Slotnick 1971 {published data only}

Slotnick VB. Management of the acutely agitated psychiatric patient with parental neuroleptics: The comparative symptom effectiveness profiles of haloperidol and chlorpromazine. *Proceedings of the 5th World Congress of Psychiatry*; 1971 Nov 28 - Dec 4; Mexico City, Mexico. 1971.

Smythies 1982 {published data only}

Smythies JR, Hain J, Crews E, Grayson G. A clinical trial of injectable molindone (Moban(R)). *Current Therapeutic Research, Clinical and Experimental* 1982;**32**(5):752-6.

Vasquez-Gomez 2001 {published data only}

Vasquez-Gomez F. The use of midazolam for control of acutely agitated outpatients in psychiatry emergency. *Proceedings of the 7th World Congress of Biological Psychiatry*; 2001 Jul 1-6; Berlin, Germany. 2001.

Wang 2005a {published data only}

Wang R-C, Li L-H, Zhao Y-C. Ziprasidone made in China and haloperidol in treating acute onset of schizophrenia: a randomized, single blinded, double modelling, parallel control study. *Chinese Journal of Clinical Rehabilitation* 2005; Vol. 9, issue 44:83-5.

Wang 2007 {published data only}

Wang J-C, Xu X-F, Ouyang H, Li W-Y. The second phase clinical trials of intramuscular ziprasidone for the treatment of acute psychotic agitation of the schizophrenia [注射用甲磺酸齐拉西酮治疗精神分裂症兴奋躁动状态的二期临床试验]. *Nervous Diseases and Mental Health [#####]* 2007;**7**(5):364-70.

Yildiz 2003 {published data only}

Yildiz A, Turgay A, Alpay M, Sachs GS. Risperidone versus haloperidol in adult psychiatric emergencies. *Proceedings of the 156th Annual Meeting of the American Psychiatric Association*; 2003 May 17-22; San Francisco, California, USA 2003:Nr259.

Zhang 2012 {published data only}

张庆娥, 王刚, 张玲, 王雪, 罗炯, 路亚洲, et al. Control study of acute agitation in schizophrenia risperidone oral solution merger clonazepam tablets treatment [Google Translate] [利培酮口服液合并氯硝西泮片治疗精神分裂症急性激越的对照研究]. *中华医学会第十次全国精神医学学术会议* 2012; Vol. 22, issue 02:89-91.

叶萌, 2009 {published data only}

叶萌, 房茂胜, 李乐华, 赵靖平. Domestic ziprasidone mesylate injection in the treatment of acute agitation of schizophrenia [国产甲磺酸齐拉西酮注射液治疗精神分裂症急性激越症状疗效观察]. *Journal of the Xinxiang Medical College [新乡医学院学报]* 2009; Vol. 26, issue 1:63-5.

周德祥, 2012 {published data only}

周德祥, 蒋幸衍, 徐清, 方馨怡, 夏鸣华, 陆雅娜. 利培酮口服液合并氯硝西泮治疗精神分裂症兴奋躁动症状的随机对照研究. *中国健康心理学杂志* 2012; Vol. 8:1123-4.

References to ongoing studies

NCT00838032 2009 {published data only}

NCT00838032. Pilot study to evaluate the efficacy and safety of quetiapine fumarate instant-release (seroquel ir) in controlling agitation and aggressive symptoms in the acute treatment of patients with schizophrenia. <http://www.clinicaltrials.gov> 2009.

Additional references

Adams 2013

Adams CE, Bergman H, Irving CB, Lawrie S. Haloperidol versus placebo for schizophrenia. *Cochrane Database of Systematic Reviews* 2013, Issue 11. [DOI: [10.1002/14651858.CD003082.pub3](https://doi.org/10.1002/14651858.CD003082.pub3); CD003082]

Ahmed 2010

Ahmed U, Jones H, Adams CE. Chlorpromazine for psychosis induced aggression or agitation. *Cochrane Database of Systematic Reviews* 2010, Issue 4. [DOI: [10.1002/14651858.CD007445.pub2](https://doi.org/10.1002/14651858.CD007445.pub2); PUBMED: 20393959]

Ahmed 2011

Ahmed U, Rehman F, Jones H, Adams CE. Risperidone for psychosis induced aggression or agitation. *Cochrane Database of Systematic Reviews* 2011, Issue 11. [DOI: [10.1002/14651858.CD009412](https://doi.org/10.1002/14651858.CD009412); CD009412]

Altman 1996

Altman DG, Bland JM. Detecting skewness from summary information. *BMJ* 1996;**313**(7066):1200.

APA 2006

American Psychiatric Association. Practice Guideline for the Treatment of Patients With Schizophrenia. 2nd Edition. Washington DC: American Psychiatric Association, 2006. [DOI: [10.1176/appi.books.9780890423363.45859](https://doi.org/10.1176/appi.books.9780890423363.45859)]

Barnes 1989

Barnes TR. A rating scale for drug-induced akathisia. *British Journal of Psychiatry* 1989;**154**:672-6.

Belgamwar 2005

Belgamwar RB, Fenton M. Olanzapine IM or velotab for acutely disturbed/agitated people with suspected serious mental illnesses. *Cochrane Database of Systematic Reviews* 2005, Issue 2. [DOI: [10.1002/14651858.CD003729.pub2](https://doi.org/10.1002/14651858.CD003729.pub2)]

Berk 2004

Berk M, Rathbone J, Mandriota-Carpenter SL. Clotiapine for acute psychotic illnesses. *Cochrane Database of Systematic Reviews* 2004, Issue 4. [DOI: [10.1002/14651858.CD002304.pub2](https://doi.org/10.1002/14651858.CD002304.pub2); CD002304]

Bland 1997

Bland JM. Statistics notes. Trials randomised in clusters. *BMJ* 1997;**315**:600.

BNF 2011

British National Formulary. British National Formulary. 61. NHS, London: Pharmaceutical Press, 2011.

Boissel 1999

Boissel JP, Cucherat M, Li W, Chatellier G, Gueyffier F, Buyse M, et al. The problem of therapeutic efficacy indices. 3. Comparison of the indices and their use [Aperçu sur la problématique des indices d'efficacité thérapeutique, 3: comparaison des indices et utilisation. Groupe d'étude des indices d'efficacité]. *Thérapie* 1999;**54**(4):405-11. [PUBMED: 10667106]

Choudhury 2011

Choudhury R, Dewsbery M, Williams K, Hovey N. Audit of the rapid tranquilization policy in the 2gether NHS foundation trust. *Journal of Psychiatric Intensive Care* 2011;**7**(1):55-61.

Corrigan 1989

Corrigan JD. Development of a scale for assessment of agitation following traumatic brain injury. *Journal of Clinical and Experimental Neuropsychology* 1989;**11**(2):261-77.

Deeks 2000

Deeks J. Issues in the selection for meta-analyses of binary data. Proceedings of the 8th International Cochrane Colloquium; 2000 Oct 25-28; Cape Town. Cape Town: The Cochrane Collaboration, 2000.

Divine 1992

Divine GW, Brown JT, Frazier LM. The unit of analysis error in studies about physicians' patient care behavior. *Journal of General Internal Medicine* 1992;**7**(6):623-9.

Donner 2002

Donner A, Klar N. Issues in the meta-analysis of cluster randomized trials. *Statistics in Medicine* 2002;**21**:2971-80.

Douglas-Hall 2015

Douglas-Hall P, Whicher EV. 'As required' medication regimens for seriously mentally ill people in hospital. *Cochrane Database of Systematic Reviews* 2015, Issue 12. [DOI: [10.1002/14651858.CD003441.pub3](https://doi.org/10.1002/14651858.CD003441.pub3)]

Du 2017

Du M, Wang X, Yin S, Shu W, Hao R, Zhao S, Rao H, Yeung WL, Jayaram MB, Xia J. De-escalation techniques for psychosis-induced aggression or agitation. *Cochrane Database of Systematic Reviews* 2017, Issue 4. [DOI: [10.1002/14651858.CD009922.pub2](https://doi.org/10.1002/14651858.CD009922.pub2)]

Easterbrook 1991

Easterbrook PJ, Berlin JA, Gopalan R, Matthews DR. Publication bias in clinical research. *Lancet* 1991;**337**(8746):867-72. [PUBMED: 1672966]

Egger 1997

Egger M, Davey Smith G, Schneider M, Minder C. Bias in meta-analysis detected by a simple, graphical test. *BMJ* 1997;**315**:629-34.

Elbourne 2002

Elbourne D, Altman DG, Higgins JPT, Curtina F, Worthington HV, Vaile A. Meta-analyses involving cross-over trials: methodological issues. *International Journal of Epidemiology* 2002;**31**(1):140-9.

Furukawa 2006

Furukawa TA, Barbui C, Cipriani A, Brambilla P, Watanabe N. Imputing missing standard deviations in meta-analyses can provide accurate results. *Journal of Clinical Epidemiology* 2006;**59**(7):7-10.

Gillies 2013

Gillies D, Sampson S, Beck A, Rathbone J. Benzodiazepines for psychosis-induced aggression or agitation. *Cochrane Database of Systematic Reviews* 2013, Issue 4. [DOI: [10.1002/14651858.CD003079.pub3](https://doi.org/10.1002/14651858.CD003079.pub3)]

Glazer 1999

Glazer WM. Does loxapine have "atypical" properties? Clinical evidence. *Journal of Clinical Psychiatry* 1999;**60**(Suppl 10):42-6. [PUBMED: 10340686]

Gulliford 1999

Gulliford MC. Components of variance and intraclass correlations for the design of community-based surveys and intervention studies: data from the Health Survey for England 1994. *American Journal of Epidemiology* 1999;**149**:876-83.

Guy 1976

Guy W. Clinical Global Impression. ECDEU Assessment Manual for Psychopharmacology. revised. National Institute of Mental Health, 1976:217-21.

Higgins 2003

Higgins JP, Thompson SG, Deeks JJ, Altman DG. Measuring inconsistency in meta-analyses. *BMJ* 2003;**327**:557-60.

Higgins 2011

Higgins JPT, Green S (editors). Cochrane Handbook for Systematic Reviews of Interventions Version 5.1.0 [updated March 2011]. The Cochrane Collaboration, 2011. Available from www.cochrane-handbook.org.

Huf 2016

Huf G, Alexander J, Gandhi P, Allen MH. Haloperidol plus promethazine for psychosis-induced aggression. *Cochrane Database of Systematic Reviews* 2016, Issue 11. [DOI: [10.1002/14651858.CD005146.pub3](https://doi.org/10.1002/14651858.CD005146.pub3)]

Isbister 2010

Isbister GK, Calver LA, Page CB, Stokes B, Bryant JL, Downes MA. Randomized controlled trial of intramuscular droperidol versus midazolam for violence and acute behavioral disturbance: the DORM study. *Annals of Emergency Medicine* 2010;**56**(4):392-401. [PUBMED: 20868907]

Janssen-Cilag Ltd 2010

Janssen-Cilag Ltd. Haldol (haloperidol) 5mg/ml Injection. Summary of product characteristics. <http://www.medicines.org.uk/emc/medicine/7267/spc/haldol+injection/> (accessed 3 June 2011).

www.medicines.org.uk/emc/medicine/7267/spc/haldol+injection/ (accessed 3 June 2011).

Jayakody 2012

Jayakody K, Gibson RC, Kumar A, Gunadasa S. Zuclopenthixol acetate for acute schizophrenia and similar serious mental illnesses. *Cochrane Database of Systematic Reviews* 2012, Issue 4. [DOI: [10.1002/14651858.CD000525.pub3](https://doi.org/10.1002/14651858.CD000525.pub3)]

Kay 1986

Kay SR, Opler LA, Fiszbein A. Positive and Negative Syndrome Scale (PANSS) Manual. North Tonawanda, NY: Multi-Health Systems, 1986.

Kay 1987

Kay SR, Fiszbein A, Opfer LA. The Positive and Negative Syndrome Scale (PANSS) for Schizophrenia. *Psychological Reports* 1987;**10**:799-812.

Khokhar 2016

Khokhar MA, Rathbone J. Droperidol for psychosis-induced aggression or agitation. *Cochrane Database of Systematic Reviews* 2016, Issue 12. [DOI: [10.1002/14651858.CD002830.pub3](https://doi.org/10.1002/14651858.CD002830.pub3); CD002830]

Khushu 2016

Khushu A, Powney MJ. Haloperidol for long-term aggression in psychosis. *Cochrane Database of Systematic Reviews* 2016, Issue 11. [DOI: [10.1002/14651858.CD009830.pub2](https://doi.org/10.1002/14651858.CD009830.pub2)]

Leucht 2005

Leucht S, Kane JM, Kissling W, Hamann J, Etschel E, Engel R. Clinical implications of brief psychiatric rating scale scores. *British Journal of Psychiatry* 2005;**187**:366-71. [PUBMED: 16199797]

Leucht 2005a

Leucht S, Kane JM, Kissling W, Hamann J, Etschel E, Engel RR. What does the PANSS mean?. *Schizophrenia Research* 2005;**79**(2-3):231-8. [PUBMED: 15982856]

Leucht 2007

Leucht S, Engel RR, Bauml J, Davis JM. Is the superior efficacy of new generation antipsychotics an artifact of LOCF?. *Schizophrenia Bulletin* 2007;**33**(1):183-91. [PUBMED: 16905632]

Lexchin 2003

Lexchin J, Bero LA, Djulbegovic B, Clark O. Pharmaceutical industry sponsorship and research outcome and quality: Systematic review. *BMJ* 2003;**326**(7400):1167-70. [PUBMED: 12775614]

Lieberman 2005

Lieberman JA, Stroup TS, McEvoy JP, Swartz MS, Rosenheck RA, Perkins DO, et al. Clinical Antipsychotic Trials of Intervention Effectiveness (CATIE) Investigators. Effectiveness of antipsychotic drugs in patients with chronic schizophrenia. *New England Journal of Medicine* 2005;**353**(12):1209-23. [PUBMED: 16172203]

López-Munoz 2009

López-Munoz F, Alamo C. The consolidation of neuroleptic therapy: Janssen, the discovery of haloperidol and its introduction into clinical practice. *Brain Research Bulletin* 2009;**79**:130–41.

Marshall 2000

Marshall M, Lockwood A, Bradley C, Adams C, Joy C, Fenton M. Unpublished rating scales: a major source of bias in randomised controlled trials of treatments for schizophrenia. *British Journal of Psychiatry* 2000;**176**:249–52.

Moher 2001

Moher D, Schulz KF, Altman DG. The CONSORT statement: revised recommendations for improving the quality of reports of parallel-group randomised trials. *Lancet* 2001;**357**(9263):1191–4.

Mohr 2005

Mohr P, Pecenek J, Svestka J, Swingle D, Treuer T. Treatment of acute agitation in psychotic disorders. *Neuro Endocrinology Letters* 2005;**26**(4):327–35. [PUBMED: 16136016]

Montoya 2011

Montoya A, Valladares A, Lizán L, San L, Escobar R, Paz S. Validation of the Excited Component of the Positive and Negative Syndrome Scale (PANSS-EC) in a naturalistic sample of 278 patients with acute psychosis and agitation in a psychiatric emergency room. *Health Quality Life Outcomes* 2011;**9**:18.

Muralidharan 2006

Muralidharan S, Fenton M. Containment strategies for people with serious mental illness. *Cochrane Database of Systematic Reviews* 2006, Issue 3. [DOI: [10.1002/14651858.CD002084.pub2](https://doi.org/10.1002/14651858.CD002084.pub2)]

NHS 2009

National Health Service. The prescribing preview report on antipsychotic drugs, available to general practitioners in August 2009, is reproduced here for readers with an interest in patterns and trends of prescribing. Impact: Prescribing and Dispensing Newsletter. http://www.nhs.uk/PrescriptionServices/Documents/PrescriptionServices/Impact_Dec_09_8p.pdf (accessed 2nd June 2011):4–6.

NICE 2005

National Institute for Clinical Excellence. The short term-management of disturbed/violent behaviour in inpatient psychiatric settings and emergency departments. Vol. **25**, London: NICE, 2005.

NICE 2015

National Institute for Clinical Excellence. Violence and aggression: short-term management in mental health, health and community settings. nice.org.uk/guidance/ng10 2015.

Olkola 2008

Olkola KT, Ahonen J. Midazolam and other benzodiazepines. *Handbook of Experimental Pharmacology* 2008;**182**:335–60. [PUBMED: 18175099]

Overall 1962

Overall JE, Gorham DR. The brief psychiatric rating scale. *Psychological Reports* 1962;**10**:799–812.

Parker 2010

Parker C, Khwaja MG. What is new in rapid tranquillisation?. *Journal of Psychiatric Intensive Care* 2010;**7**(2):1–11.

Pereira 2007

Pereira S, Fleischacker W, Allen M. Management of behavioural emergencies. *Journal of Psychiatric Intensive Care* 2007;**2**(2):71–83.

Pratt 2008

Pratt P, Chandler-Oatts J, Nelstrop L, Branford D, Periera S, Johnston S. Establishing gold standard approaches to rapid tranquillisation: A review and discussion of the evidence of the safety and efficacy of medications used. *Journal of Psychiatric Intensive Care* 2008;**4**(1–2):43–57.

Reid 2003

Reid G, Hughson M. Droperidol dropped; consultants not consulted: A survey of the practice of rapid tranquillisation by consultant psychiatrists in the west of Scotland. *Psychiatric Bulletin* 2003;**27**:301–4.

Sailas 2000

Sailas E, Fenton M. Seclusion and restraint for people with serious mental illnesses. *Cochrane Database of Systematic Reviews* 2000, Issue 2. [DOI: [10.1002/14651858.CD001163](https://doi.org/10.1002/14651858.CD001163)]

Schleifer 2011

Schleifer JJ. Management of acute agitation in psychosis: an evidence-based approach in the USA. *Advances in Psychiatric Treatment* 2011;**17**:91–100.

Schünemann 2008

Schünemann HJ, Oxman AD, Vist GE, Higgins JPT, Deeks JJ, Glasziou P, et al. Chapter 12: Interpreting results and drawing conclusions. In: Higgins JPT, Green S editor(s). *Cochrane Handbook for Systematic Reviews of Interventions*. The Cochrane Collaboration, 2008:359–83.

Simpson 1970

Simpson GM, Angus JSW. A rating scale for extrapyramidal side effects. *Acta Psychiatrica Scandinavica. Supplementum* 1970;**212**:11–9.

Thorpe 2009

Thorpe KE, Zwarenstein M, Oxman AD, Treweek S, Furberg CD, Altman DG, et al. A pragmatic-explanatory continuum indicator summary (PRECIS): a tool to help trial designers. *Journal of Clinical Epidemiology* 2009;**62**(5):464–75. [PUBMED: 19348971]

TREC 2003

TREC Collaborative Group. Rapid tranquillisation for agitated patients in emergency psychiatric rooms: a randomised trial of midazolam versus haloperidol plus promethazine. *BMJ* 2003;**327**(7417):708–13. [PUBMED: 14512476]

Ukoumunne 1999

Ukoumunne OC, Gulliford MC, Chinn S, Sterne JAC, Burney PGJ. Methods for evaluating area-wide and organisation-based intervention in health and health care: A systematic review. *Health Technology Assessment* 1999;**3**(5):1-75.

Vangala 2012

Vangala R, Ahmed U, Ahmed R. Loxapine inhaler for psychosis-induced aggression. *Cochrane Database of Systematic Reviews* 2012.

WHO 2011

World Health Organization. WHO Essential Medicines List. http://whqlibdoc.who.int/hq/2011/a95053_eng.pdf (accessed 2nd June 2011).

WHO 2015

World Health Organization. WHO Model List of Essential Medicines. http://www.who.int/medicines/publications/essentialmedicines/EML_2015_FINAL_amended_NOV2015.pdf?ua=1 2015.

Wilkie 2012

Wilkie F, Fenton M. Quetiapine for psychosis-induced aggression or agitation. *Cochrane Database of Systematic Reviews* 2012, Issue 4. [DOI: [10.1002/14651858.CD009801](https://doi.org/10.1002/14651858.CD009801)]

Xia 2009

Xia J, Adams CE, Bhagat N, Bhagat V, Bhoopathi P, El-Sayeh H, et al. Loss to outcomes stakeholder survey: the LOSS study. *Psychiatric Bulletin* 2009;**33**(7):254-7.

Xiberas 2001

Xiberas X, Martinot JL, Mallet L, Artiges E, Loc'h C, Maziere B, et al. Extrastriatal and striatal D2 dopamine receptor blockade with haloperidol or new antipsychotic drugs in patients with schizophrenia. *British Journal of Psychiatry* 2001;**179**:503-8.

Yudofsky 1986

Yudofsky SC, Silver JM, Jackson W, Endicott J, Williams D. The Overt Aggression Scale for the objective rating of verbal and physical aggression. *American Journal of Psychiatry* 1986;**143**:35-9.

Yudofsky 1997

Yudofsky SC, Kopecky HJ, Kunik M, Silver JM, Endicott J. The Overt Agitation Severity Scale for the objective rating of agitation. *Journal of Neuropsychiatry* 1997;**9**(4):541-8.

References to other published versions of this review

Powney 2011

Powney MJ, Adams CE, Jones H. Haloperidol (rapid tranquillisation) for psychosis induced aggression or agitation. *Cochrane Database of Systematic Reviews* 2011, Issue 10. [DOI: [10.1002/14651858.CD009377](https://doi.org/10.1002/14651858.CD009377)]

Powney 2012

Powney MJ, Adams CE, Jones H. Haloperidol for psychosis-induced aggression or agitation (rapid tranquillisation). *Cochrane Database of Systematic Reviews* 2012, Issue 11. [DOI: [10.1002/14651858.CD009377.pub2](https://doi.org/10.1002/14651858.CD009377.pub2)]

* Indicates the major publication for the study

CHARACTERISTICS OF STUDIES

Characteristics of included studies [ordered by study ID]

Bailine 1987

Methods	Allocation: randomised. Blindness: blind for some outcomes. Duration: 72 hours.
Participants	Diagnosis: Research Diagnostic Criteria (11) diagnosis of schizophrenia, any type, or manic depressive illness, manic type and grossly abnormal behavior. N = 34. Age: mean 25.2 years. Sex: 26 males, 8 females. History: "violent or assaultive or too psychologically disorganized to be maintained on a unit without significant extra supervision." (p.128). Excluded: not reported. Setting: inpatient, USA.

Bailine 1987 (Continued)

Interventions	<p>1. Haloperidol IM: dose 5 mg or 10 mg/IM, maximum cumulative dose of 100 mg during 24 hours (mean number of IMs not reported). N = 20.</p> <p>2. Chlorpromazine IM: dose 50 mg or 100 mg/IM, maximum cumulative dose of 500 mg during 24 hours (mean number of IMs not reported). N = 14.</p>
Outcomes	<p>Adverse events: leaving the study early.</p> <p>Unable to use:</p> <p>Mental state: BPRS (mean and overall P value only for significant findings reported, SD/SE/CI not reported).</p> <p>Global state: GAS (not published, mean and overall P value only for significant findings reported, SD/SE/CI not reported).</p> <p>Global state: NOSIE (overall P value only for significantly findings reported, mean/SD/SE/CI not reported).</p> <p>Adverse events: SAS (overall P value only for significantly findings reported, mean/SD/SE/CI not reported).</p>
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"random number sequence".
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Not stated.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	BPRS was rater-blinded. However, it is not clear whether other outcomes were blinded.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	It is reported that no one discontinued due to adverse effects, however it does not state if there were discontinuations due to any other reason.
Selective reporting (reporting bias)	High risk	Only the significant findings/factors are reported for a number of outcomes.
Other bias	High risk	Small short study.

Baldacara 2011

Methods	<p>Allocation: randomised.</p> <p>Blindness: single (rater-blinded)*.</p>
---------	---

Baldacara 2011 (Continued)

Duration: 12 hours.

Participants	<p>Diagnosis: DSM-IV-TR criteria for bipolar (manic or mixed episode) or psychotic disorder.</p> <p>N = 150.</p> <p>Age: 18-50 years.</p> <p>Sex: 91 males, 59 females.</p> <p>History: "admitted to the Psychiatric Emergency Room" (p.31).</p> <p>Excluded: "disorders due to drug abuse, organic disorder, anxiety or personality disorder (DSM-IV-TR criteria), failure to agree to participate in the study, incapability of completing all steps and unstable clinical disease" (p.31).</p> <p>Setting: emergency room, São Paulo.</p>
Interventions	<p>1. Haloperidol IM: fixed dose 5 mg/IM. N = 30.</p> <p>2. Olanzapine IM: fixed dose 10 mg/IM. N = 30.</p> <p>3. Ziprasidone IM: fixed dose 20 mg/IM. N = 30.</p> <p>4. Haloperidol IM: fixed dose 5 mg/IM + promethazine IM: fixed dose 50 mg/IM. N = 30.</p> <p>5. Haloperidol IM: fixed dose 5 mg/IM + midazolam IM: fixed dose 15 mg/IM. N = 30.</p> <p>"Only additional doses of haloperidol + promethazine could be used" (p.31).</p>
Outcomes	<p>Need for additional drugs for tranquillisation.</p> <p>Agitation: OASS**.</p> <p>Aggression: OAS**.</p> <p>Global state: restriction.</p> <p>Adverse events: death, RSS, EPS, excessive sedation, hypotension.</p>
Notes	<p>**"patients were instructed not to reveal their current treatment to the investigators" (p.31).</p> <p>**endpoint scores at 1 hour are discordant between text (p.32-33) and Table 2 (p.35); scores from Table 2 were used, due to compatibility with Figure 2 (p.36).</p> <p>Authors were contacted (3rd July 2016, 7th July 2016, 9th July 2016).</p>

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	"randomization employed in this clinical trial was allocation by permuted blocks" (p.31); no further informations on the randomisation process.
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	High risk	"study medications were packaged in identical color-coded boxes" (p.31); stated as double blind, however "patients were instructed not to reveal their current treatment to the investigators" (p.31); no information whether the blindness was effective.

Baldacara 2011 (Continued)

Blinding of outcome assessment (detection bias) All outcomes	High risk	"Patients were assessed by two psychiatrists. The psychiatrists were all masked with regard to the patient's treatment assignment" (p.31); no information whether the blindness was effective.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete outcome data.
Selective reporting (reporting bias)	High risk	Endpoint scores at 1 hour are discordant between text (p.32-33) and Table 2 (p.35).
Other bias	Low risk	None obvious.

Battaglia 1997

Methods	Allocation: randomised. Blindness: double. Duration: 24 hours.
Participants	Diagnosis: mania (N = 13), psychoactive substance use (N = 16), psychosis not otherwise specified (N = 27), schizophrenia (N = 47), schizophreniform disorder (N = 1).* N = 100.** Age: range 18-57 years. Sex: 73 males, 25 females. History: "psychosis and behavioral dyscontrol (agitated, aggressive, destructive, assaultive, or restless behavior." Excluded: "clinically obvious alcohol intoxication, central nervous system (CNS) depression, pregnancy, allergic hypersensitivity, delirium, neuroleptic malignant syndrome, airway obstruction, severe hypotension or hypertension, acute narrow angle glaucoma and treatment with a benzodiazepine or neuroleptic within the previous 24 hours (previous two weeks for fluphenazine decanoate and previous 4 weeks for haloperidol decanoate)." Setting: Emergency department, USA hospitals.
Interventions	1. Haloperidol IM: dose up to 6 injections of 5 mg (mean number of IMs 2.86). N = 35. 2. Lorazepam IM: dose up to 6 injections of 2 mg (mean number of IMs 2.87). N = 31. 3. Lorazepam IM: dose up to 6 injections of 2 mg + haloperidol 5 mg (mean number of IMs 2.41). N = 32. First 3 injections - at least 1 hour apart, remainder 2 hours apart. Need for subsequent doses made by blinded evaluator.
Outcomes	Tranquillisation or asleep. Global state: number of additional injections.** Adverse events: EPS, hypertension, others in over $\geq 9\%$ of people. Unable to use - Global state: CGI (reported, out of necessity, only for those awake), Efficacy Index (no data reported).

Battaglia 1997 (Continued)

Agitation: ABS (mean, SE/SD not reported, P values of significant findings reported, overall F value).

Mental state: MBPRS (modified, unpublished; mean, SE/SD not reported, P values of significant findings reported, overall F value), Alertness Scale (no mean, SE/SD not reported, P values of significant findings reported).

Physiological: vital signs (not reported).

Notes

* 6 people had more than one diagnosis.

** 2 people excluded from efficacy analysis - after enrolment they had received proscribed antipsychotic medication.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"Computer-generated table of random numbers".
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	"The ED [Emergency Department] psychiatrist who <i>treated</i> and rated the patient remained blinded to the identity of the patient's medication throughout treatment" (p.336). [italic added by reviewers] "Double blind" - participant blinding not specifically mentioned.
Blinding of outcome assessment (detection bias) All outcomes	Low risk	"The ED [Emergency Department] psychiatrist who treated and <i>rated</i> the patient remained blinded to the identity of the patient's medication throughout treatment" (p.336). [italic added by reviewers]. Not tested.
Incomplete outcome data (attrition bias) All outcomes	High risk	"two patients were excluded from the efficacy analysis. It was determined after enrolment that both received proscribed antipsychotic medication before entering the study."
Selective reporting (reporting bias)	High risk	More details on level of significance for outcomes with $P < 0.05$. Adverse effects for central nervous system reported only if present in $\geq 9\%$.
Other bias	High risk	Sponsored by drug company.

Battaglia 2002

Methods

Allocation: randomised.

Blindness: double.

Duration: 24 hours, preceded by a 2 hour screening period.

Participants

Diagnosis: DSM-IV schizophrenia, schizophreniform disorder, or schizoaffective disorder.

N = 311

Age: mean 38.2 years.

Sex: 204 males, 107 females.

Battaglia 2002 (Continued)

History: mean age of illness onset, years (SD); haloperidol 25.1 (8.3), olanzapine 23.5 (8.9), placebo 24.9 (8.0).

Excluded: Pregnant or lactating women, people with serious medical illnesses or people currently pre-scribed antipsychotic medication via depot injection.

Setting: Multi-centre (Australia, Austria, Belgium, Canada, the Czech Republic, France, Greece, Hungary, Israel, the Republic of South Africa, Spain, the UK, and the USA).

Interventions	<p>1. Haloperidol IM: dose up to 3 injections of 7.5 mg (mean number of IMs 1.27). N = 126.</p> <p>2. Olanzapine IM: dose up to 3 injections of 10 mg (mean number of IMs 1.3). N = 131.</p> <p>3. Placebo IM: up to 3 injections (mean number of IMs ~ 1.5). N = 54.</p> <p>Need for subsequent doses at the discretion of the investigator.</p>
Outcomes	<p>Mental state: BPRS.</p> <p>Global state: CGI, number of additional injections, use of benzodiazepine.</p> <p>Adverse effects: SAS, BAS, COSTART, QT changes, use of anticholinergic drugs.</p> <p>Agitation: ACES, PANSS-EC.</p>
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	"Both drugs and placebo were administered in identical, color-blinded, translucent standard syringes, and raters and study <i>personnel</i> were blind to treatment assignment" (Wright 2001, p.1149). [italic added by reviewers]
Blinding of outcome assessment (detection bias) All outcomes	Low risk	"Both drugs and placebo were administered in identical, color-blinded, translucent standard syringes, and <i>raters</i> and study personnel were blind to treatment assignment" (Wright 2001, p.1149). [italic added by reviewers]
Incomplete outcome data (attrition bias) All outcomes	High risk	The authors only account for 5 of the 26 people who did not complete the 24 hours study.
Selective reporting (reporting bias)	Low risk	No indication of selective reporting.
Other bias	High risk	Sponsored by the manufacturers of olanzapine.

Breier 2002

Methods	Allocation: randomised.
---------	-------------------------

Haloperidol for psychosis-induced aggression or agitation (rapid tranquillisation) (Review)

Copyright © 2017 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.

Breier 2002 (Continued)

	<p>Blindness: double.</p> <p>Duration: 24 hours.</p>
Participants	<p>Diagnosis: schizophrenia, schizophreniform or schizoaffective disorder.</p> <p>N = 270.</p> <p>Age: range 18-73 years.</p> <p>Sex: 155 males, 115 females.</p> <p>History: "recently hospitalised with acute agitation." Mean age at onset of illness 25.1 (SD 7.3) years.</p> <p>Excluded: people with "significant medical disorders, including alcohol and/or drug dependency, were excluded from this trial."</p> <p>Setting: Multi-centre (Croatia, Italy, Romania and South Africa).</p>
Interventions	<p>1. Haloperidol IM: dose up to 3 injections of 7.5 mg/IM (mean number of IMs not reported, mean dose 9.9 mg (SD 4.6). N = 40.</p> <p>2. Olanzapine IM: dose up to 3 injections of 2.5 mg/IM (mean number of IMs not reported, mean dose 4 mg (SD 1.5). N = 48.</p> <p>3. Olanzapine IM: dose up to 3 injections of 5 mg/IM (mean number of IMs not reported, mean dose 6.9 mg (SD 2.7). N = 45.</p> <p>4. Olanzapine IM: dose up to 3 injections of 7.5 mg/IM (mean number of IMs not reported, mean dose 9.8 mg (SD 3.8). N = 46.</p> <p>5. Olanzapine IM: dose up to 3 injections of 10 mg/IM (mean number of IMs not reported, mean dose 12.6 mg (SD 4.9). N = 46*.</p> <p>6. Placebo IM. N = 50.</p> <p>2nd injection allowed 2 hours after 1st, 3rd injection allowed 4 hours after 2nd injection. Both to be administered within 20 hours of 1st injection.</p> <p>Need for subsequent doses at the discretion of the investigator.</p>
Outcomes	<p>Mental state: BPRS Total, BPRS Positive.</p> <p>Global state: CGI-S, additional injections, need for benzodiazepine.</p> <p>Agitation: ABS, PANSS-EC.</p> <p>Adverse events: QT interval at 24 hours, hypotension**, acute dystonia**, EPS**.</p> <p>Unable to use*** -</p> <p>Leaving the study early: reasons are not given for why participants did not complete the study.</p> <p>Adverse effects: SAS (mean, SE/SD not reported).</p> <p>Adverse effects: BAS (mean, SE/SD not reported).</p> <p>Adverse effects: need for anticholinergic therapy (olanzapine 10 mg/IM not reported).</p> <p>Adverse effects: QT interval at 2 hours (mean, SE/SD not reported).</p>
Notes	<p>* For data analysis, where continuous outcomes were used, haloperidol was compared with Olanzapine 10 mg/IM as this is the dose referred to as usual by the BNF (BNF 2011).</p> <p>** It is possible that this binary data was derived from the BAS scale.</p>

Breier 2002 (Continued)

*** Additional data are reported by dose but not used in this review (Table 4; Table 5; Table 6; Table 7; Table 8)

Risk of bias		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Low risk	"Allocation concealed placebo-controlled trial." Method of randomisation not reported.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	"unblinded third-party personnel, who played no role in evaluating patients, were trained to handle and administer injections in identical, unmarked syringes." "Investigators, patients, clinical carers and raters were blinded."
Blinding of outcome assessment (detection bias) All outcomes	Low risk	"Investigators, patients, clinical carers and raters were blinded."
Incomplete outcome data (attrition bias) All outcomes	High risk	Reasons why all participants that left the study early were not reported.
Selective reporting (reporting bias)	Unclear risk	It is not clear whether all side effects are reported.
Other bias	High risk	Trial sponsored by manufacturers of olanzapine IM.

Bristol Myers 2004a

Methods	Allocation: randomised, 2:2:1 ratio. Blindness: double. Duration: 2-hour screening period, 24 hours initial phase, 4-day second phase, 35 days follow-up.
Participants	Diagnosis: DSM-IV diagnosis of schizophrenia (N = 324) or schizoaffective disorder (N = 124). N = 448. Age: range 18-69 years. Sex: males and females. History: voluntarily hospitalised, experiencing acute agitation, mean age of diagnosis ~ 26.3 years. Excluded: a diagnosis of schizophreniform disorder; a psychiatric disorder other than schizophrenia requiring pharmacotherapy, those at significant risk of suicide, neurological diagnoses (including migraine, Parkinson's disease, epilepsy, Alzheimer's disease, residual of stroke, multiple sclerosis, transient cerebral ischaemic attacks, learning disability, cerebral palsy); a history of seizures, severe head trauma, abnormal ECG, stroke or evidence of other unstable medical conditions or adverse event, substance or alcohol dependence within 2 months of the study, history of neuroleptic malignant syndrome from antipsychotic agents, suspected substance-induced psychiatric disorder or behavioural disturbance, benzodiazepine or anticholinergics within 4 hours before the first injection of study medication; lack of response to previous antipsychotic medication.

Bristol Myers 2004a (Continued)

Setting: Multi-centre (USA, Czech Republic, France, Estonia, Latvia, Poland, Croatia, Italy, Puerto Rico, South Africa, and Spain).

Interventions	<p>1. Haloperidol: dose 6.5 mg/IM, maximum 3 doses during first 24 hours (mean number of IMs 1.43). N = 185. Followed by haloperidol: max dose 7 mg/oral to 10 mg/oral in 24 hours for 4 days. N = 151.</p> <p>2. Aripiprazole: dose 10 mg/IM, maximum 3 doses during first 24 hours (mean number of IMs 1.54). N = 175. Followed by aripiprazole: max dose 10 mg/oral to 15 mg/oral in 24 hours for 4 days. N = 153.</p> <p>3. Placebo IM, maximum 2 doses during first 24 hours. If 3rd dose needed aripiprazole: dose 10 mg/IM administered (mean number of IMs 1.92). N = 88. Followed by aripiprazole: max dose 10 mg/oral to 15 mg/oral in 24 hours for 4 days. N = 76.</p>
Outcomes	<p>Global state: CGI-S, CGI-I, additional medication, need for benzodiazepine.</p> <p>Leaving the study early.</p> <p>Adverse events: concomitant medication, death, ECG, physical examination, vital signs.</p> <p>Leaving the study early.</p> <p>Unable to use:</p> <p>Adverse effects: SAS (SE/SD/CI not reported).</p> <p>Adverse effects: BAS (SE/SD/CI not reported).</p>

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Low risk	"Randomization was conducted via a centralized call-in system and was organized by study center using permuted block randomization".
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	During the IM phase, haloperidol, aripiprazole and placebo were delivered in 1.3 mL injections. However, in the oral phase a tablet was used for aripiprazole and a capsule was used for haloperidol and so it is possible that this presented a risk to blinding.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Described as "double blind", however no further details are given.
Incomplete outcome data (attrition bias) All outcomes	High risk	Reasons why all participants that left the study early were not reported.
Selective reporting (reporting bias)	High risk	The author mentions that there is increased glucose levels in both aripiprazole and haloperidol group but does not provide data. The authors only report the side effects that occurred in $\geq 2\%$ of people.
Other bias	High risk	Sponsored by the manufacturers of aripiprazole.

Bristol-Myers 2005b

Methods	<p>Allocation: randomised.</p> <p>Blindness: double.</p> <p>Duration: 24 hours (preceded by 2-hour screening period).</p>
Participants	<p>Diagnosis: DSM-IV diagnosis of schizophrenia (N = 237), schizoaffective disorder (N = 113) or schizophreniform disorder (N = 7).</p> <p>N = 357*.</p> <p>Age: 18-66 years.</p> <p>Sex: 214 males, 143 females.</p> <p>History: acute agitation.</p> <p>Excluded: psychoactive substance dependence 2 months prior to the start of the study, required involuntary restraint, were suicidal, had a neurological or psychiatric condition other than schizophrenia, schizoaffective disorder or schizophreniform disorder, had a significant medical condition or were known non responders to antipsychotic medication.</p> <p>Setting: Multi-centre (USA, Canada, Czech Republic, France, Estonia, Latvia, Lithuania, Spain).</p>
Interventions	<ol style="list-style-type: none"> 1. Haloperidol IM: dose up to 3 injections of 7.5 mg/IM (mean number of IMs 1.33). N = 60. 2. Aripiprazole IM: dose up to 3 injections of 1 mg/IM (mean number of IMs not reported). N = 57. 3. Aripiprazole IM: dose up to 3 injections of 5 mg/IM (mean number of IMs not reported). N = 63. 4. Aripiprazole IM: dose up to 3 injections of 10 mg/IM (mean number of IMs not reported). N = 57*. 5. Aripiprazole IM: dose up to 3 injections of 15 mg/IM (mean number of IMs not reported). N = 58. 6. Placebo IM (mean number of IMs not reported). N = 50.
Outcomes	<p>Mental state: BPRS Total and Positive Scores.</p> <p>Global state: CGI-I, CGI-S, number of re-injections*.</p> <p>Agitation: ACES, CABS, PANSS-EC.</p> <p>Leaving the study early.</p> <p>Adverse events: adverse event reports. Table 9</p> <p>Unable to use***:</p> <p>Adverse effects: SAS (SE/SD/CI not reported).</p> <p>Adverse effects: BAS (SE/SD/CI not reported).</p> <p>Physiological: QTc is not reported for 9.75 mg. Results of clinical laboratory tests, physical examination, vital sign measurement are not reported.</p>
Notes	<p>* For data analysis, where continuous outcomes were used, haloperidol was compared with aripiprazole 10 mg/IM as this is the dose referred to as usual by the BNF (BNF 2011).</p> <p>**The number of additional injections needed was not reported individually for each of the aripiprazole doses but given as a range (i.e. 56% to 60% received 1 injection), therefore the author chose the mean value (i.e. 58%).</p> <p>*** Additional data are reported by dose but not used in this review (Table 9; Table 10)</p>

Bristol-Myers 2005b (Continued)

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	High risk	No blinding was used during the preparation of injections and in most cases the person preparing the drug administered the injection.
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Study investigators conducting the assessments were blinded to treatment.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	The reasons for leaving the study early were reported for those participants who did not complete the study. However, the number of participants completing different assessments at the same time point vary and reasons are not given.
Selective reporting (reporting bias)	High risk	The authors only report the adverse effects that occurred in $\geq 5\%$ of people. The QTc abnormality for 9.75 mg is not reported.
Other bias	High risk	Sponsored by the manufacturers of aripiprazole.

Brook 2000

Methods	Allocation: randomised. Blindness: open. Duration: 7 days.
Participants	Diagnosis: diagnosis of schizophrenia, schizoaffective disorder, bipolar disorder, brief psychotic disorder, schizophreniform disorder, delusional disorder, or psychotic disorders not otherwise specified as defined by DSM-III-R. N = 132. Age: range 19-66 years. Sex: 123 males, 9 females. History: recently hospitalised with acute psychosis. Excluded: substance-induced psychosis (confirmed by urinalysis), psychosis related to organic origin, clinically relevant medical illness, abnormal ECG, imminent risk of suicide or homicide, history of substance abuse or dependence in the previous 2 months. Setting: Multi-centre (7 countries).
Interventions	1. Haloperidol: flexible dose 2.5 mg/IM to 10 mg/IM (mean number of IMs not reported), maximum 4 doses in 24 hours for 3 days. Followed by haloperidol: flexible dose 10 mg/oral to 80 mg/oral during 24 hours for 4 days. N = 42.

Brook 2000 (Continued)

2. Ziprasidone: dose 10 mg/IM, maximum 4 doses in 24 hours for 3 days (mean number of IMs not reported). Followed by ziprasidone: flexible dose 80 mg/oral to 200 mg/oral in 24 hours for 4 days. N = 90.

Outcomes	<p>Mental state: BPRS.</p> <p>Global state: CGI-S.</p> <p>Leaving the study early.</p> <p>Adverse events: COSTART, BAS, modified SAS, investigators assessment of severity, blood pressure, laboratory tests, concomitant medication, ECG, urinalysis, others in ≥ 10 of participants.</p> <p>Unable to use</p> <p>Specific behaviours: BPRS agitation items (not published).</p> <p>Global state: CGI-I (SE/SD not reported).</p> <p>Sedation score: 5-point categorical scale (not published, present mean/SD rather than binary data).</p>
----------	---

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"Computerized randomization assigned".
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	High risk	Open trial.
Blinding of outcome assessment (detection bias) All outcomes	High risk	Open trial.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete outcome data. A table is provided which presents a summary of discontinuations.
Selective reporting (reporting bias)	High risk	The number of participants reported as discontinuing due to adverse effects is reported inconsistently (see Brook 1998c table 2 and Brook 2000 table 4). It is not clear whether all adverse events are reported. Brook 2000 table 4 only reports adverse events occurring in ≥ 10 of people, however there are more adverse events mentioned throughout the text.
Other bias	High risk	Sponsored by drug company.

Calver 2015

Methods	<p>Allocation: randomised.</p> <p>Blindness: double.</p>
---------	--

Calver 2015 (Continued)

Duration: 2 hours.

Participants	<p>Diagnosis: mental illness (N = 114), drug-induced psychosis (N = 70), intoxication (N = 17), threatened self-harm (N = 6), other/unknown (N = 19) (p.225)*.</p> <p>N = 228.</p> <p>Age: major than 18 years.</p> <p>Sex: 144 males, 84 females.</p> <p>History: admitted involuntarily to the psychiatric intensive care unit from the psychiatric emergency care centre (p.223).</p> <p>Excluded: patients willing to take oral medication for sedation without physical restraint or seclusion.</p> <p>Setting: psychiatry intensive care unit.</p>
Interventions	<p>1. Haloperidol: fixed dose 10 mg/IM. N = 110.</p> <p>2. Droperidol: fixed dose 10 mg/IM. N = 118.</p>
Outcomes	<p>Tranquillisation or asleep, time to sedation.</p> <p>Need for additional benzodiazepines.</p> <p>Adverse events: respiratory rate less than 12 breaths/min, systolic blood pressure less than 90 mmHg, heart rate less than 60 bpm, oxygen saturation less than 90% or the presence of extrapyramidal side effects.</p>
Notes	*when available, outcomes are reported for the mental illness sub-group; gently provided by the authors (contacted 5th July 2016).

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"Block randomisation was used. Microsoft Excel was used to randomly create blocks of four (ABAB, AABBB, etc.) or six (ABABAB, AAABBB, etc.)" (p.224).
Allocation concealment (selection bias)	Low risk	"Pre-packed treatment kits [...] contained either droperidol (10mg in 2ml) or haloperidol (10mg in 2ml) [...] transferred into vials identical" (p.224).
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Double-blind study.
Blinding of outcome assessment (detection bias) All outcomes	Low risk	"labels A or B and not the drug names were known to the investigator. This investigator analysed the data independently and presented this to the other investigators" (p.224).
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete outcome data.
Selective reporting (reporting bias)	Unclear risk	"In breach of the study protocol midazolam was given nine times simultaneously with the study drug" (p.226); no further evidence of selective reporting.
Other bias	High risk	Short study.

Currier 2004

Methods	Allocation: randomised. Blindness: single-blind (rater-blinded). Duration: 24 hours.
Participants	Diagnosis: DSM-IV diagnosis of paranoid schizophrenia (N = 54), schizoaffective disorder (N = 36), bipolar disorder (N = 13), psychotic disorder not otherwise specified (N = 31), other (N = 28). N = 162. Age: range 18-65 years. Sex: 105 males, 57 females. History: acute agitation. Excluded: pregnancy, delirium, epilepsy, learning disability, intoxication, symptoms of withdrawal from alcohol or other psychoactive substances, medical illness, treatment with any anti-psychotic or benzodiazepine within 6 hours of screening, history of neuroleptic malignant syndrome, known hypersensitivity to any of the trial medications, treatment with depot injection, use of disallowed medication. Setting: multi-centre, USA (24 sites)
Interventions	1. Haloperidol: dose 5 mg/IM + lorazepam 2 mg/oral (mean number of IMs not reported). N = 79. 2. Risperidone: dose 2 mg/oral + lorazepam 2 mg/oral (mean number of doses not reported). N = 83.
Outcomes	Tranquillisation or asleep. Leaving the study early. Global state: CGI-S, additional lorazepam. Agitation: PANSS-EC. Aggression: OAS. Adverse effects: BAS, SAS, EPS, heart monitoring, sedation.

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Low risk	"A telephone-based central service was used to randomly assign eligible patients to receive a single dose of oral treatment or IM injection." (p.387)
Blinding of participants and personnel (performance bias) All outcomes	High risk	Single-blind study.
Blinding of outcome assessment (detection bias)	Low risk	"independent raters blinded to the treatment arm conducted efficacy assessments". (p.387)

Haloperidol for psychosis-induced aggression or agitation (rapid tranquillisation) (Review)

Currier 2004 (Continued)

All outcomes

Incomplete outcome data (attrition bias) All outcomes	Unclear risk	A flow diagram is provided with the reasons for leaving the study early reported. However, the number of participants completing different assessments at the same time point vary and reasons are not given.
Selective reporting (reporting bias)	High risk	The authors only report the side effects that occurred in $\geq 5\%$ of people.
Other bias	High risk	Sponsored by drug company (Janssen Pharmaceutica).

Dorevitch 1999

Methods	Allocation: randomised. Blindness: double. Duration: 2 hours.
Participants	Diagnosis: DSM-IV diagnosis of paranoid schizophrenia (N = 19), schizoaffective disorder (N = 7), bipolar disorder (N = 2). N = 28. Age: range 20-60 years. Sex: 13 males, 15 females. History: "actively psychotic inpatients". (p.142). Excluded: Not reported. Setting: Psychiatric hospital, Tel Aviv.
Interventions	1. Haloperidol: dose 5 mg/IM (mean number of IMs not reported). N = 13. 2. Flunitrazepam: dose 1 mg/IM (mean number of IMs not reported). N = 15.
Outcomes	Global state: need for seclusion or restraint. Aggression: OAS. Adverse effects.

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"assigned by a table of random numbers." (p.142).
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias)	Unclear risk	Described as double-blind, however further details regarding the blinding of participants and personnel are not reported.

Haloperidol for psychosis-induced aggression or agitation (rapid tranquillisation) (Review)

Dorevitch 1999 (Continued)

All outcomes

Blinding of outcome assessment (detection bias) All outcomes	Low risk	"All ratings were completed by the same rater (N.K.), who was blind to the study medications." (p.143).
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete outcome data.
Selective reporting (reporting bias)	Unclear risk	It is not clear whether all side effects are reported.
Other bias	Unclear risk	Small short study.

Eli Lilly 2004

Methods	Allocation: randomised. Blindness: double. Duration: 24-72 hours.
Participants	Diagnosis: DSM-IV diagnosis of schizophrenia. N = 49. Age: range 18-65* years. Sex: 35 males, 14 females. History: "The mean length of current psychotic episode was 13.8 and 17.9 days in the olanzapine and haloperidol groups respectively." (p.6). Excluded: dementia, documented allergy to study drugs, treatment with injectable depot within 1 injection interval, unstable illnesses where death is anticipated within 1 year, or treatment in intensive care expected within 6 months, previous participation in a Lilly sponsored intra-muscular olanzapine clinical trial. Setting: multi-centre, Taiwan.
Interventions	1. Haloperidol: dose 7.5 mg/IM (mean number of IMs 1.13). N = 24. 2. Olanzapine: dose 10 mg/IM (mean number of IMs 1.32). N = 25.
Outcomes	Mental state: BPRS total score, BPRS positive sub-scale. Global state: CGI-I, additional medication. Agitation: ACES, PANSS-EC. Leaving the study early. Adverse effects: SAS, BAS, COSTART, death, laboratory analytes, ECG. Unable to use: Global state: CGI-S. Adverse effects: change in vital signs (mean, SD CI not reported).

Eli Lilly 2004 (Continued)

Notes

*The age difference between the two groups was statistically significant ($P = 0.011$).

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Described as double-blind, however no further details reported regarding the blinding of participants and personnel are not reported.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Described as double-blind, however no further details reported regarding rater-blinding.
Incomplete outcome data (attrition bias) All outcomes	High risk	The number of patients that completed the study is unclear: authors assert 45 in a study, 46 participants in the other one out of a possible 49 participants. However, only 42 participants are included in the analysis for the CGI-I (at the end of the study), SAS, BAS, clinical laboratory evaluations, vital signs and ECG, without explanation for the missing participants.
Selective reporting (reporting bias)	High risk	Only the CGI-S baseline score is reported.
Other bias	High risk	Sponsored by the manufacturers of olanzapine.

Fang 2012

Methods	Allocation: randomised. Blindness: open. Duration: 120 hours (as for session I).
Participants	Diagnosis: DSM-IV diagnosis of schizophrenia ($N = 192$), schizophreniform ($N = 11$) or schizoaffective ($N = 0$) disorders, no information* ($N = 2$); PANSS-EC ≥ 14 ; PANSS ≥ 60 . $N = 208$ screened, 205 randomised. Age: range 18-45 years. Sex: 99 males, 106 females. History: acute agitation. Excluded: "women pregnant, breastfeeding or planning to become pregnant during the study", "psychotic agitation caused by delirium, epilepsy, mental retardation, or affective disorder", "intoxication or symptoms of withdrawal from alcohol or other psychoactive substances", "serious medical illness", "known hypersensitivity to any of the study medications or no response to risperidone or haloperidol during previous treatment", "treatment with a depot antipsychotic with one cycle of screening", "use of disallowed medication".

Fang 2012 (Continued)

Setting: multi-centre (6), China.

Interventions	<p>1. Haloperidol: flexible dose IM 10 mg/day to 20 mg/day (mean prescribed daily dose 12.2 ± 3.7 mg), N = 101.</p> <p>2. Risperidone: flexible dose oral 2 mL/day to 6 mL/day (mean prescribed daily dose 3.4 ± 0.7 mg), N = 104 + clonazepam: flexible dose oral 0 mg/day to 8 mg/day (mean prescribed daily dose 2.9 ± 1.5 mg), N = 99.</p>
Outcomes	<p>Not asleep.</p> <p>Mental State: PANSS total.</p> <p>Agitation: PANSS-EC.</p> <p>Adverse events: insomnia, tachycardia, EPS, akathisia.</p> <p>Unable to use:</p> <p>Adverse effects: SAS (mean, SE/SD not reported).</p> <p>Adverse effects: BAS (mean, SE/SD not reported).</p>
Notes	<p>*Author assume a typing error in the demographic-clinical "table 1".</p> <p>An attempt to contact the authors was made (2nd July 2016).</p>

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"A telephone-based central service was used to randomly assign eligible patients" (p.108).
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	High risk	Open trial.
Blinding of outcome assessment (detection bias) All outcomes	High risk	Open trial.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete outcome data.
Selective reporting (reporting bias)	High risk	SAS and BAS scores at 2, 4, 24, 72, and 120 hours are not reported. Only side effects occurred in $\geq 5\%$ of people are reported.
Other bias	High risk	Sponsored by drug company (Xian-Janssen Pharmaceutical Ltd).

Fitzgerald 1969

Methods	Allocation: randomised.
---------	-------------------------

Fitzgerald 1969 (Continued)

Blindness: double.

Duration: 48 hours.

Participants	<p>Diagnosis: "acute psychotic behavior". "Eleven of the haloperidol recipients and 12 of the perphenazine recipients exhibited paranoid reactions". (p.515).</p> <p>N = 44.</p> <p>Age: mean 41 years (haloperidol group), 36 years (perphenazine group).</p> <p>Sex: 17 males, 27 females.</p> <p>History: "acutely disturbed. Newly admitted patients exhibiting severe flat affect, inappropriateness, agitation, hostility, overactivity or various combinations". (p.515).</p> <p>Excluded: Not reported.</p> <p>Setting: inpatient, USA.</p>	
Interventions	<p>1. Haloperidol: dose 5 mg/IM (mean number of IMs not reported, mean dose 25.7 mg). N = 23.</p> <p>2. Perphenazine: dose 5 mg/IM (mean number of IMs not reported, mean dose 27.6 mg). N = 21.</p>	
Outcomes	<p>Global state: clinical observation of global improvement.</p> <p>Leaving the study early.</p> <p>Adverse effects: vital signs, need for anti-parkinsonian medication.</p> <p>Not usable:</p> <p>Mental state: Target symptoms (unpublished).</p>	
Notes		
Risk of bias		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	"haloperidol and perphenazine were supplied in identically appearing 1-mL ampoules each containing 5 mg of drug." (p.515).
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Described as double-blind - no further details given regarding rater-blinding.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	It is reported that 3 participants were discontinued on the medication. According to Table 9, not all participants completed all outcome measures, it is not clear if this is due to attrition or because people were not available for assessment.

Fitzgerald 1969 (Continued)

Selective reporting (reporting bias)	Unclear risk	"Most adverse reactions disappeared with the use of control measures" - author suspects that only the adverse effects severe enough for discontinuation are reported.
Other bias	Unclear risk	No clear interested funding.

Foster 1997

Methods	Allocation: randomised. Blindness: double. Duration: 4 hours.
Participants	Diagnosis: DSM-III diagnosis of schizophrenia (N = 13), schizoaffective disorder (N = 4), bipolar disorder (N = 13), psychotic disorder not otherwise specified (N = 7). N = 37. Age: range 18-61 years. Sex: 26 males, 11 females. History: "patients presenting at a psychiatric emergency room service." (p.177). Excluded: Not reported. Setting: psychiatric emergency room, USA.
Interventions	1. Haloperidol: dose 5 mg/oral/IM, repeated at 30 minute intervals if necessary for up to 4 hours (mean number of IMs 2.25). N = 17. 2. Lorazepam: dose 2 mg/oral/IM, repeated at 30 minute intervals if necessary for up to 4 hours (mean number of IMs 1.82). N = 20.
Outcomes	Mental state: BPRS. Unable to use: Global State: need for additional oral/IM medication of study drug (N/SD not reported, mean doses reported). Global state: GCI not published. Behaviour: use of additional benzodiazepine (not reported). Physiological: vital signs (not reported).
Notes	*Author assumes GCI is the same as CGI.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Unclear risk	Not stated.

Foster 1997 (Continued)

Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Described as double-blind, however it is not clear whether the study drugs appeared identical.
Blinding of outcome assessment (detection bias) All outcomes	Low risk	"All raters were unaware of the specific medication received by the patient". (p.176).
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete outcome data.
Selective reporting (reporting bias)	High risk	"patients continued to receive treatment with a benzodiazepine, if necessary, for the duration of their stay in the emergency department." (p.176). The amount was not reported.
Other bias	High risk	Small short study.

Fruensgaard 1977

Methods	Allocation: randomised. Blindness: double. Duration: 72 hours.
Participants	Diagnosis: acute schizophrenia (N = 12), psychogenic psychosis (N = 18). N = 30. Age: range 19-65 years. Sex: 7 males, 23 females. History: "newly admitted patients with a diagnosis of acute psychosis and characterized by symptoms such as agitation, excitement, aggressiveness, delusions and hallucinations." (p.257). Onset < 7 days (N = 14), 1 week - 1 month (N = 13), 1-6 months (N = 3). Excluded: "pregnancy, manic depressive illness, electroconvulsive therapy within the preceding 8 weeks, organic brain syndrome with marked dementia, convulsive disorders, alcoholism or drug dependence, serious impairment of renal, hepatic, cardiovascular or metabolic functions, and present or former increased intra-ocular pressure." (p.257). Setting: psychiatric hospital, Denmark.
Interventions	1. Haloperidol: flexible dose 2.5 mg/IM to 5 mg/IM + biperiden: dose 2.5 mg/IM to 5 mg/IM (mean number of IMs not reported). N = 15. 2. Loxapine: flexible dose 25 mg/IM to 50 mg/IM + biperiden: dose 2.5 mg/IM (mean number of IMs not reported). N = 15.
Outcomes	Adverse effects: recording of side effects, laboratory tests.. Agitation: observation*. Not able to use: Mental state: BPRS (mean reported, SD/CI not reported).

Fruensgaard 1977 (Continued)

Global state: CGI (mean reported, SD/CI not reported).

Adverse effects: blood pressure, pulse rate (reports a decrease in blood pressure/pulse rate in both groups but does not give N/mean/SD/CI).

Notes * Scale derived data for agitation/excitation scores was not included as the measure was not validated.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	"Both drugs were supplied in ampules of identical appearance." (p.257).
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Described as double-blind, however no further details reported regarding rater-blinding.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete data.
Selective reporting (reporting bias)	Low risk	No evidence of selective reporting.
Other bias	High risk	Small study.

Garza-Trevino 1989

Methods	Allocation: randomised. Blindness: open trial. Duration: not reported.
Participants	Diagnosis: Schizophrenia (N = 16), mania (N = 22), "atypical psychotic patients" (N = 6), "miscellaneous" (N = 14). N = 68. Age: Not reported. Sex: 41 males, 27 females. History: "judged to require immediate treatment for agitated or assaultive behavior, with either drugs, seclusion or both." (p.1599). Excluded: Not reported. Setting: psychiatric hospital, USA.

Garza-Trevino 1989 (Continued)

Interventions	1. Haloperidol: dose 5 mg/IM (mean number of IMs not reported). N = 21. 2. Haloperidol: dose 5 mg/IM + lorazepam 4 mg/IM (mean number of IMs not reported). N = 24. 3. Lorazepam: dose 4 mg/IM (mean number of IMs not reported). N = 23.	
Outcomes	Agitation: VAS.	
Notes		
<i>Risk of bias</i>		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	High risk	Open trial.
Blinding of outcome assessment (detection bias) All outcomes	High risk	Open trial.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete data.
Selective reporting (reporting bias)	High risk	There is no reporting regarding the incidence of adverse effects.
Other bias	Unclear risk	The duration of the study is not reported.

Guo 2007

Methods	Allocation: randomised. Blindness: open trial. Duration: 14 days.
Participants	Diagnosis: CCMD-3 diagnosis of schizophrenia with agitation/aggression. N = 60. Age: range 18-49 years. Sex: males and females. History: mean length of illness 56.7 (SD 7.8) months. Excluded: severe physical illness, drug/alcohol dependence.

Guo 2007 (Continued)

Setting: inpatient, China.

Interventions	<p>Haloperidol: dose 5 mg/day to 10 mg/day for 3 days, 10 mg/IM/day to 30 mg/IM/day afterwards (mean number of IMs not reported). N = 30.</p> <p>Quetiapine: dose 100 mg/day to 800 mg/day + magnesium valproate dose: 0.5 g/p.o./day to 1 g/p.o./day (mean number of IMs not reported) N = 30.</p>
Outcomes	<p>Agitation: PANSS-EC.</p> <p>Not able to use:</p> <p>Global outcome: CGI (not outcome data reported).</p> <p>Adverse effects: TESS (no outcome data reported).</p>
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	High risk	Open-label.
Blinding of outcome assessment (detection bias) All outcomes	High risk	Open-label.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No incomplete outcome data
Selective reporting (reporting bias)	High risk	CGI and TESS were measured, but not reported.
Other bias	Unclear risk	None obvious.

Higashima 2004

Methods	<p>Allocation: randomised.</p> <p>Blindness: open trial.</p> <p>Duration: 8 weeks.</p>
Participants	<p>Diagnosis: DSM-IV diagnosis of schizophrenia.</p> <p>N = 19.</p>

Higashima 2004 (Continued)

Age: mean 29.5 years (monotherapy), mean 26.6 years (combined therapy).

Sex: 12 males, 7 females.

History: "acute exacerbation of psychotic symptoms." (p.380).

Excluded: Not reported.

Setting: Psychiatric hospital, Japan.

Interventions	Haloperidol: average endpoint dose 5.4 (2.7) mg/p.o./day. N = 10. Haloperidol + levomepromazine: average endpoint dose ratio 1:10, 5.4 (2.4) mg/p.o./day. N = 9.
Outcomes	Global state: no overall improvement. Unable to use: Mental state: BPRS. Adverse effects: not reported. Adverse effects: the quantity of anticholinergic medication prescribed was not reported.

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	High risk	Open trial.
Blinding of outcome assessment (detection bias) All outcomes	High risk	Open trial.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete data.
Selective reporting (reporting bias)	High risk	The discussion describes how the combination drug "produced moderate sedation without any cardiovascular adverse effect". However, there is no other reporting regarding the incidence adverse effects.
Other bias	High risk	Small N study.

Huf 2007

Methods	Allocation: randomised.
---------	-------------------------

Huf 2007 (Continued)

Blindness: open trial.

Duration: 14 days.

Participants	<p>Diagnosis: psychosis (N = 244), substance misuse (N = 58), others (N = 11), no information (N = 3).</p> <p>N = 316.</p> <p>Age: mean 40.1 years.</p> <p>Sex: males.</p> <p>History: people presenting at a psychiatric emergency room service. 1st attendance (N = 59), previous attendance (N = 224), unknown history (N = 33).</p> <p>Excluded: where the "clinician believed that one of the treatment options represented an additional risk for the patient." (p.3).</p> <p>Setting: psychiatric emergency room, Brazil.</p>
Interventions	<p>1. Haloperidol: dose 5 mg/IM to 10 mg/IM. N = 156.</p> <p>2. Haloperidol: dose 5 mg/IM to 10 mg/IM + promethazine 25 mg/IM to 50 mg/IM. N = 160.</p>
Outcomes	<p>Tranquillisation or asleep: tranquil/asleep, time to tranquillisation/sleep, need for additional tranquillising drug.</p> <p>Specific behaviours: another episode of aggression by 24 hours.</p> <p>Global state: use of restraints, needing extra visits from the doctor, refusing oral medication.</p> <p>Leaving the study early.</p> <p>Service outcomes: duration of hospital stay.</p> <p>Adverse effects: serious, specific adverse effects.</p>

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	A computer-generated randomised sequence was applied to a table of numbers.
Allocation concealment (selection bias)	Low risk	"Boxes were consecutively numbered, sealed, opaque, and identical appearance and weight; on the outside was a form with questions to be completed by the attending doctor while "blind" to the contents of the box." (p.2).
Blinding of participants and personnel (performance bias) All outcomes	High risk	Open trial.
Blinding of outcome assessment (detection bias) All outcomes	High risk	Open trial.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete outcome data.

Huf 2007 (Continued)

Selective reporting (reporting bias)	Low risk	No evidence of selective reporting.
Other bias	Unclear risk	The study was stopped early due to adverse effects, therefore it is possible that the available data may be biased.

Kewala 1984

Methods	Allocation: randomised. Blindness: double. Duration: 5 hours.
Participants	Diagnosis: DSM-III diagnosis of schizophrenia (N = 13), schizoaffective disorder (N = 4), bipolar disorder (N = 13), psychotic disorder not otherwise specified (N = 7). N = 44. Age: range 19-56 years. Sex: 21 males, 23 females. History: "newly admitted patients...mean duration of illness 7.75 years." (p.77). Excluded: Not reported. Setting: psychiatric hospital, USA.
Interventions	1. Haloperidol: flexible dose 2.5 mg/IM to 10 mg/IM, maximum 4 injections/cumulative dosage of 27.5 mg during 24 hours (mean number of IMs not reported, mean cumulative dose ~ 12.1 mg). N = 24. 2. Thiothixene: flexible dose 2.5 mg/IM to 10 mg/IM, maximum 4 injections/cumulative dosage of 27.5 mg during 24 hours (mean number of IMs not reported, mean cumulative dose ~ 16.1 mg). N = 20.
Outcomes	Global state: global evaluation. Adverse effects. Unable to use: Mental state: BPRS (SE/SD/CI not reported, mean reported). Mental state: Study Symptom Profile (mean, SE/SD not reported, P values of significant findings reported). Physiological: the authors report that there were no abnormal laboratory results, although no data was given.

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported

Kewala 1984 (Continued)

Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Described as double-blind, however no further details reported regarding the blinding of participants and personnel.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Described as double-blind, however no further details reported regarding rater-blinding.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete outcome data.
Selective reporting (reporting bias)	Unclear risk	No evidence of selective reporting.
Other bias	High risk	Received grant from Roerig, a division of Pfizer Pharmaceuticals (thiothixene manufacturers), small short study.

Kinon 2001d

Methods	Allocation: randomised. Blindness: double. Duration: 21 days.
Participants	Diagnosis: schizophrenia (N = 56), schizophreniform disorder (N = 42), schizoaffective disorder (N = 2). N = 100. Age: range 18-50 years. Sex: 71 males, 29 females. History: "newly admitted patients who were acutely agitated." (p.181). Excluded: pregnant or lactating women, people with "serious unstable illnesses, including hepatic, renal, gastroenterologic, respiratory, cardiovascular, endocrinologic, neurologic, immunologic, or hematologic disease, in which pharmacotherapy posed a substantial clinical risk or confounded diagnosis were excluded from this trial." (p.181). Setting: not reported.
Interventions	1. Haloperidol: accelerated dose titration 10 mg/p.o. to 20 mg/p.o. during 24 hours. N = 48. 2. Olanzapine: accelerated dose titration 10 mg/p.o. to 20 mg/p.o. during 24 hours. N = 52.
Outcomes	Mental state: PANSS total, need for increased dose of study drug. Global state: CGI-I, time in restraints, seclusion and special observation, need for increased dose of study drug, use of benzodiazepine. Agitation: Seclusion and special observation. Leaving the study early.

Kinon 2001d (Continued)

Adverse effects: Tranquillisation Scale, recording rates of discontinuation, recording treatment emergent adverse events in $\geq 10\%$ of participants, need for parkinsonian drugs.

Unable to use:

Global state: CGI (between-group data was described as not significant but figures not reported - only within-group data reported), CGI-S (not reported).

Agitation: OASS (only present the data from 2 items of the scale), NOSIE (between-group data were described as not significant but figures not reported - only within-group data reported), PANSS Agitation sub-scale (only present data from certain items of the PANSS), SAS (between-group data were described as not significant but figures not reported - only within-group data reported).

Adverse effects: BAS (between-group data was described as not significant but figures not reported - only within-group data reported), vital signs (not reported), laboratory analytes (described as no clinically relevant changes, although no data given), weight gain (mean, P values of significant findings, overall F value reported, SD/SE/CI not reported).

Satisfaction: DAI-10 (significant within-group data reported, for between-group P values of non significant findings reported, overall F value).

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Described as double-blind, however no further details reported regarding the blinding of participants and personnel.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Described as double-blind, however no further details reported regarding rater-blinding.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete outcome data.
Selective reporting (reporting bias)	High risk	Significant findings from the within-group data are presented on a number of occasions, whilst acknowledging but omitting the non-significant between-group findings. The OASS is a 16 item rating scale, yet only the results for two items where the olanzapine group had significantly greater improvement than the haloperidol group were reported. Treatment-emergent adverse effects were only reported if they occurred with a frequency $\geq 10\%$ or $P < .05$ through the end of the study.
Other bias	High risk	Sponsored by the manufacturers of olanzapine.

Li 2006

Methods	Allocation: randomised. Blindness: single. Duration: 72 hours.
Participants	Diagnosis: CCMD-3 diagnosis of schizophrenia with agitation/aggression. N = 231. Age: mean 34 (SD 12) years. Sex: males and females. History: mean length of illness, haloperidol group 30 ($\pm$ 68) months, ziprasidone 18 ($\pm$ 35) months. Excluded: severe physical or neurological impairment/diseases; drug or alcohol abuse, dependence, pregnant women. Setting: not reported.
Interventions	1. Haloperidol: flexible dose 5 mg/IM to 10 mg/IM, repeated every 4-6 hours if necessary, maximum dose 30 mg during 24 hours. N = 116. 2. Ziprasidone: flexible dose 10 mg/IM to 20 mg/IM, repeated every 4-6 hours if necessary, maximum dose 40 mg during 24 hours. N = 115.
Outcomes	Mental state: PANSS endpoint. Agitation: PANSS-EC. Adverse effects: TESS.
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Single-blind – however the author did not state who had been blinded.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Single-blind – however the author did not state who had been blinded.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No incomplete data.

Li 2006 (Continued)

Selective reporting (reporting bias)	Unclear risk	TESS and SAS were used for assessment of adverse events. No continuous data were reported from either scales, but binary outcomes for adverse events were reported.
Other bias	Unclear risk	None obvious.

Liang 2013

Methods	Allocation: randomised. Blinding: not stated. Duration: 28 days.
Participants	Diagnosis: ICD-10 diagnosis of schizophrenia. N = 80. Age: range 18-60 years. Sex: 42 males, 38 females. History: acute agitation and excitement; length of illness 7.1 ± 7.7 (quetiapine group), 7.0 ± 6.3 (haloperidol group). Excluded: "pregnancy, serious physical sickness, addiction to alcohol or drugs, intolerant to quetiapine, haloperidol or orally taking medicines, taking quetiapine or haloperidol or other long-term schizophrenia medicines one month before recruitment, neutrophil cells $\leq 1.5 \times 10^9/L$ ". Setting: inpatients, China.
Interventions	1. Haloperidol: dose 2 mg/oral/day to 4 mg/oral/day, by 3rd day 6 mg/day, by 7th day 10 mg/day to 12 mg/day. N = 40. 2. Quetiapine: dose 50 100 mg/oral/day to 100 mg/oral/day, by 3rd day 300 mg/day, by 7th day 600 mg/day to 750 mg/day. N = 40.
Outcomes	Mental state: PANSS total and sub-scale. Agitation: PANSS-EC. Adverse effects: general. Adverse effects: movement - SAS. Dropout from the study.
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"randomised by using random number table".
Allocation concealment (selection bias)	Unclear risk	Not stated.

Liang 2013 (Continued)

Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Not stated.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Not stated.
Incomplete outcome data (attrition bias) All outcomes	High risk	A total of 5 dropped out in haloperidol group (adverse events = 4, lost to contact = 1) and 4 dropped out in quetiapine group (adverse events = 1, lost to contact = 3); missing data have not been imputed using appropriate methods such as LOCF.
Selective reporting (reporting bias)	Low risk	No evidence of selective reporting.
Other bias	Unclear risk	None obvious.

Lim 2010

Methods	Allocation: randomised. Blindness: open-label, rater-blinded. Duration: 24 hours.
Participants	Diagnosis: DSM-IV diagnosis of schizophrenia (N = 78), bipolar disorder (N = 43), others (N = 3). N = 124. Age: range 18-65 years. Sex: 66 males, 58 females. History: "acute psychotic agitation in the emergency room." (p.82). Excluded: neurological disorder, severe medical disease, alcohol or other psychoactive substance misusers, history of neuromalignant syndrome or hypersensitivity to trial medications, pregnant or lactating women, people treated with antipsychotics or benzodiazepines within 6 hours to the start of the trial or with depot antipsychotic within one treatment cycle of enrolment. Setting: psychiatric emergency room, Korea.
Interventions	1. Haloperidol: dose 5 mg/IM, maximum 15 mg during 24 hours (mean number of IMs not reported). N = 62. 2. Risperidone: dose 2 mg/p.o, maximum 6 mg during 24 hours (mean dose not reported). N = 62.
Outcomes	Global state: need for benzodiazepine. Agitation: PANSS-EC. Leaving the study early. Adverse effects. Unable to use:

Lim 2010 (Continued)

Mental state: PANSS Total (mean, SE/SD not reported, P values of significant findings reported, overall F value).

Mental state: YMRS Total (mean, SE/SD/CI not reported).

Global state: CGI-S (mean, SE/SD not reported, P values of significant findings reported, overall F value).

Adverse effects: use of antiparkinsonian drugs not reported.

Adverse effects: AIMS (mean, SE/SD/CI not reported).

Adverse effects: SAS (mean, SE/SD/CI not reported).

Behaviour: Behaviorally Anchored Rating Scale (mean, SE/SD/CI not reported).

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Participants were assigned "according to a predefined randomization code that was balanced to ensure even distribution of patients in each treatment group." (p.82).
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	High risk	This was an open-label study where one group received IM and the other group received an orodispersible tablet.
Blinding of outcome assessment (detection bias) All outcomes	Low risk	The study was rater-blinded.
Incomplete outcome data (attrition bias) All outcomes	Low risk	There is no evidence of incomplete outcome data. The authors give reasons for the participants who discontinued the study.
Selective reporting (reporting bias)	High risk	The authors report that antiparkinsonian drugs could be given at the lowest effective dose, however the amount administered is not described in the results.
Other bias	High risk	Sponsored by drug company (Janssen Pharmaceutica Korea).

Liu 2012

Methods	Allocation: randomised. Blinding: not stated. Duration: 2 weeks.
Participants	Diagnosis: CCMD-3 diagnosis of schizophrenia. N = 150.

Haloperidol for psychosis-induced aggression or agitation (rapid tranquillisation) (Review)

Liu 2012 (Continued)

Age: range 18-59 years.

Sex: 69 males, 81 females.

History: agitated behaviour; length of illness 0- 30 months.

Excluded: "serious physical sickness, additive to alcohol and drugs, intolerant to schizophrenia treatment, pregnancy".

Setting: inpatients, China.

Interventions	<p>1. Haloperidol: dose 10 mg/IM/day to 20 mg/IM/day (mean dose 16.15 ± 4.67 mg/day). N = 50.</p> <p>2. Risperidone: dose 3 mg/oral/day to 6 mg/oral/day (mean dose 4.57 ± 0.65 mg/day) + haloperidol dose 10 mg/IM/day to 20 mg/IM/day (mean dose 15.73 ± 4.26 mg/day). N = 50.</p> <p>3. Risperidone: dose 3 mg/oral/day to 6 mg/oral/day + clonazepam dose 3 mg/IM/day to 6 mg/IM/day (mean dose 4.05 ± 0.81 mg/day). N = 50.</p> <p>All IM injections changed to oral administration by 10 days.</p>
Outcomes	<p>Global state: clinical observation of global improvement.</p> <p>Agitation: PANSS-EC.</p> <p>Adverse effects: TESS.</p>
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"randomised by using random number table".
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Not stated.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Not stated.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No incomplete data.
Selective reporting (reporting bias)	Low risk	No evidence of selective reporting.
Other bias	Low risk	None obvious.

Man 1973

Methods	Allocation: not reported. Blindness: double. Duration: 72 hours.
Participants	Diagnosis: acute exacerbation of chronic psychosis - "severe agitation with marked psychomotor hyperactivity, extreme assaultiveness, hostility and mania. The patients presented a clear danger to themselves, to other patients and to staff personnel". N = 30. Age: mean 33 years. Sex: 15 males and 15 females. History: acute agitation, 90% had previous admission to a psychiatric hospital. Excluded: anyone who used anti-psychotic or anti-convulsant within one week prior to the trial or anyone who used long lasting drug within one month prior to the trial. Setting: inpatients, China.
Interventions	1. Haloperidol: dose 5 mg/IM, repeated every 30 minutes if necessary, maximum 15 mg during 24 hours (mean number of IMs 3.33). N = 15. 2. Chlorpromazine: dose 50 mg/IM, repeated every 30 minutes if necessary, maximum 600 mg during 24 hours (mean number of IMs 2.93). N = 15.
Outcomes	Global state: need for additional injections. Adverse effects: reporting of EPS. Leaving the study early. Unable to use: Mental state: BPRS (SD/SE/CI not reported, mean not reported). Global state: TSRS (unpublished, mean reported, SD/SE/CI not reported). Adverse effects: it is reported that there were no significant changes in blood pressure, pulse rate and respirations, however no data given.

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	High risk	Not explicitly described as randomised.
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Described as double-blind, however it is unclear whether the syringes appeared identical given the varied doses.

Man 1973 (Continued)

Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Described as double-blind, however no further details reported regarding rater blinding.
Incomplete outcome data (attrition bias) All outcomes	High risk	The authors give reasons for the participants who discontinued the study. However, the results are only presented for the 27 participants who completed the study. Missing data have not been imputed using appropriate methods such as LOCF.
Selective reporting (reporting bias)	Low risk	No evidence of selective reporting.
Other bias	High risk	Small study.

Nobay 2004

Methods	Allocation: randomised. Blindness: double. Duration: not reported.
Participants	Diagnosis: violent behaviour and severe agitation. N = 111. Age: mean 40.7 years. Sex: not reported. History: prior psychiatric history (N = 54), no psychiatric history (N = 8), unknown psychiatric history (N = 49). Excluded: known allergy to study drugs, hypotensive, unable to maintain own airway, tachycardia, bradycardia, respiratory distress, <18 or pregnant. Setting: emergency psychiatric department, USA.
Interventions	1. Haloperidol: fixed dose 5 mg/IM (mean number of IMs not reported). N = 42. 2. Midazolam: fixed dose 5 mg/IM (mean number of IMs not reported). N = 42.*
Outcomes	Tranquil or asleep: modified TCS. Global state: need for additional medication. Adverse effects. Unable to use: Adverse effects: it is reported that there were no significant differences between groups for changes to blood pressure, heart rate and oxygen. However, data not given.
Notes	*There was also a lorazepam group, however we cannot include the data because randomisation was broken.

Risk of bias

Bias	Authors' judgement	Support for judgement
------	--------------------	-----------------------

Haloperidol for psychosis-induced aggression or agitation (rapid tranquillisation) (Review)

139

Nobay 2004 (Continued)

Random sequence generation (selection bias)	Low risk	"The administered drugs were prepared in the pharmacy using a computer generated randomisation code." (p.745).
Allocation concealment (selection bias)	Low risk	"The administered drugs were prepared in the pharmacy using a computer generated randomisation code." (p.745).
Blinding of participants and personnel (performance bias) All outcomes	Low risk	"The research assistant [rater], the administering physician, and the patient were all blinded to the drug delivered." (p.745).
Blinding of outcome assessment (detection bias) All outcomes	Low risk	"The research assistant [rater], the administering physician, and the patient were all blinded to the drug delivered." (p.745).
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete outcome data.
Selective reporting (reporting bias)	Low risk	No evidence of selective reporting.
Other bias	High risk	The duration of the study was not reported, therefore it is difficult to establish how long participants were monitored for adverse effects.

Paprocki 1977

Methods	Allocation: randomised. Blindness: double. Duration: 4-day IM phase followed by up to 4 week oral phase.
Participants	Diagnosis: "acute psychosis with manifest disturbed psychotic symptoms characterised by agitation, excitement, aggressiveness, hostility, delusions and hallucinations." (p.81). N = 35. Age: range 18-49 years. Sex: females. History: onset of present episode: < 1 week (N = 10), > 1 week < 1 month (N = 26)*. Excluded: psychotropic drugs prior to 24 hours of the start of the study, "known hypersensitivity to dibenzazepine compounds, electroconvulsive therapy or insulin coma or subcoma therapy within the preceding eight weeks, presence of organic brain syndrome with marked dementia and inability to communicate during interviews, history of convulsive disorders, alcoholism or drug dependence as a significant feature of clinical history, presence of serious impairment of renal or hepatic function, increased intra ocular pressure or history of narrow angle glaucoma or urinary retention, cardiovascular or metabolic disease, pregnancy suspected or confirmed." (p.82). Setting: emergency psychiatric department, USA.
Interventions	1. Haloperidol: dose 2.5 mg/IM to 5 mg/IM, repeated every 6 to 12 hours if necessary (IM phase), dose 5 mg to 15 mg (oral phase) (mean number of IMs not reported, mean daily dose 11.5 mg). N = 18.

Paprocki 1977 (Continued)

2. Loxapine: dose 25 mg/IM to 50 mg/IM, repeated every 6 to 12 hours if necessary (IM phase), dose 50 mg to 150 mg (oral phase) (mean number of IMs not reported, mean daily dose 115.4 mg). N = 17.

Outcomes	<p>Global state: CGI.</p> <p>Leaving the study early</p> <p>Adverse effects: observations, laboratory tests.</p> <p>Unable to use:</p> <p>Mental state: BPRS (overall P value and level of significance reported, SD/SE/CI not reported).</p> <p>Global state: NOSIE (overall P value and level of significance reported, SD/SE/CI not reported).</p> <p>Adverse effects: sedation score (mean reported, SD/SE/CI not reported).</p>
Notes	*This would suggest there were 36 participants. It is not clear if this is a typing error because it is reported in the rest of the paper that there were 35 participants.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Described as double-blind - "supplied in 1ml ampules...of identical appearance, individually packaged and labelled with the code number assigned to each patient". Capsules used in the oral phase were also of identical appearance.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Described as double-blind - no further details given regarding rater blinding.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	The number of participants reported is inconsistent (reported at 36 in one instance and 35 at other times) and it is not clear if this is a typing error.
Selective reporting (reporting bias)	Low risk	No evidence of selective reporting.
Other bias	High risk	Small study.

Reschke 1974

Methods	<p>Allocation: randomised.</p> <p>Blinding: double for IM phase, open for oral phase.</p> <p>Duration: approximately 2 hours.</p>
Participants	<p>Diagnosis: schizophrenia (no specified criteria).</p> <p>N = 50.</p> <p>Sex: 2 males, 48 females.</p> <p>Age: range 19-57, mean 35.9 years.</p>

Haloperidol for psychosis-induced aggression or agitation (rapid tranquillisation) (Review)

Reschke 1974 (Continued)

History: severe agitation - "Psychiatric emergencies with symptoms such as agitation, excitement, and assaultiveness of such severity that rapid tranquillisation was necessary".

Excluded: not reported.

Setting: hospital.

Interventions	<p>1. Haloperidol IM: dose up to 4 injections of 1 mg (mean number of IMs not reported). N = 10.</p> <p>2. Haloperidol IM: dose up to 4 injections of 2 mg (mean number of IMs 3.7). N = 11.</p> <p>3. Haloperidol IM: dose up to 4 injections of 5 mg (mean number of IMs 2.8). N = 8.</p> <p>4. Chlorpromazine IM: dose up to 4 injections of 25 mg (mean number of IMs not reported). N = 10.</p> <p>5. Placebo IM: dose up to 4 injections of 1 mg (mean number of IMs not reported). N = 11.</p>
Outcomes	<p>Not tranquil or asleep: not asleep at 2 hours.</p> <p>Global state: own 5-point rating scale.</p> <p>Adverse effects.</p> <p>Leaving the study early.</p> <p>Unable to use:</p> <p>Mental state: BPRS and own target symptoms scale (no mean or SD).</p>
Notes	<p>Data for the haloperidol interventions were combined within the paper for some outcomes, for the binary outcomes that were not already presented in this way we added events for the different doses together following the protocol (see Unit of analysis issues).</p>

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Described as double-blind, however no further details are given. It is not clear whether the study drugs appeared identical.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Described as double-blind, however no further details reported regarding rater blinding.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete outcome data.
Selective reporting (reporting bias)	Low risk	No evidence of selective reporting.
Other bias	Unclear risk	Small study.

Resnick 1984

Methods	Allocation: randomised. Blindness: double. Duration: 24 hours.
Participants	Diagnosis: acute agitation. N = 27. Age: range 18-65 years. Sex: not reported. History: involuntary hospitalised. Excluded: intoxicated, known sensitivity to droperidol or haloperidol, evidence of active renal, hepatic, or cardiac disease. Setting: emergency and psychiatric crisis unit, USA.
Interventions	1. Haloperidol: dose up to 4 injections of 5 mg (mean number of IMs not reported). N = 16. 2. Droperidol: dose up to 4 injections of 5 mg (mean number of IMs 1.36). N = 11.
Outcomes	Global state: need for additional injection. Adverse effects: reporting of EPS. Unable to use: Mental state: BPRS (mean/SD/SE/CI not reported). Adverse effects: it is reported that there were no significant changes in blood pressure, pulse rate and respirations, however no data given.

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Low risk	"packages of medication were identified only by a code which was kept in the pharmacy until the conclusion of the study" (p.298).
Blinding of participants and personnel (performance bias) All outcomes	Low risk	"packages of medication were identified only by a code which was kept in the pharmacy until the conclusion of the study" (p.298).
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Described as double-blind, however no further details reported regarding rater blinding.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete data.

Resnick 1984 (Continued)

Selective reporting (re-reporting bias)	Low risk	No evidence of selective reporting.
Other bias	High risk	Small short study.

Ritter 1972

Methods	Allocation: randomised, no further details. Blindness: double, identical appearing ampoules. Duration: 48 hours.
Participants	Diagnosis: paranoid schizophrenia (N = 48), involuntal melancholia (N = 1), manic-depression (N = 1). N = 50. Age: range 21-65 years, mean 36 years. Sex: males. History: all exhibited symptoms such as aggressiveness, assaultiveness or uncooperativeness that disrupted normal hospital operation and interfered with attempts at therapy. Excluded: not reported. Setting: not reported.
Interventions	1. Haloperidol: fixed dose 5 mg/IM, repeated every 6-8 hours as necessary (mean number of IMs not reported). N = 25. 2. Chlorpromazine: fixed dose 50 mg/IM, repeated every 6-8 hours as necessary (mean number of IMs not reported). N = 25.
Outcomes	Global state: own 5-point scale. Adverse events: Leucopenia, SGPT increase. Unable to use: Adverse effects: EPS, hypotension (no data).
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Described as double-blind - the test drugs were supplied in identically appearing ampoules.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Described as double-blind, however no further details reported regarding rater blinding.
Incomplete outcome data (attrition bias)	Low risk	No evidence of incomplete outcome data.

Ritter 1972 (Continued)

All outcomes

Selective reporting (reporting bias)	High risk	Do not report all adverse effects. The authors do not report the mean dosages/number of injections needed.
Other bias	Unclear risk	None obvious.

Salzman 1991

Methods	Allocation: not reported. Blindness: double. Duration: 48 hours.
Participants	Diagnosis: schizophrenia (N = 26), schizoaffective disorder (N = 4), bipolar disorder (N = 11), organic mental disorder (N = 6), other (N = 13). N = 60. Age: mean 30.5 years (haloperidol group), mean 37.9 years (lorazepam group). Sex: not reported. History: involuntary hospitalised - "Aggressive, assaultive or disruptive behaviour, thought to be a danger to the patient or other people". Excluded: intoxicated, known sensitivity to droperidol or haloperidol, evidence of active renal, hepatic, or cardiac disease. Setting: emergency and psychiatric crisis unit, USA.
Interventions	1. Haloperidol: fixed dose 5 mg/IM (mean number of IMs 1.1). N = 30. 2. Lorazepam: fixed dose 2 mg/IM (mean number of IMs 1.13). N = 30.
Outcomes	Adverse effects. Unable to use: Mental state: BPRS (mean change given, SD/SE/CI not reported). Global state: CGI-S (mean change given, SD/SE/CI not reported). Aggression: OAS (mean change given, SD/SE/CI not reported). Global state: additional injection (mean dose given, SD/SE/CI not reported).

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	High risk	Not explicitly described as randomised.
Allocation concealment (selection bias)	Unclear risk	Not stated.

Salzman 1991 (Continued)

Blinding of participants and personnel (performance bias) All outcomes	Low risk	"Parenteral medication was prepared in advance by a member of staff who had no direct clinical responsibility for the patient." (p.177).
Blinding of outcome assessment (detection bias) All outcomes	Low risk	"Raters were blind to the treatment condition received by the patient." (p.178).
Incomplete outcome data (attrition bias) All outcomes	High risk	It is reported that the number of participants declined after the baseline assessment because they were discharged or transferred. However, exact numbers are not given and it is not clear if the participants were transferred because they had a positive or negative outcome. Attrition approaching 50% for some outcomes at 24 hours.
Selective reporting (reporting bias)	Low risk	No evidence of selective reporting.
Other bias	High risk	Sponsored by drug company.

Shi 2010

Methods	Allocation: randomised. Blindness: not stated. Duration: 2 weeks.
Participants	Diagnosis: CCMD-3 diagnosis of schizophrenia. N = 64. Age: range 18-50 years. Sex: 37 males, 27 females. History: agitation and aggression. Excluded: serious physical illness; addiction to alcohol or drugs; pregnancy. Setting: inpatients, China.
Interventions	1. Haloperidol IM (mean dose 15 ± 2.5 mg/day). N = 31. 2. Olanzapine oral (mean dose 10 ± 2.5 mg/day) + magnesium valproate oral (mean 1.25 ± 0.25 g/day). N = 33.
Outcomes	Global state: clinical observation of global improvement. Mental state: PANSS total score. Agitation: PANSS-EC. Adverse effects.
Notes	
Risk of bias	

Shi 2010 (Continued)

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"randomised by using random number table".
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Not stated.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Not stated.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete outcome data.
Selective reporting (reporting bias)	Low risk	No evidence of selective reporting.
Other bias	Low risk	None obvious.

Shu 2010

Methods	Allocation: randomised. Blindness: open-label, rater-blinded. Duration: 72 hours.
Participants	Diagnosis: ICD-10 diagnosis of schizophrenia. N = 376. Age: range 18-65 years. Sex: males and females. History: mean duration of acute agitation 3.4 days (haloperidol group), 3.2 days (ziprasidone group); mean duration of schizophrenia 4 years (haloperidol group), 5.6 years (ziprasidone group). Excluded: history of clinically significant physical illness especially myocardial infarction, non compensatory heart failure, people receiving an investigational agent in the previous 3 months to screening, use of antipsychotic agents within 12 hours or parenteral benzodiazepines within 4 hours prior to randomisation and during the study. Setting: multi-centre, China.
Interventions	1. Haloperidol: dose 5 mg/IM, maximum 20 mg during 24 hours (mean number of IMs not reported). N = 187. 2. Ziprasidone: flexible dose 10 mg/IM to 20 mg/IM, maximum 40 mg in 24 hours (mean number of IMs not reported). N = 189.

Shu 2010 (Continued)

Outcomes	<p>Mental state: BPRS.</p> <p>Adverse effects: laboratory tests, ECG's, physical examination.</p> <p>Leaving the study early.</p> <p>Unable to use:</p> <p>Specific behaviour: agitation - BPRS agitation items (modified from BPRS, only report LS means; means/SD not reported).</p> <p>Specific behaviour: agitation - BARS (only give upper quartile value and LS means; means/SD not reported).</p> <p>Global state: CGI-I (only report CI and LS means, means/SD not reported).</p> <p>Global state: CGI-S (only give upper quartile value and LS means, means/SD not reported).</p> <p>Adverse effects: BAS and SAS (only report LS means).</p>
----------	--

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	"randomized in a 1:1 ratio (in blocks of 4)" (p.179); no further informations on the randomization procedure is stated.
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	High risk	Open-label.
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Rater-blinded.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	The reasons for leaving the study early were reported for those participants who did not complete the study. However, the number of participants completing different assessments at the same time point vary and reasons are not given.
Selective reporting (reporting bias)	Unclear risk	Only report side effects experienced by $\geq 2\%$ of people.
Other bias	High risk	Sponsored by drug company.

Stotsky 1977

Methods	<p>Allocation: randomised.</p> <p>Blindness: double.</p> <p>Duration: 6 hours.</p>
---------	--

Stotsky 1977 (Continued)

Participants	<p>Diagnosis: schizophrenia (N = 24), bipolar disorder (N = 2), psychotic depressive reaction (N = 4).</p> <p>N = 30.</p> <p>Age: range 18-66 years.</p> <p>Sex: 13 males, 17 females.</p> <p>History: disturbed, acutely disruptive symptomology outpatients, 16 of whom had prior inpatient treatment, and 21 of whom had a history of outpatient treatment.</p> <p>Excluded: known hypersensitivity to haloperidol or thiothixene, active hepatic, renal or cardiac disease or organic brain disease.</p> <p>Setting: emergency and psychiatric crisis unit, USA.</p>
Interventions	<p>1. Haloperidol: flexible dose 4 mg/IM or 8 mg/IM, maximum dose 32 mg during 6 hours (mean number of IMs 3.07, mean dose 15 mg). N = 15.</p> <p>2. Thiothixene: flexible dose 4 mg/IM or 8 mg/IM, maximum dose 32 mg during 6 hours (mean number of IMs 2.6, mean dose 10.1 mg). N = 15.</p>
Outcomes	<p>Global state: need for additional injection.</p> <p>Adverse effects.</p> <p>Unable to use:</p> <p>Mental state: BPRS (mean scores given, SD/SE/CI not reported).</p> <p>Mental state: symptom profile (not a validated scale).</p> <p>Adverse effects: vital signs (reports significant between-group changes but does not give data).</p> <p>Global state: mean dose given (SD/SE/CI not reported).</p>
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Randomised in blocks of 10. Method of sequence generation not clear.
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	"Medication was provided in identically appearing 2cc. ampules each containing 4 mg's. Prior to the administration of the drug labels were removed by a person not involved either in the prescription or administration of the drug." (p.968).
Blinding of outcome assessment (detection bias) All outcomes	Low risk	"Ratings were performed by the psychiatrist ordering the medication who was not aware which drug was administered to the patient. When asked to guess which drug was given, the psychiatrists administering the drug were unable to guess significantly better than chance." (p.968).
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete outcome data.

Stotsky 1977 (Continued)

Selective reporting (reporting bias)	Low risk	No evidence of selective reporting.
Other bias	High risk	Small short study, sponsored by drug company.

Taymeeyapradit 2002

Methods	Allocation: randomised. Blindness: present for some outcomes. Duration: 7 days.
Participants	Diagnoses: ICD-10 diagnosis of schizophrenia with acute exacerbation of symptoms (N = 39), mania (N = 1), other (N = 6). N = 70. Age: range 18-65 years. Sex: males and females. History: disturbed and aggressive behaviour, unresponsive to verbal intervention. Mean onset of acute episode, 20.09 days (haloperidol group), 13.63 days (zuclopenthixol group). Excluded: psychosis caused by medication, medical condition, learning disability, people who had anti-psychotic in the previous 2 weeks. Setting: psychiatric hospital, Thailand.
Interventions	1. Haloperidol IM: flexible dose 5 mg to 10 mg repeated every 6 hours as required (mean number of IMs 5). N = 32. 2. Zuclopenthixol acetate IM: flexible dose 50 mg to 100 mg repeated every 12 hours as required (mean number of IMs 2). N = 38. Oral antipsychotics and mood stabilisers allowed.
Outcomes	Global state: additional injections. Adverse effects: presence of tremor, reaction at injection site. Unable to use - Mental state: BPRS, BRMS (means, inadequate data on SD). Global state: CGI (means, no SD).

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Unclear risk	Not stated.

Taymeeyapradit 2002 (Continued)

Blinding of participants and personnel (performance bias) All outcomes	High risk	Open-label.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	The BPRS was completed by a psychiatric nurse who was blind to the intervention. However, it is not reported whether the rater for CGI and adverse effects was blinded.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete outcome data.
Selective reporting (reporting bias)	Low risk	No evidence of selective reporting.
Other bias	Unclear risk	None obvious.

Tuason 1986

Methods	Allocation: randomised. Blinding: "modified double" - staff administering drugs not blinded, assessments blinded. Duration: 24-72 hours (IM phase) - oral phase data not included.
Participants	Diagnosis: schizophrenia, acutely psychotic (DSM-III used beyond 3 days). N = 54*. Age: range 18-65, mean 35 (SD 10) years. Sex: 33 males, 19 females, 2 not reported. History: newly admitted. Excluded: ill health, co-existing mental illness condition. Setting: psychiatric hospital, USA.
Interventions	1. Haloperidol: dose 5 mg/IM, then flexible dose 2.5 mg/hour/IM to 5 mg/hour/IM, maximum 100 mg/day (mean daily dose reported only for day one). N = 29. 2. Loxapine: dose 25 mg/IM, then flexible dose 12.5 mg/hour/IM to 25 mg/hour/IM, maximum 250 mg/day (mean daily dose reported only for day one). N = 25. Antiparkinsonian drugs and chloral hydrate as required.
Outcomes	Mental state: BPRS.* Global state: CGI-I, needing an extended period of medication > 24 hours. Leaving the study early. Adverse effects: sedation - ESBE. Unable to use: Global state: CGI-S (no usable data).

Tuason 1986 (Continued)

Adverse effects: amount of antiparkinsonian drugs/sleeping tablets administered not reported.

Notes *BPRS scores reported with SD. Data thought not to be SD but SE and review authors have converted to SD.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	High risk	Staff administering the medication were not blind to the intervention.
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Rater-blinded.
Incomplete outcome data (attrition bias) All outcomes	High risk	Participants who were discharged/transferred were not followed up. The number of participants reported as leaving the study is contradictory as follows; "On Day 3, 19% of the haloperidol patients were withdrawn from the study while all loxapine patients remained" "the others (6 loxapine and 11 haloperidol) were dropped on Day 2, when all were still rated severely or very severely ill." (p.127). By day 10, over 50% attrition in both groups.
Selective reporting (reporting bias)	High risk	The amount of anticholinergic and sleeping medication administered was not reported. Table 9 and 3 present data from 48 hours after the 1st injection, rather than from 24 hours, despite referring to two of the outcomes in the text; "Twenty-four hours after the first I.M. dose, 72% (18 of 25) of the loxapine patients and 59% (12 of 27) of the haloperidol patients showed moderate or marked improvement" (p.128).
Other bias	Unclear risk	None obvious.

Walther 2014

Methods	Allocation: randomised. Blinding: not stated. Duration: 96 hours.
Participants	Diagnosis: DSM-IV criteria for schizophrenia (N = 27), schizoaffective (N = 6), or schizophreniform disorder (N = 10); PANSS-PES ≥20. N = 43*. Age: range 18-55. Sex: 31 males, 12 females. History: admitted to acute care inpatient unit.

Walther 2014 (Continued)

Excluded: relevant medical or neurological disorders, pregnancy, ongoing intake of illicit drugs and alcohol.

Setting: psychiatric hospital of Bern, Switzerland.

Interventions	<p>1. Haloperidol: fixed dose 10 mg/oral for the first day**, then fixed dose 7.5 mg/twice daily/oral***. N = 14.</p> <p>2. Risperidone: fixed dose 2 mg/oral for the first day**, fixed dose 2 mg/twice daily/oral***, then fixed dose 3 mg/twice daily/oral***. N = 14.</p> <p>3. Olanzapine: fixed dose 15 mg/oral for the first day**, then fixed dose 20 mg/oral***. N = 15.</p>
Outcomes	<p>Specific behaviour: agitation - PANSS-PAS.</p> <p>Adverse effects: general - death.</p> <p>Adverse effects: movement - AIMS.</p> <p>Adverse effects: movement - BARS.</p> <p>Adverse effects: movement - SAS.</p> <p>Leaving the study early.</p> <p>Unable to use:</p> <p>Mental state: PANSS (mean scores and SD/SE/CI not provided).</p> <p>Adverse effects: amount of patients that needed antiparkinsonian drugs/benzodiazepine not reported.</p>
Notes	<p>*Eligible patients were 52 participants but 9 refused to provide post-hoc consent.</p> <p>**Additional use of diazepam up to 30 mg was permitted in day 1.</p> <p>***Additional use of diazepam up to 60 mg was permitted from day 2 to day 5.</p> <p>Authors were contacted (2nd July 2016, 9th July 2016).</p>

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"Randomization was performed by creating 3 sets of random numbers between 1 and 60 using a computer-based research randomizer" (p.125).
Allocation concealment (selection bias)	Low risk	"The randomization list was locked in the office of the principal investigator (T.J.M.), who was engaged neither in treatment nor in study assessments" (p.125).
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Not stated.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Not stated.
Incomplete outcome data (attrition bias) All outcomes	High risk	9 participants were excluded post-randomisation due to refusal to provide post-hoc consent; their treatment assignment was not provided.

Walther 2014 (Continued)

Selective reporting (reporting bias)	Unclear risk	PANSS-PAS scores at 24, 48, 72, and 96 hours not provided. PANSS scores at 96 hours not provided. Number of patients that needed additional BDZ not provided. Number of patients that needed additional biperiden not provided.
Other bias	Unclear risk	Small study.

Yin 2012

Methods	Allocation: randomised. Blinding: researcher-blinded. Duration: 72 hours.
Participants	Diagnosis: ICD-10 diagnosis of schizophrenia. N = 60 Age: range 20-54 years. Sex: 33 males, 27 females. History: acute agitation; length of illness: 1- 23 years. Excluded: "taking ziprasidone 3 months before recruitment; taking other schizophrenia medications 12 hours before recruitment; taking long-term schizophrenia medication 2 weeks before recruitment; additive to drugs 3 months before recruitment; ECG irregular (QTc > 500msec)". Setting: inpatients, China.
Interventions	1. Haloperidol: starting dose 5 mg/IM/day; maximum dose 20 mg. N = 30. 2. Ziprasidone: starting dose 10 mg/IM/day; maximum dose 40 mg. N = 30.
Outcomes	Mental state: BPRS total. Adverse effects. Adverse effects: movement - SAS.
Notes	*It is not clear if this paper presents the results from one centre of the Fang 2012 multi-centre trial. Attempted to contact author to clarify.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Described as randomised - method of randomisation is not reported.
Allocation concealment (selection bias)	Unclear risk	Not stated.

Yin 2012 (Continued)

Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Researcher blinded to assignment.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Not stated.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No evidence of incomplete outcome data.
Selective reporting (reporting bias)	Low risk	No evidence of selective reporting.
Other bias	Low risk	None obvious.

Zhou 2014

Methods	Allocation: randomised. Blinding: not stated. Duration: 4 weeks*.
Participants	Diagnosis: CCMD-3 diagnosis of schizophrenia. N = 76. Age: range 16-60 years. Sex: 39 males, 37 females. History: "excited/agitated symptoms of acute-stage schizophrenia"; "length of illness 1-32 months". Excluded: "serious physical sickness, additive to alcohol and drugs, pregnant, history of taking long-term schizophrenia medicine, suicide tendency and allergic to treatment". Setting: inpatients, China.
Interventions	1. Haloperidol: flexible dose 5 mg/IM/day to 30 mg/IM/day (mean dose 16.4 ± 6.1 mg/day). N = 38. 2. Ziprasidone: flexible dose 40 mg/oral/day to 160 mg/oral/day (mean dose 90.6 ± 20.8 mg/day) + clonazepam flexible dose 2 mg/oral/day to 6 mg/oral/day. N = 38.
Outcomes	Mental state: PANSS total score, positive sub-scale, negative sub-scale, general psychopathology sub-scale. Agitation: PANSS-EC. Global outcome: CGI-S. Leaving the study early.
Notes	*Data were extracted only for the first 7 days because at the 8th day research group stopped clonazepam and switched from haloperidol to ziprasidone.

Zhou 2014 (Continued)

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"randomised by using random number table".
Allocation concealment (selection bias)	Unclear risk	Not stated.
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Not stated.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Not stated.
Incomplete outcome data (attrition bias) All outcomes	High risk	A total of 3 dropped out in haloperidol group (serious adverse events = 2, not following protocol = 1) and 2 dropped out in ziprasidone + clonazepam (lost to contact = 1, not following protocol = 1); missing data have not been imputed using appropriate methods such as LOCF.
Selective reporting (reporting bias)	Low risk	No evidence of selective reporting.
Other bias	Low risk	None obvious.

ABS: Agitated Behaviour Scale
 ACES: Agitation-Calmness Evaluation Scale
 AIMS: Abnormal Involuntary Movement Scale
 BARS: Behavioural Activity Rating Scale
 BAS: Barnes Akathisia Scale
 BDZ: benzodiazepine
 BPRS: Brief Psychiatric Rating Scale
 BRMS:
 CABS: Corrigan Agitated Behavior Scale
 CCMD: Chinese Classification of Mental Disorders
 CGI: Clinical Global Impression
 CGI-I: Clinical Global Impression - Improvement
 CGI-S: Clinical Global Impression - Severity
 CI: confidence interval
 COSTART: Coding Symbols and Thesaurus for Adverse Reaction Terms
 DAI-10: Drug Attitude Inventory
 DSM: Diagnostic and Statistical Manual
 ECG: electrocardiogram
 EPS: Extrapyramidal Side Effects
 ESBE:
 GAS: Global Adjustment Scale
 ICD: International Classification of Diseases
 IM: intramuscular
 LOCF: last observation carried forward
 LS:
 MBPRS: Modified Brief Psychiatric Rating Scale
 NOSIE: Nurses Observation Scale for Inpatient Evaluation
 OAS: Overt Aggression Scale
 OASS: Overt Agitation Severity Scale

PANSS: Positive and Negative Syndrome Scale

PANSS-EC: Positive and Negative Syndrome Scale Excited Component

PANSS-PAS: Positive and Negative Syndrome Scale Psychotic Agitation Component

PANSS-PES:

p.o.: orally

RSS: Ramsay Sedation Scale

SAS: Simpson-Angus Scale

SD: standard deviation

SE: standard error

SGPT: serum glutamic pyruvic transaminase

TCS:

TESS: Treatment Emergent Symptom Scale

TSRS: Target Rating Symptom Scale

VAS: Visual analogue scale

YMRS: Young Mania Rating Scale

Characteristics of excluded studies *[ordered by study ID]*

Study	Reason for exclusion
Addington 1996	Allocation: randomised. Participants: people with acute schizophrenia, but not specifically overtly aggressive.
Alexander 2004	Allocation: randomised. Participants: acute agitation and dangerous behaviour. Intervention: haloperidol + promethazine vs olanzapine, no haloperidol alone group.
Allen 2007	Allocation: not randomised, review.
Asadollahi 2015	Allocation: randomised. Participants: psychotic disorders (N = 52), mood disorders (N = 23), cognitive impairment (N = 23), adjustment disorders (N = 17), others (infection, substance intoxication or withdrawal) (N = 35), and unknown aetiology (N = 10). Therefore, eligible patients are < 50% of the sample. An attempt to contact the authors for partial outcomes was made (3rd July 2016).
Beasley 1998	Allocation: randomised. Participants: people with first episode psychosis, but not specifically aggressive.
Bieniek 1998	Allocation: randomised. Participants: newly admitted patients to a psychiatric emergency service who were uncooperative, threatening, and unresponsive to verbal interventions. Intervention: haloperidol + lorazepam vs lorazepam, no haloperidol alone group.
Brook 1998	Allocation: randomised. Participants: people with acute psychosis, but not specifically aggressive.
Campbell 1982	Allocation: randomised. Participants: children with aggressive behaviour, but the number of those with suspected psychosis < 50% of the sample.
Chen 2004	Allocation: quasi-randomised.

Study	Reason for exclusion
Chen 2008	Allocation: quasi-randomised.
Chouinard 1991	Allocation: randomised. Participants: people with acute psychosis, but not specifically aggressive.
Citrome 2001	Allocation: randomised. Participants: people with schizophrenia and persistent aggressive behaviour, not acute exacerbation of symptoms.
Colonna 1998	Allocation: randomised. Participants: people with acute schizophrenia, but not specifically aggressive.
Conde 2011	Allocation: randomised. Participants: admitted patients to an emergency setting experiencing agitation associated with active psychosis; inpatients experiencing exacerbation or agitation with active psychosis. Intervention: haloperidol + clonazepam vs risperidone + clonazepam, no haloperidol alone group.
Crandall 2005	Allocation: randomised. Participants: people with chronic schizophrenia, acute exacerbation of symptoms, not specifically aggressive.
Currier 2000	Allocation: not randomised.
Daniel 2000	Allocation: not randomised, review.
Daniel 2003	Allocation: randomised. Participants: people with acute schizophrenia, not acutely agitated or aggressive.
Daniel 2004	Allocation: randomised. Participants: people with psychosis, not acutely agitated or aggressive.
Fan 2012	Quasi-randomised controlled trial: "the participants were assigned by the order of admission".
Faretra 1970	Allocation: randomised. Participants: children with suspected schizophrenia, not specifically aggressive.
Freeman 2009	Allocation: not randomised, retrospective case note study.
Goldstein 1966	Allocation: randomised. Participants: people with acute psychosis, not aggressive.
Harvey 2004	Allocation: randomised. Participants: people with mild symptoms of schizophrenia, not aggressive.
Huang 2004	Allocation: quasi-randomised.
Jiang 2009	Quasi-randomised controlled trial: "the participants were assigned by the order of admission".
Jones 2001	Allocation: not randomised, review.

Study	Reason for exclusion
Kane 2000	Allocation: randomised. Participants: people with acute schizophrenia, not specifically aggressive.
Kinon 2001a	Allocation: randomised. Participants: people with schizophrenia, but not specifically overtly aggressive.
Kinon 2003	Allocation: not randomised.
Krakowski 2006	Allocation: randomised. Participants: people with schizophrenia and persistent aggressive behaviour, not an emergency situation.
Krakowski 2008	Allocation: randomised. Participants: people with schizophrenia and persistent aggressive behaviour, not an emergency situation.
Lewis 2006	Allocation: randomised. Participants: people with schizophrenia and persistent aggressive behaviour, not an emergency situation.
Li 2005	Allocation: randomised. Participants: people with acute excitement in schizophrenia. Intervention: haloperidol vs tiapride. Outcome: there was no specific outcome for acute agitation (T1 at 7 days).
Li 2007	Allocation: quasi-randomised.
Liang 2003	Quasi-randomised controlled trials: "the participants were randomly assigned by the order of admission".
Liu 2012b	Allocation: randomised. Participants: people with schizophrenia and acute agitation. Intervention: haloperidol vs modified electroconvulsive therapy (MECT), no comparison drug.
Mandel 2008	Allocation: randomised. Participants: people with schizophrenia, not specifically aggressive.
Mantua 2006	Allocation: not randomised, naturalistic observational study.
McEvoy 1994	Allocation: randomised. Participants: people with schizophrenia, not specifically aggressive.
Mendelowitz 2004	Allocation: not randomised, review.
NCT00189995 2005	Allocation: was to be randomised, but trial not conducted because ethics approval not granted.
NCT00631722 2008	Allocation: randomised.

Study	Reason for exclusion
	Participants: people with acute schizophrenia, not specifically aggressive.
NCT00797277 2008	Allocation: randomised. Participants: people with schizophrenia and agitation. Intervention: haloperidol plus lorazepam vs olanzapine - no haloperidol alone group.
Pan 2005	Allocation: quasi-randomised.
Pathiraja 1995	Allocation: randomised. Participants: people with schizophrenia, not specifically aggressive.
Pedros 2004	Allocation: not randomised - cohort study.
Pei 2009	Quasi-randomised controlled trial: "the participants were assigned by the order of admission".
Puech 1998	Allocation: randomised. Participants: people with acute schizophrenia, not specifically aggressive.
Romeo 2009	Allocation: randomised. Participants: people with intellectual disability and challenging behaviour, not psychosis-induced aggression.
Simpson 2010	Allocation: randomised. Participants: people with agitation in the emergency setting. Intervention: haloperidol plus benztropine mesylate vs lorazepam vs quetiapine - no haloperidol alone group.
Singh 1981	Allocation: not reported, described as double-blind. Participants: people with schizophrenia, not acute exacerbation of symptoms.
Smith 1974	Allocation: randomised. Participants: people with senile psychosis, not specifically aggressive.
Srinath 2010	Allocation: randomised. Participants: "patients who presented with acute psychotic agitation". Intervention: haloperidol + promethazine vs lorazepam, no haloperidol alone group.
Teja 1975	Allocation: randomised. Participants: people with chronic schizophrenia, not specifically aggressive.
Thomas 1992	Allocation: randomised. Participants: violent and agitated people, mostly intoxicated, not necessarily psychosis-induced aggression.
TREC CG	Allocation: randomised. Participants: acute agitation and dangerous behaviour.

Study	Reason for exclusion
	Intervention: haloperidol + promethazine vs midazolam (no haloperidol alone group).
Ungvari 1982	Allocation: randomised. Participants: people with acute schizophrenia, not specifically aggressive.
Vaisanen 1981	Allocation: randomised. Participants: people with intellectual disability and agitation, not psychosis-induced aggression.
Veser 2006	Allocation: randomised. Participants: people with psychosis-induced agitation, not specifically aggressive (exclusion criteria included an inability to give informed consent).
Vives 2015	Allocation: randomised. Participants: agitated schizophrenic patients in the emergency setting. Intervention: haloperidol intranasal vs haloperidol intramuscular, no other drugs except haloperidol.
Wan 2005	Allocation: randomised. Participants: people with schizophrenia and agitation/aggression. Intervention: haloperidol vs chlorpromazine vs risperidone + clonazepam. Outcome: excitement and agitation scale (not validated), also the data for the haloperidol and chlorpromazine groups are combined.
Wang 2004	Allocation: quasi-randomised.
Wang 2005	Allocation: quasi-randomised.
Wyant 1990	Allocation: randomised. Participants: people with schizophrenia and acute agitation. Intervention: haloperidol vs sodium amytal vs midazolam. Outcome: modified CGI - no usable data, presented in graphs, no binary data.
Zapletalek 1986	Allocation: not reported, described as double-blind. Participants: people with schizophrenia, not specifically aggressive.

CGI: Clinical Global Impression
vs: versus

Characteristics of studies awaiting assessment *[ordered by study ID]*

Daniel 2006

Methods	Allocation: randomised. Blindness: not stated. Duration: 24-hour IM phase followed by 4-day oral phase.
Participants	Diagnosis: schizophrenia with agitation. N = 360.

Haloperidol for psychosis-induced aggression or agitation (rapid tranquillisation) (Review)

Copyright © 2017 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.

Daniel 2006 (Continued)

	Age: range 18-60 years.
	Sex: not stated.
	History: not stated.
	Exclusion: not stated.
	Setting: not stated.
Interventions	<p>1. Haloperidol: dose 6.5 mg/IM, maximum 3 injections during 24 hours, followed by haloperidol 7 mg/p.o./day to 10 mg/p.o./day for 4 days.</p> <p>2. Aripiprazole: dose 10 mg/IM, maximum 3 injections during 24 hours, followed by haloperidol 10 mg/p.o./day to 15 mg/p.o./day for 4 days.</p>
Outcomes	<p>Global state: CGI-I, CGI-S.</p> <p>Agitation: PANSS-EC, ACES, CABS.</p>
Notes	Conference proceeding, full characteristics and outcome data not reported. Additionally, unclear whether this is the same study as Bristol-Myers 2005b - attempted to contact Bristol Myers.

Davis 2008

Methods	<p>Allocation: randomised.</p> <p>Blindness: not stated.</p> <p>Duration: 24 hours.</p>
Participants	<p>Diagnosis: schizophrenia.</p> <p>N = 62.</p> <p>Age: not stated.</p> <p>Sex: not stated.</p> <p>History: "agitated schizophrenic patient admitted to hospital."</p> <p>Exclusion: not stated.</p> <p>Setting: not stated.</p>
Interventions	<p>1. Haloperidol: dose 5 mg/IM/day to 15 mg/IM/day. N = 21.</p> <p>2. Haloperidol: dose 5 mg/IM/day to 15 mg/IM/day + clonazepam: dose 2 mg/IM/day to 6 mg/IM/day. N = 21.</p> <p>3. Clonazepam: dose 2 mg/IM/day to 6 mg/IM/day. N = 20.</p>
Outcomes	<p>Mental state: BPRS Total, BPRS Psychosis sub-scale.</p> <p>Global state: CGI.</p> <p>Agitation: BPRS agitation sub-scale.</p> <p>Adverse effects.</p>

Davis 2008 (Continued)

Notes	Conference proceeding, full characteristics and outcome data not reported. Attempted to contact author.
-------	---

Dubin 1985

Methods	Allocation: randomised. Blindness: single. Duration: not stated.
Participants	Diagnosis: schizophrenia (N = 77), mania (N = 37), psychosis aetiology unknown (N = 25), amphetamine intoxication (N = 11), alcohol intoxication (N = 5), brief reactive psychosis (N = 3), brain organic syndrome (N = 1). N = 159. Age: range 18-65 years. Sex: 77 males and 82 females. History: newly admitted - previous history not stated. Exclusion: pregnancy, central nervous system depression, history of urinary retention or glaucoma, known sensitivity to phenothiazines, conditions such as epilepsy, severe hypotension or hypertension, head injuries, and haematologic, cardiovascular, renal or hepatic impairments, and ingestion of antipsychotic, sedative-hypnotic, or antianxiety drugs within 24 hours of admission. Setting: psychiatric emergency hospital, USA.
Interventions	1. Haloperidol: dose 5 mg/oral concentrate (mean number of doses 3.8). N = 15. 2. Haloperidol: dose 5 mg/IM (mean number of IMs not reported 3.2). N = 14. 3. Haloperidol: dose 10 mg/oral concentrate (mean number of doses 3.5). N = 14. 4. Haloperidol: dose 10 mg/IM (mean number of IMs 2.2). N = 17. 5. Thiothixene: dose 10 mg/IM (mean number of IMs 3.1). N = 24. 6. Thiothixene: dose 10 mg/oral concentrate (mean number of doses 4.7). N = 14. 7. Thiothixene: dose 20 mg/oral concentrate (mean number of doses 3.5). N = 14. 8. Thioridazine: dose 25 mg/oral suspension (mean number of doses 3.8). N = 16. 9. Thioridazine: dose 50 mg/oral suspension (mean number of doses 3.8). N = 13. 10. Thioridazine: dose 100 mg/oral suspension (mean number of doses 3.2). N = 18.
Outcomes	Global state: additional injections. Mental state: BPRS (mean/SD/SE/CI not reported). Adverse effects: % of whole sample experiencing side effects reported.*
Notes	*Attempted to contact author to see if there are any usable data available.

Esmailian 2015

Methods	Allocation: randomised. Blindness: double. Duration: not stated.
Participants	Diagnosis: patients who referred to the emergency department because of medical diseases, drug poisoning, or trauma, and need the sedative agent to sedation. N = 48. Age: 18-65 years. Sex: 36 males and 12 females. History: internal (N = 36)*, drug poisoning (N = 2), trauma (N = 10). Exclusion: sensitivity to haloperidol or midazolam, contraindications to these drugs, sympathomimetic agents, agitation because of reversible factors (such as hypotension, hypoxia, and hypoglycaemia), tachycardia or bradycardia, respiratory distress, pregnancy, symptoms of withdrawal syndrome, and receiving sedative agents within the past 12 hours. Setting: emergency department, Iran.
Interventions	1. Haloperidol: fixed dose 5 mg/IM. 2. Midazolam: flexible doses 2.5 mg/IM to 5 mg/IM according to weight.
Outcomes	General: time to sedation, time to full consciousness, need for additional doses, need for re-sedation within 60 minutes. Adverse effects.
Notes	*not clear which diagnosis includes; an attempt to contact the authors was made (3rd July 2016).

Herrera 2005

Methods	Allocation: randomised. Blindness: double. Duration: not stated.
Participants	Diagnosis: psychosis with agitation and/or violence. N = 20. Age: not stated. Sex: not stated. History: not stated. Exclusion: not stated. Setting: psychiatric emergency department.
Interventions	1. Haloperidol: dose 10 mg/IM. 2. Risperidone: dose 2 mg/liquid.

Herrera 2005 (Continued)

Outcomes	Agitation: PANSS-EC.
Notes	Conference proceeding, full characteristics and outcome data not reported. Attempted to contact author.

Hsu 2010

Methods	Allocation: randomised. Blindness: single. Duration: 24 hours.
Participants	Diagnosis: DSM-IV diagnosis of schizophrenia (N = 20), bipolar disorder (N = 18), schizoaffective disorder (N = 1), delusional disorder or others (N = 3). N = 42. Age: range 18-65 years. Sex: 20 males and 22 females. History: within 24 hours of admission - previous psychiatric history not stated. Exclusion: "pregnant or lactating women; patients with serious medical illnesses; patients with closed-angle glaucoma; patients with an allergic reaction to olanzapine, risperidone, or haloperidol; or patients who had received a long-acting antipsychotic agent injection within 30 days were excluded". Setting: acute medical centre, Taiwan.
Interventions	1. Haloperidol: dose 7.5 mg/IM. N = 11. 2. Risperidone: dose 3 mg/liquid. N = 10. 3. Olanzapine: dose 10 mg/IM. N = 11. 4. Olanzapine: dose 10 mg/velotab. N = 10.
Outcomes	Agitation: PANSS-EC.*
Notes	*Attempted to contact author to see if there is any usable data available.

Istikoglou 2010

Methods	Allocation: randomised. Blindness: rater-blinded. Duration: not stated.
Participants	Diagnosis: psychosis with agitation. N = 120. Age: not stated. Sex: males and females.

Istikoglou 2010 (Continued)

	History: not stated.
	Exclusion: not stated.
	Setting: psychiatric emergency room, Greece.
Interventions	1. Haloperidol IM (dose not reported). 2. Aripiprazole IM (dose not reported).
Outcomes	Mental state: PANSS, YMRS. Global state: CGI-I. Agitation: AIMS. Adverse effects: SAS. Quality of life: QoL
Notes	Conference proceeding, full characteristics and outcome data not reported. Unable to establish contact details at this time.

Kong 2009

Methods	Allocation: randomised. Blindness: open-label. Duration: 2 hours.
Participants	Diagnosis: acutely agitated patients. N = 37. Age: 18 years or older. Sex: not stated. History: "DSM-IV diagnosis of schizophrenia, schizophreniform disorder, brief psychotic disorder, or schizoaffective disorder, bipolar disorder" (p.S525). Exclusion: not stated. Setting: psychiatry ward, Inje University, China.
Interventions	1. Haloperidol: fixed dose 5 mg/IM. N = 10. 2. Olanzapine: fixed dose 10 mg/IM. N = 12. 3. Haloperidol: fixed dose 5 mg/IM + lorazepam fixed dose 4 mg/IM. N = 15.
Outcomes	Global State: CGI-S, CGI-I. Agitation: PANSS-EC. Adverse effects. Seclusions.
Notes	Abstract, full characteristics and outcome data not reported. Additionally, it is probably the same study as Shim 2009 (discordant doses and reported adverse effects).

Haloperidol for psychosis-induced aggression or agitation (rapid tranquillisation) (Review)

Kong 2009 (Continued)

An attempt to contact the authors was made (3rd July 2016).

Lasic 2006

Methods	Allocation: randomised. Blindness: not stated. Duration: up to 3 months.
Participants	Diagnosis: ICD X diagnosis of acute exacerbation of schizophrenia or schizoaffective disorder, mania with psychotic features, acute paranoid reaction, or delusional disorders. N = 60. Age: > 18. Sex: males and females. History: not stated. Exclusion: "delirium, epilepsy, or mental retardation; intoxication or symptoms of withdrawal from alcohol or other psychoactive substances; clinical laboratory values indicating serious medical illness; treatment with any antipsychotic or benzodiazepine within 6 hours of screening; a history of neuroleptic malignant syndrome or known hypersensitivity to any of the trial medications; treatment with a depot antipsychotic within 1 treatment cycle of screening and use of disallowed medications". Setting: acute psychiatric inpatient ward, Croatia.
Interventions	1. Haloperidol IM (dose not reported). 2. Risperidol liquid (dose not reported).
Outcomes	Mental state: PANSS. Global state: CGI-I. Agitation: BARS, PANSS agitation cluster. Adverse effects
Notes	Conference proceeding, full characteristics and outcome data not reported. Attempted to contact author.

Li 2007a

Methods	Allocation: not clearly stated; "each unit of patients were randomly assigned to one of three treatment groups by random number and hospital serial number". Blindness: double-blind. Duration: 24 hours.
Participants	Diagnosis: CCMD-3 diagnosis of schizophrenia. N = 45.

Li 2007a (Continued)

	<p>Age: range 18-45 years.</p> <p>Sex: males (20) and females (25).</p> <p>History: acute agitation.</p> <p>Exclusion: serious physical illness; additive to alcohol, drugs; taking schizophrenia medication one month before recruitment.</p> <p>Setting: psychiatric inpatient ward, China.</p>
Interventions	<p>1. Haloperidol: dose 5 mg/IM/day to 15 mg/IM/day. N = 15.</p> <p>2. Haloperidol: dose 5 mg/IM/day to 15 mg/IM/day + clonazepam dose 2 mg/IM/day to 6 mg/IM/day. N = 15.</p> <p>3. Clonazepam: dose 2 mg/IM/day to 6 mg/IM/day. N = 15.</p>
Outcomes	<p>Mental state: BPRS.</p> <p>Adverse effects.</p> <p>Unable to use:</p> <p>CGI: data not available.</p>
Notes	Allocation methodology not clearly stated.

Lu 2006

Methods	<p>Allocation: randomised.</p> <p>Blindness: rater-blinded.</p> <p>Duration: 72 hours.</p>
Participants	<p>Diagnosis: CCMD-3 diagnosis of schizophrenia.</p> <p>N = 60.</p> <p>Age: mean 33.5 (SD 12.3) years.</p> <p>Sex: male and female.</p> <p>History: acute agitation.</p> <p>Excluded: anyone who used anti-psychotic or anti-convulsant within one week prior to the trial or anyone who used long lasting drug within one month prior to the trial.</p> <p>Setting: inpatients, China*.</p>
Interventions	<p>1. Haloperidol: dose 5 mg/IM, to 10 mg/IM, maximum dose 30 mg during 24 hours. N = 30.</p> <p>2. Ziprasidone: dose 10 mg/IM, to 20 mg/IM, maximum dose 40 mg during 24 hours. N = 30.</p>
Outcomes	<p>Global state: CGI-IS.</p> <p>Adverse effects: TESS.</p> <p>Agitation: PANSS-EC.</p> <p>Unable to use:</p>

Lu 2006 (Continued)

Adverse effects: the amount of anticholinergic drugs used was not reported.

Notes	*It is not clear if this paper presents the results from one centre of the Li 2006 multi-centre trial. Attempted to contact author to clarify.
-------	--

NCT00859872 2009

Methods	Allocation: randomised. Blindness: single. Duration: up to 47 days.
Participants	Diagnosis: DSM-IV diagnosis of acute exacerbation of schizophrenia or schizoaffective disorder with agitation. N = not stated. Age: range > 18 < 45 years. Sex: males and females. History: not stated. Exclusion: pregnant or lactating women, serious medical illness, known sensitivity to study medication, treatment with a depot antipsychotic with 1 cycle of screening, use of disallowed medication, psychosis caused by "delirium, epilepsy, mental retardation and affective disorder; intoxication or symptoms of withdrawal from alcohol of other psychoactive substances." Setting: psychiatric inpatient ward, China.
Interventions	1. Haloperidol: dose 5 mg/IM/day to 20 mg/IM/day. 2. Risperidone: dose 2 mg/oral/day to 6 mg/oral/day + clonazepam: dose 4 mg/oral/day to 8 mg/oral/day.
Outcomes	Mental state: PANSS. Agitation: PANSS-EC.
Notes	*Protocol, full characteristics and outcome data not reported. Unable to establish contact details at this time.

NCT00866645 2009

Methods	Allocation: randomised. Blindness: single. Duration: 72 hours.
Participants	Diagnosis: DSM-IV diagnosis of schizophrenia or schizophreniform psychosis. N = not stated. Age: range > 18 < 65 years. Sex: males and females.

NCT00866645 2009 (Continued)

History: not stated.

Exclusion: investigator and his/her relatives, participation in another drug trial within 3 months prior to enrolment into this study, pregnant or lactating women, serious medical illness, a significantly abnormal value in ECG or lab results, family history of sudden death, meet the DSM-IV criteria for substance abuse within 1 year prior to enrolment, regular use of antipsychotics (clozapine within 90 days), antidepressants, mood stabilisers, anti-epileptics or prolonged-action preparations within 2 weeks prior to enrolment, use of ECT within 30 days prior to enrolment, systematic use of sulpiride, levosulpiride, or haloperidol therapy within 30 days prior enrolment, history of neuroleptic malignant syndrome, severe EPS or significant tardive dyskinesia, severe suicide attempt, known hypersensitivity to sulpiride, levosulpiride or haloperidol, history of hypersensitivity to more than 2 drugs, use of psychotropics (except for permitted drugs) within 12 hours prior to enrolment, known lack of efficacy to levosulpiride or haloperidol by formal treatment before, organic mental disorders including learning disability, history of psychosurgery treatment, people who could not comply with the study protocol.

Setting: psychiatric inpatient ward, China.

Interventions	<p>1. Haloperidol: dose 5 mg/IM, ≥ 4 hours apart, maximum 10 mg during 24 hours.</p> <p>2. Levosulpiride: dose 50 mg/IM, ≥ 4 hours apart, maximum 100 mg during 24 hours.</p>
Outcomes	<p>Mental state: BPRS, PANSS Total.</p> <p>Global State: CGI, CGI-I, CGI-S.</p> <p>Agitation: ACES, PANSS-EC.</p> <p>Adverse effects: BAS, ECG, laboratory tests, physical examination, RSESE.</p>
Notes	Protocol, full characteristics and outcome data not reported. Attempted to contact author to enquire whether there is any available data.

Shim 2009

Methods	<p>Allocation: randomised.</p> <p>Blindness: open-label.</p> <p>Duration: 2 hours.</p>
Participants	<p>Diagnosis: acutely agitated patients.</p> <p>N = 37.</p> <p>Age: not stated.</p> <p>Sex: not stated.</p> <p>History: "hospitalized patients with acute psychotic symptom in association with schizophrenia, bipolar I disorder, schizoaffective disorder, schizophreniform disorder, brief psychotic disorder, delusional disorder" (p.S512).</p> <p>Exclusion: not stated.</p> <p>Setting: psychiatry ward, Inje University, China.</p>
Interventions	<p>1. Haloperidol: fixed dose 4 mg/IM. N = 10.</p> <p>2. Olanzapine: fixed dose 10mg/IM. N = 12.</p>

Shim 2009 (Continued)

	3. Haloperidol: fixed dose 4 mg/IM + lorazepam fixed dose 5 mg/IM. N = 15.
Outcomes	Global State: CGI-S. Agitation: PANSS-EC, BARS. Adverse effects.
Notes	Abstract, full characteristics and outcome data not reported. Additionally, it is probably the same study as Kong 2009 (discordant doses and reported adverse effects). An attempt to contact the authors was made (3rd July 2016).

Si 2009

Methods	Allocation: randomised. Blindness: open-label. Duration: 28 days.
Participants	Diagnosis: acute exacerbation of schizophrenia with agitation and hostility. N = 80. Age: not stated. Sex: not stated. History: not stated. Exclusion: not stated. Setting: psychiatric inpatient ward, China.
Interventions	1. Haloperidol: dose 10 mg/oral/day to 12 mg/oral/day. N = 40. 2. Quetiapine: dose 600 mg/oral/day to 750 mg/oral/day. N = 40.
Outcomes	Mental state: BPRS, PANSS Total. Global State: CGI, CGI-I, CGI-S. Agitation: ACES, PANSS-EC. Adverse effects: BAS, ECG, laboratory tests, physical examination, RSESE.
Notes	Conference proceeding, full characteristics and outcome data not reported. Attempted to contact author.

Slotnick 1971

Methods	Allocation: randomised. Blindness: double-blind. Duration: 48 hours.
---------	--

Slotnick 1971 *(Continued)*

Participants	<p>Diagnosis: "acutely agitated psychiatric patients".</p> <p>N = 150 (three trials combined).</p> <p>Age: not stated.</p> <p>Sex: not stated.</p> <p>History: not stated.</p> <p>Exclusion: not stated.</p> <p>Setting: not stated.</p>
Interventions	<p>1. Haloperidol IM (dose not reported). N = 75 (three trials combined).</p> <p>2. Chlorpromazine IM (dose not reported). N = 75 (three trials combined).</p>
Outcomes	Mental state: BPRS.
Notes	Conference proceeding, full characteristics and outcome data for the three trials not reported. Unable to establish contact details at this time.

Smythies 1982

Methods	<p>Allocation: not reported.</p> <p>Blindness: double.</p> <p>Duration: 10 days.</p>
Participants	<p>Diagnosis: DSM-III diagnosis of schizophrenia with acute agitation.</p> <p>N = 16.</p> <p>Age: not stated.</p> <p>Sex: not stated.</p> <p>History: not stated.</p> <p>Exclusion: at risk of pregnancy, significant medical disorders, history of EPS or hypotensive crisis upon administration of other injectable antipsychotic agents, significantly abnormal laboratory test results, recent alcoholism or drug dependency.</p> <p>Setting: inpatient, USA.</p>
Interventions	<p>1. Haloperidol: dose 2.5 mg/IM to 80 mg/IM, for 2 to 10 days.</p> <p>2. Molindone: dose 12.5 mg/IM to 150 mg/IM, for 4-7 days.</p>
Outcomes	<p>Mental state: BPRS.</p> <p>Global state: CGI.</p> <p>Agitation: TSRS*.</p>
Notes	*Attempted to contact author to see if there is any available usable data available.

Vasquez-Gomez 2001

Methods	Allocation: randomised. Blindness: simple. Duration: not stated.
Participants	Diagnosis: DSM-IV diagnosis of acute psychosis - "schizophrenic or affective". N = 30. Age: range 18-83 years. Sex: not stated. History: not stated. Exclusion: not stated. Setting: emergency room, Peru.
Interventions	1. Haloperidol: dose 5 mg/IM. N = 15. 2. Midazolam: dose 15 mg/IM. N = 15.
Outcomes	Aggression: OAS.
Notes	Conference proceeding, full characteristics and outcome data for the three trials not reported. Unable to establish contact details at this time.

Wang 2005a

Methods	Allocation: randomised. Blinding: single-blind. Duration: 72 hours.
Participants	Diagnosis: CCMD-3 diagnosis of schizophrenia. N = 60. Age: mean 27.40 ± 9.64 years (ziprasidone group), mean 27.40 ± 9.64 years (haloperidol group). Sex: 39 males and 21 females. History: total score of PANSS ≥ 60 ; total score of excitement component of PANSS ≥ 14 and at least one section ≥ 4 . Exclusion: physical illness; additive to alcohol, drugs; pregnant or plan to be pregnant. Setting: inpatients*, China.
Interventions	1. Haloperidol: dose 5 mg/IM to 10 mg/IM, maximum 30 mg/day. N = 30. 2. Ziprasidone: dose 10 mg/IM to 20 mg/IM, maximum 40 mg/day. N = 30.
Outcomes	Mental state: PANSS-EC. Adverse effects: TESS.

Wang 2005a (Continued)

Adverse effects: movement - SAS.

Notes	*It is not clear if this paper presents the results from one centre of the Li 2006 multi-centre trial. Attempted to contact author to clarify.
-------	--

Wang 2007

Methods	Allocation: randomised. Blindness: single. Duration: 72 hours.
Participants	Diagnosis: CCND-3 diagnosis of schizophrenia with agitation/aggression. N = 32. Age: mean 25.8 (SD 10.4) years (haloperidol group), 28.6 (SD 9.4) years (ziprasidone group). Sex: 16 males and 16 females. History: first episode (N = 27), relapse (N = 5). Excluded: direct blood relatives of staff in current trial, participation in other drug trials 30 days prior to enrolment, pregnant or lactating women, severe disease/unstable medical condition, clinically significant abnormal EEG or laboratory results, QTC interval ≥ 450 , family history of sudden death, drug abuse in past year, regular user of antipsychotic, antidepressant, depot medication, ziprasidone or haloperidol in past 30 days, people with drug-induced malignant syndrome, suicidal ideation, or known allergy to ziprasidone or haloperidol. Setting: inpatient, China*.
Interventions	1. Haloperidol: flexible dose 5 mg/IM, to 10 mg/IM, repeated every 4-6 hours if necessary, maximum dose 30 mg during 24 hours. N = 16. 2. Ziprasidone: flexible dose 10 mg/IM, to 20 mg/IM, repeated every 4-6 hours if necessary, maximum dose 40 mg during 24 hours. N = 16.
Outcomes	Mental state: PANSS. Global State: CGI. Agitation: PANNS-EC, ACES. Adverse effects: BAS.
Notes	*It is not clear if this paper presents the results from one centre of the Li 2006 multi-centre trial. Attempted to contact author to clarify.

Yildiz 2003

Methods	Allocation: not stated. Blindness: not stated. Duration: 2 hours.
Participants	Diagnosis: "acutely agitated psychiatric patients".

Yildiz 2003 (Continued)

	<p>N = not stated.</p> <p>Age: not stated.</p> <p>Sex: not stated.</p> <p>History: not stated.</p> <p>Excluded: not stated.</p> <p>Setting: acute psychiatry service.</p>
Interventions	<p>1. Haloperidol: flexible dose 2 mg/IM to 5 mg/IM. N = not stated.</p> <p>2. Risperidone: flexible dose 1 mg/oral to 2 mg/oral. N = not stated.</p> <p>Concomitant use of lorazepam was allowed.</p>
Outcomes	<p>Unable to use:</p> <p>Agitation: "four different agitation rating scales", not reported.</p>
Notes	<p>Abstract of poser, full characteristics and outcome data not reported. Unable to establish contact details at this time.</p>

Zhang 2012

Methods	<p>Allocation: randomised.</p> <p>Blinding: not stated.</p> <p>Duration: 7 days.</p>
Participants	<p>Diagnosis: DSM-IV diagnosis of schizophrenia.</p> <p>N = 60.</p> <p>Age: range 18-65 years.</p> <p>Sex: 36 males, 24 females.</p> <p>History: acute agitation; length of illness 124.7 ± 115.3 months (haloperidol group), 157.5 ± 113.0 months (risperidone+clonazepam group).</p> <p>Excluded: "allergy to compound utilised in the treatment, serious physical sickness, addiction to alcohol or drugs, history of taking long-term schizophrenia medicine, suicide tendency, pregnancy".</p> <p>Setting: inpatients, China.</p>
Interventions	<p>1. Haloperidol: dose 10 mg/IM/day to 20 mg/IM/day. N = 30.</p> <p>2. Risperidone: dose 2 mg/oral/day to 6 mg/oral/day + clonazepam dose 2 mg/oral/day to 8 mg/oral/day. N = 30.</p>
Outcomes	<p>Mental state: PANSS total and positive sub-scale.</p> <p>Agitation: PANSS-EC.</p> <p>Adverse effects: TESS.</p>
Notes	

叶萌, 2009

Methods	Allocation: randomised. Blindness: single. Duration: 72 hours.
Participants	Diagnosis: DSM-IV diagnosis of schizophrenia. N = 60. Age: mean 28.37 ± 1 0.20 years (haloperidol group), mean 27.40 ± 9.63 years (ziprasidone group). Sex: 39 males, 21 females. History: acute agitation. Excluded: "physical illness; addiction to alcohol or drugs; pregnancy or plan to be pregnant". Setting: inpatients, China*.
Interventions	1. Haloperidol: dose 5 mg/IM to 10 mg/IM, maximum 30 mg/day. N = 30. 2. Ziprasidone: dose 10 mg/IM to 20 mg/IM, maximum 40 mg/day. N = 30.
Outcomes	Mental state: PANSS total. Agitation: PANSS-EC. Adverse events.
Notes	*It is not clear if this paper presents the results from one centre of the Li 2006 multi-centre trial. Authors contact unavailable.

周德祥, 2012

Methods	Allocation: randomised. Blindness: not stated. Duration: 7 days.
Participants	Diagnosis: CCMD-3 diagnosis of schizophrenia. N = 65. Age: range 18-60 years. Sex: 36 males, 29 females. History: acute psychotic agitation; length of illness of 1.1 to 22.3 months. Excluded: pregnancy, serious physical illness, addiction to alcohol or drugs. Setting: inpatients, China*.
Interventions	1. Haloperidol: dose 10 mg/IM/day to 20 mg/IM/day. N = 32.

周德祥, 2012 (Continued)

2. Risperidone: dose 4 mg/oral/day to 6 mg/oral/day + clonazepam dose 2 mg/oral/day to 4 mg/oral/day. N = 33.

Outcomes	<p>Unable to use:</p> <p>Mental state: PANSS total (data not reported).</p> <p>Agitation: PANSS-EC (data not reported).</p> <p>Adverse events (data not reported).</p>
Notes	<p>*It is not clear if this paper presents the results from one centre of the Fang 2012 multi-centre trial. Authors contact unavailable.</p>

ACES: Agitation-Calmness Evaluation Scale
 AIMS: Abnormal Involuntary Movement Scale
 BARS: Behavioural Activity Rating Scale
 BAS: Barnes Akathisia Scale
 BPRS: Brief Psychiatric Rating Scale
 CABS: Corrigan Agitated Behavior Scale
 CCMD: Chinese Classification of Mental Disorders
 CGI: Clinical Global Impression
 CGI-I: Clinical Global Impression – Improvement
 CGI-S: Clinical Global Impression – Severity
 CI: confidence interval
 DSM: Diagnostic and Statistical Manual
 ECG: electrocardiogram
 ECT: electroconvulsive therapy
 EPS: Extrapyramidal Side Effects
 ICD: International Classification of Diseases
 IM: intramuscular
 OAS: Overt Aggression Scale
 PANSS: Positive and Negative Syndrome Scale
 PANSS-EC: Positive and Negative Syndrome Scale Excited Component
 p.o.: by mouth
 RSESE: Rating Scale for Extrapyramidal Side Effects.
 SAS: Simpson-Angus Scale
 SD: standard deviation
 SE: standard error
 TESS: Treatment Emergent Symptom Scale
 TSRS: TSRS - Target Rating Symptom Scale
 YMRS - Young Mania Rating Scale

Characteristics of ongoing studies [ordered by study ID]

NCT00838032 2009

Trial name or title	Pilot study to evaluate the efficacy and safety of quetiapine fumarate instant-release (Seroquel IR) in controlling agitation and aggressive symptoms in the acute treatment of patients with schizophrenia
Methods	<p>Allocation: randomised.</p> <p>Blindness: single.</p> <p>Duration: 14 days</p>
Participants	<p>Diagnosis: CCMD-3 diagnosis of schizophrenia with agitation.</p> <p>N = not stated.</p> <p>Age: > 18 < 65.</p>

Haloperidol for psychosis-induced aggression or agitation (rapid tranquillisation) (Review)

Copyright © 2017 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.

NCT00838032 2009 (Continued)

Sex: males and females.

History: not stated.

Exclusion: pregnancy or lactation, risk of suicide, self-harm or harm to others, known intolerance or lack of response to quetiapine fumarate or haloperidol, use of ketonazole, itraconazole, fluconazole, erythromycin, clarithromycin, troleandomycin, indinavir, nelfinavir, ritonavir, fluvoxamine, saquinavir, phenytoin, carbamazepine, rifampin, St. John's Wort and glucocorticoids two weeks prior to the trial, depot anti-psychotic within one dosing interval (for the depot) before randomisation, CCMD diagnosis of substance or alcohol dependence at enrolment (exception of caffeine or nicotine), use of opiates, amphetamine, cocaine, cannabis, barbiturates or hallucinogens within four weeks prior, medical conditions that would affect the absorption, distribution, metabolism, or excretion of study treatment, unstable or inadequately treated medical illnesses, involvement in the planning and conduct of the study, previous enrolment or randomisation in the present study, participation in another drug trial within four weeks prior to enrolment or in accordance with local requirements, a person with diabetes mellitus, an absolute neutrophil count of 1.5×10^9 per litre, 2 x higher than normal upper limit of ALT or AST, use of clozapine within 28 days prior to randomisation.

Setting: psychiatric inpatient ward, China.

Interventions	1. Haloperidol. 2. Quetiapine.
Outcomes	Not stated.
Starting date	August 2008.
Contact information	Bo Zhang, PhD 13808203275 zb_73@126.com
Notes	Protocol, full characteristics and outcome data not reported. Attempted to contact author to enquire whether there are any available data.

ALT: alanine aminotransferase

AST: aspartate transaminase

CCMD: Chinese Classification of Mental Disorders

DATA AND ANALYSES

Comparison 1. HALOPERIDOL vs PLACEBO/NIL

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Tranquillisation or asleep - asleep	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
1.1 not asleep up to 2 hours	2	220	Risk Ratio (M-H, Fixed, 95% CI)	0.88 [0.82, 0.95]
2 Repeated need for tranquillisation	4		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
2.1 needing additional injection during 24 hours (agitation only)	4	660	Risk Ratio (M-H, Fixed, 95% CI)	0.51 [0.42, 0.62]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
3 Specific behaviour: 1a. Agitation - various measures	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
3.1 PANSS-EC response at 2 hours (at least 40% change on PANSS-EC from baseline)	2	395	Risk Ratio (M-H, Fixed, 95% CI)	1.62 [1.28, 2.07]
4 Specific behaviour: 1b. Agitation - average scores - i. up to 2 hours	4		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
4.1 change score (ABS, high=worse)	3	474	Mean Difference (IV, Fixed, 95% CI)	-4.48 [-5.78, -3.18]
4.2 change score (ACES, low=agitated, high=sedated)	2	390	Mean Difference (IV, Fixed, 95% CI)	0.83 [0.39, 1.26]
4.3 score (PANSS-EC, high=worse)	3	514	Mean Difference (IV, Fixed, 95% CI)	-3.67 [-4.75, -2.59]
4.4 change score (PANSS-EC, high=worse, schizophrenia subgroup)	1	199	Mean Difference (IV, Fixed, 95% CI)	-2.57 [-4.05, -1.09]
4.5 change score (PANSS-EC, high=worse, non-sedated patients subgroup based on ACES)	1	240	Mean Difference (IV, Fixed, 95% CI)	-2.64 [-3.95, -1.33]
4.6 change score (PANSS-EC, high=worse, non-sedated patients subgroup based on ACES)	1	263	Mean Difference (IV, Fixed, 95% CI)	-2.97 [-4.27, -1.67]
5 Specific behaviour: 1c. Agitation - average scores - ii. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
5.1 change score (ABS, high=worse)	1	85	Mean Difference (IV, Fixed, 95% CI)	-2.4 [-4.13, -0.67]
5.2 change score (PANSS-EC, high=worse)	1	85	Mean Difference (IV, Fixed, 95% CI)	-1.4 [-2.97, 0.17]
6 Global outcome: 1. Not improved	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
6.1 not marked improvement	1	40	Risk Ratio (M-H, Fixed, 95% CI)	0.61 [0.44, 0.84]
6.2 not any improvement	1	40	Risk Ratio (M-H, Fixed, 95% CI)	0.28 [0.08, 1.07]
7 Global outcome: 2. Need for benzodiazepine up to 24 hours	4		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
7.1 need for benzodiazepine up to 24 hours	4	660	Risk Ratio (M-H, Fixed, 95% CI)	0.44 [0.31, 0.62]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
8 Global outcome: 3a. Average scores - i. up to 2 hours	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
8.1 endpoint score (CGI-I, high=worse)	2	390	Mean Difference (IV, Fixed, 95% CI)	-0.73 [-0.96, -0.51]
8.2 change score (CGI-I, high=worse, schizophrenia subgroup)	1	199	Mean Difference (IV, Fixed, 95% CI)	-0.64 [-0.98, -0.30]
8.3 change score (CGI-S, high=worse)	2	390	Mean Difference (IV, Fixed, 95% CI)	-0.48 [-0.78, -0.18]
9 Global outcome: 3b. Average scores - ii. up to 24 hours	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
9.1 change score (CGI-I, high=worse)	1	169	Mean Difference (IV, Fixed, 95% CI)	-0.4 [-0.62, -0.18]
9.2 change score (CGI-S, high=worse)	1	85	Mean Difference (IV, Fixed, 95% CI)	-0.2 [-0.46, 0.06]
10 Mental state: 1a. Average scores - i. up to 2 hours	3		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
10.1 change score up to 2 hours (BPRS positive sub-scale, high=worse)	3	372	Mean Difference (IV, Fixed, 95% CI)	-1.04 [-1.54, -0.54]
10.2 change score up to 2 hours (BPRS total, high=worse)	3	371	Mean Difference (IV, Fixed, 95% CI)	-5.69 [-7.38, -4.00]
11 Mental state: 1b. Average scores - ii. up to 24 hours	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
11.1 change score up to 24 hours (BPRS positive sub-scale, high=worse)	2	264	Mean Difference (IV, Fixed, 95% CI)	-1.61 [-2.34, -0.89]
11.2 change score up to 24 hours (BPRS total, high=worse)	2	264	Mean Difference (IV, Fixed, 95% CI)	-4.81 [-6.82, -2.81]
12 Adverse effects: 1a. General	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
12.1 one or more drug related adverse events during 24 hours	2	395	Risk Ratio (M-H, Fixed, 95% CI)	1.64 [1.22, 2.20]
12.2 increased severity of adverse effects after 2nd injection	1	273	Risk Ratio (M-H, Fixed, 95% CI)	3.25 [1.88, 5.63]
12.3 overall adverse events during 72 hours	1	273	Risk Ratio (M-H, Fixed, 95% CI)	1.78 [1.23, 2.59]
13 Adverse effects: 1b. General - serious	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
13.1 death	1	273	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
13.2 rated as serious	1	122	Risk Ratio (M-H, Fixed, 95% CI)	0.34 [0.01, 8.29]
13.3 tonic clonic seizure	1	117	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
14 Adverse effects: 2a. Specific - arousal level	4		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
14.1 insomnia during 24 hours (only reported if occurred in $\geq 5\%$)	1	273	Risk Ratio (M-H, Fixed, 95% CI)	1.31 [0.61, 2.82]
14.2 "over" sedated	2	313	Risk Ratio (M-H, Fixed, 95% CI)	3.36 [1.42, 7.99]
14.3 somnolence during 24 hours	4	615	Risk Ratio (M-H, Fixed, 95% CI)	2.28 [0.97, 5.36]
15 Adverse effects: 2b. Specific - cardiovascular - miscellaneous outcomes	4		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
15.1 dizziness during 24 hours (only reported if occurred in $\geq 5\%$)	2	392	Risk Ratio (M-H, Fixed, 95% CI)	1.33 [0.48, 3.65]
15.2 hypotension during 24 hours	2	125	Risk Ratio (M-H, Fixed, 95% CI)	1.2 [0.05, 27.44]
15.3 QTc abnormality	1	122	Risk Ratio (M-H, Fixed, 95% CI)	3.1 [0.33, 28.98]
15.4 sinus tachycardia during 24 hours (only reported if occurred in $\geq 5\%$)	1	122	Risk Ratio (M-H, Fixed, 95% CI)	3.1 [0.33, 28.98]
15.5 tachycardia during 24 hours (only reported if occurred in $\geq 5\%$)	1	122	Risk Ratio (M-H, Fixed, 95% CI)	1.03 [0.07, 16.15]
16 Adverse effects: 2c. Specific - cardiovascular - QTc interval (average change at 24 hours)	2	265	Mean Difference (IV, Fixed, 95% CI)	3.63 [-2.67, 9.93]
17 Adverse effects: 2d. Specific - movement disorders	5		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
17.1 akathisia during 24 hours (only reported if occurred in $\geq 5\%$)	1	122	Risk Ratio (M-H, Fixed, 95% CI)	13.43 [0.77, 233.23]
17.2 dystonia during 24 hours	2	207	Risk Ratio (M-H, Fixed, 95% CI)	7.49 [0.93, 60.21]
17.3 extrapyramidal effects - use of antiparkinson drugs during 24 hours	1	180	Risk Ratio (M-H, Fixed, 95% CI)	5.57 [1.37, 22.65]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
17.4 EPS during 24 hours	3	398	Risk Ratio (M-H, Fixed, 95% CI)	6.79 [2.19, 21.07]
18 Adverse effects: 2e. Specific - movement disorders: i. average scores - i. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
18.1 change score at 24 hours (BAS, high=worse)	1	168	Mean Difference (IV, Fixed, 95% CI)	0.09 [-0.17, 0.35]
18.2 change score at 24 hours (SAS, high=worse)	1	167	Mean Difference (IV, Fixed, 95% CI)	1.89 [0.77, 3.01]
19 Adverse effects: 2f. Specific - miscellaneous	3		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
19.1 agitation during 24 hours (only reported if occurred in $\geq 5\%$)	2	395	Risk Ratio (M-H, Fixed, 95% CI)	0.80 [0.29, 2.19]
19.2 dry mouth	1	40	Risk Ratio (M-H, Fixed, 95% CI)	3.6 [0.21, 61.86]
19.3 headache during 24 hours (only reported if occurred in $\geq 5\%$)	2	395	Risk Ratio (M-H, Fixed, 95% CI)	1.28 [0.55, 3.00]
19.4 pain at injection site	1	122	Risk Ratio (M-H, Fixed, 95% CI)	0.21 [0.01, 4.22]
19.5 pain (1st injection)	1	273	Risk Ratio (M-H, Fixed, 95% CI)	0.10 [0.00, 1.97]
19.6 pain (2nd injection)	1	273	Risk Ratio (M-H, Fixed, 95% CI)	3.25 [1.88, 5.63]
19.7 nausea during 24 hours (only reported if occurred in $\geq 5\%$)	2	395	Risk Ratio (M-H, Fixed, 95% CI)	0.69 [0.13, 3.67]
19.8 vomiting during 24 hours (only reported if occurred in $\geq 5\%$)	1	122	Risk Ratio (M-H, Fixed, 95% CI)	1.03 [0.07, 16.15]
20 Leaving the study early	3		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
20.1 any reason	3	435	Risk Ratio (M-H, Fixed, 95% CI)	1.06 [0.53, 2.11]
20.2 lack of efficacy	3	435	Risk Ratio (M-H, Fixed, 95% CI)	0.12 [0.03, 0.49]
20.3 withdrew	1	273	Risk Ratio (M-H, Fixed, 95% CI)	3.35 [0.17, 64.15]
20.4 consent	2	395	Risk Ratio (M-H, Fixed, 95% CI)	6.2 [0.77, 49.98]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
20.5 adverse effects	3	435	Risk Ratio (M-H, Fixed, 95% CI)	1.78 [0.21, 15.39]
20.6 could not be evaluated due to being asleep	1	40	Risk Ratio (M-H, Fixed, 95% CI)	1.2 [0.05, 27.44]
20.7 other	2	395	Risk Ratio (M-H, Fixed, 95% CI)	1.81 [0.27, 12.07]

Analysis 1.1. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 1 Tranquillisation or asleep - asleep.

Analysis 1.2. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 2 Repeated need for tranquillisation.

Analysis 1.3. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 3 Specific behaviour: 1a. Agitation - various measures.

Analysis 1.4. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 4 Specific behaviour: 1b. Agitation - average scores - i. up to 2 hours.

Analysis 1.5. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 5 Specific behaviour: 1c. Agitation - average scores - ii. up to 24 hours.

Analysis 1.6. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 6 Global outcome: 1. Not improved.

Analysis 1.7. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 7 Global outcome: 2. Need for benzodiazepine up to 24 hours.

Analysis 1.8. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 8 Global outcome: 3a. Average scores - i. up to 2 hours.

Analysis 1.9. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 9 Global outcome: 3b. Average scores - ii. up to 24 hours.

Analysis 1.10. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 10 Mental state: 1a. Average scores - i. up to 2 hours.

Analysis 1.11. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 11 Mental state: 1b. Average scores - ii. up to 24 hours.

Analysis 1.12. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 12 Adverse effects: 1a. General.

Analysis 1.13. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 13 Adverse effects: 1b. General - serious.

Analysis 1.14. Comparison 1 HALOPERIDOL vs PLACEBO/ NIL, Outcome 14 Adverse effects: 2a. Specific - arousal level.

Analysis 1.15. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 15 Adverse effects: 2b. Specific - cardiovascular - miscellaneous outcomes.

Analysis 1.16. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 16 Adverse effects: 2c. Specific - cardiovascular - QTc interval (average change at 24 hours).

Analysis 1.17. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 17 Adverse effects: 2d. Specific - movement disorders.

Analysis 1.18. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 18 Adverse effects: 2e. Specific - movement disorders: i. average scores - i. up to 24 hours.

Analysis 1.19. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 19 Adverse effects: 2f. Specific - miscellaneous.

Analysis 1.20. Comparison 1 HALOPERIDOL vs PLACEBO/NIL, Outcome 20 Leaving the study early.

Study or subgroup	haloperidol n/N	placebo n/N	Risk Ratio M-H, Fixed, 95% CI	Weight	Risk Ratio M-H, Fixed, 95% CI
1.20.2 withdrawal					
Bristol Myers 2004a	1/185	1/88		11.19%	0.48[0.03,7.52]
Bristol-Myers 2005b	0/60	1/62		12.18%	0.34[0.01,8.29]
Reschke 1974	0/29	6/11		76.63%	0.03[0,0.5]
Subtotal (95% CI)	274	161		100%	0.12[0.03,0.49]
Total events: 1 (haloperidol), 8 (placebo)					
Heterogeneity: Tau²=0; Chi²=2.3, df=2(P=0.32); I²=12.91%					
Test for overall effect: Z=2.95(P=0)					
1.20.3 withdrew					
Bristol Myers 2004a	3/185	0/88		100%	3.35[0.17,64.15]
Subtotal (95% CI)	185	88		100%	3.35[0.17,64.15]
Total events: 3 (haloperidol), 0 (placebo)					
Heterogeneity: Not applicable					
Test for overall effect: Z=0.8(P=0.42)					
1.20.4 consent					
Bristol Myers 2004a	0/185	0/88			Not estimable
Bristol-Myers 2005b	6/60	1/62		100%	6.2[0.77,49.98]
Subtotal (95% CI)	245	150		100%	6.2[0.77,49.98]
Total events: 6 (haloperidol), 1 (placebo)					
Heterogeneity: Not applicable					
Test for overall effect: Z=1.71(P=0.09)					
1.20.5 adverse effects					
Bristol Myers 2004a	2/185	0/88		48.64%	2.39[0.12,49.31]
Bristol-Myers 2005b	0/60	0/62			Not estimable
Reschke 1974	1/29	0/11		51.36%	1.2[0.05,27.44]
Subtotal (95% CI)	274	161		100%	1.78[0.21,15.39]
Total events: 3 (haloperidol), 0 (placebo)					
Heterogeneity: Tau²=0; Chi²=0.1, df=1(P=0.75); I²=0%					
Test for overall effect: Z=0.52(P=0.6)					
1.20.6 could not be evaluated due to being asleep					
Reschke 1974	1/29	0/11		100%	1.2[0.05,27.44]
Subtotal (95% CI)	29	11		100%	1.2[0.05,27.44]
Total events: 1 (haloperidol), 0 (placebo)					
Heterogeneity: Not applicable					
Test for overall effect: Z=0.11(P=0.91)					
1.20.7 other					
Bristol Myers 2004a	1/185	0/88		40.75%	1.44[0.06,34.89]
Bristol-Myers 2005b	2/60	1/62		59.25%	2.07[0.19,22.2]
Subtotal (95% CI)	245	150		100%	1.81[0.27,12.07]
Total events: 3 (haloperidol), 1 (placebo)					
Heterogeneity: Tau²=0; Chi²=0.03, df=1(P=0.86); I²=0%					
Test for overall effect: Z=0.61(P=0.54)					
Test for subgroup differences: Chi²=12.95, df=1 (P=0.04), I²=53.65%					

Comparison 2. HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Repeated need for rapid tranquillisation: needing additional injection	2	473	Risk Ratio (M-H, Fixed, 95% CI)	0.78 [0.62, 0.99]
2 Specific behaviour: 1a. Agitation - PANSS-EC response up to 2 hours (at least 40% change on PANSS-EC from baseline)	2	477	Risk Ratio (M-H, Fixed, 95% CI)	1.07 [0.92, 1.26]
3 Specific behaviour: 1b. Agitation - average scores - up to 2 hours	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
3.1 change score (ACES, low=agitated, high=sedated)	2	470	Mean Difference (IV, Fixed, 95% CI)	0.12 [-0.30, 0.55]
3.2 change score (CABS, high=worse)	2	470	Mean Difference (IV, Fixed, 95% CI)	-0.55 [-2.10, 1.01]
3.3 change score (PANSS-EC, high=worse)	1	357	Mean Difference (IV, Fixed, 95% CI)	-0.48 [-2.12, 1.16]
3.4 change score (PANSS-EC, high=worse, schizophrenia subgroup)	1	257	Mean Difference (IV, Fixed, 95% CI)	-0.26 [-1.48, 0.96]
3.5 change score (PANSS-EC, high=worse, non-sedated patients subgroup based on ACES)	1	316	Mean Difference (IV, Fixed, 95% CI)	-0.33 [-1.42, 0.76]
3.6 change score (PANSS-EC, high=worse, non-sedated patients subgroup based on AEs)	1	346	Mean Difference (IV, Fixed, 95% CI)	-0.34 [-1.40, 0.72]
4 Global outcome: 1. Need for benzodiazepine	2	477	Risk Ratio (M-H, Fixed, 95% CI)	1.26 [0.74, 2.16]
5 Global outcome: 2. Average scores - up to 2 hours	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
5.1 change score (CGI-S, high=worse)	2	470	Mean Difference (IV, Fixed, 95% CI)	0.07 [-0.22, 0.36]
5.2 endpoint score (CGI-I, high=worse)	2	470	Mean Difference (IV, Fixed, 95% CI)	-0.02 [-0.23, 0.19]
5.3 change score (CGI-I, high=worse, schizophrenia subgroup)	1	257	Mean Difference (IV, Fixed, 95% CI)	-0.02 [-0.30, 0.26]
6 Mental state: 1. Average scores - up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
6.1 change score at 2 hours (BPRS positive sub-scale, high=worse)	1	103	Mean Difference (IV, Fixed, 95% CI)	-0.23 [-1.28, 0.82]
6.2 change score at 2 hours (BPRS total, high=worse)	1	102	Mean Difference (IV, Fixed, 95% CI)	-2.03 [-5.76, 1.70]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
7 Adverse effects: 1a. General	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
7.1 one or more drug related adverse effects during 24 hours	2	477	Risk Ratio (M-H, Fixed, 95% CI)	1.18 [0.95, 1.46]
7.2 increased severity of adverse effects after 2nd injection	1	360	Risk Ratio (M-H, Fixed, 95% CI)	1.34 [1.03, 1.74]
7.3 overall adverse events during 72 hours	1	360	Risk Ratio (M-H, Fixed, 95% CI)	1.33 [1.04, 1.70]
8 Adverse effects: 1b. General - serious	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
8.1 any	2	477	Risk Ratio (M-H, Fixed, 95% CI)	0.57 [0.18, 1.81]
8.2 tonic clonic seizure	1	117	Risk Ratio (M-H, Fixed, 95% CI)	0.32 [0.01, 7.62]
8.3 death	1	360	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
9 Adverse effects: 2a. Specific - arousal level	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
9.1 insomnia - between 0-1 days (IM phase) (only reported if occurred in $\geq 5\%$)	1	360	Risk Ratio (M-H, Fixed, 95% CI)	2.08 [1.01, 4.27]
9.2 insomnia - between 3 - 7 days (oral phase) (only reported if occurred in $\geq 2\%$)	1	360	Risk Ratio (M-H, Fixed, 95% CI)	0.82 [0.47, 1.44]
9.3 "over" sedated	1	360	Risk Ratio (M-H, Fixed, 95% CI)	1.66 [0.93, 2.95]
9.4 somnolence - between 0-1 days	1	360	Risk Ratio (M-H, Fixed, 95% CI)	0.71 [0.25, 2.00]
9.5 somnolence - between 0-1 days (IM phase) (only reported if occurred in $\geq 5\%$)	1	117	Risk Ratio (M-H, Fixed, 95% CI)	2.22 [0.60, 8.16]
9.6 somnolence - between 3 - 7 days (oral phase) (only reported if occurred in $\geq 2\%$)	1	360	Risk Ratio (M-H, Fixed, 95% CI)	2.84 [0.30, 27.02]
10 Adverse effects: 2b. Specific - cardiac	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
10.1 dizziness - between 0 -1 day (IM phase) (only reported if occurred in $\geq 5\%$)	2	477	Risk Ratio (M-H, Fixed, 95% CI)	0.69 [0.33, 1.48]
10.2 sinus tachycardia - between 0 -1 day (IM phase) (only reported if occurred in $\geq 5\%$)	1	117	Risk Ratio (M-H, Fixed, 95% CI)	6.66 [0.35, 126.06]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
10.3 tachycardia - between 0 -1 day (IM phase) (only reported if occurred in $\geq 5\%$)	1	117	Risk Ratio (M-H, Fixed, 95% CI)	0.24 [0.03, 2.06]
11 Adverse effects: 2c. Specific - gastrointestinal	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
11.1 dyspepsia - between 3 - 7 days (oral phase) (only reported if occurred in $\geq 2\%$)	1	360	Risk Ratio (M-H, Fixed, 95% CI)	10.41 [1.36, 79.76]
11.2 diarrhoea - between 3 - 7 days (oral phase) (only reported if occurred in $\geq 2\%$)	1	360	Risk Ratio (M-H, Fixed, 95% CI)	0.95 [0.28, 3.21]
11.3 nausea - between 0-1 days (IM phase) (only reported if occurred in $\geq 5\%$)	2	477	Risk Ratio (M-H, Fixed, 95% CI)	0.18 [0.05, 0.60]
11.4 nausea - between 3 - 7 days (oral phase) (only reported if occurred in $\geq 2\%$)	1	360	Risk Ratio (M-H, Fixed, 95% CI)	0.14 [0.02, 1.09]
11.5 vomiting - between 0-1 days (IM phase) (only reported if occurred in $\geq 5\%$)	1	117	Risk Ratio (M-H, Fixed, 95% CI)	0.48 [0.04, 5.10]
11.6 vomiting - between 3 - 7 days (oral phase) (only reported if occurred in $\geq 2\%$)	1	360	Risk Ratio (M-H, Fixed, 95% CI)	0.38 [0.07, 1.92]
12 Adverse effects: 2d. Specific - movement disorder	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
12.1 akathisia - between 0 -1 day (IM phase) (only reported if occurred in $\geq 5\%$)	2	477	Risk Ratio (M-H, Fixed, 95% CI)	2.85 [0.94, 8.61]
12.2 akathisia - between 3 - 7 days (oral phase) (only reported if occurred in $\geq 2\%$)	1	360	Risk Ratio (M-H, Fixed, 95% CI)	2.21 [0.58, 8.40]
12.3 dystonia - between 0 -1 day (IM phase) (only reported if occurred in $\geq 5\%$)	2	477	Risk Ratio (M-H, Fixed, 95% CI)	6.63 [1.52, 28.86]
12.4 EPS - between 0 -1 day (IM phase) (only reported if occurred in $\geq 5\%$)	1	360	Risk Ratio (M-H, Fixed, 95% CI)	9.46 [1.22, 73.13]
12.5 EPS - between 3 - 7 days (oral phase) (only reported if occurred in $\geq 2\%$)	1	360	Risk Ratio (M-H, Fixed, 95% CI)	7.09 [1.65, 30.58]
12.6 Use of antiparkinson drugs - between 3 - 7 days (oral phase)	1	360	Risk Ratio (M-H, Fixed, 95% CI)	4.94 [2.50, 9.78]
12.7 dystonia	1	360	Risk Ratio (M-H, Fixed, 95% CI)	2.84 [0.12, 69.22]
13 Adverse effects: 2e. Specific - miscellaneous	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
13.1 agitation - between 0 -1 day (IM phase) (only reported if occurred in $\geq 5\%$)	2	477	Risk Ratio (M-H, Fixed, 95% CI)	1.22 [0.46, 3.22]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
13.2 agitation - between 3 - 7 days (oral phase) (only reported if occurred in $\geq 2\%$)	1	360	Risk Ratio (M-H, Fixed, 95% CI)	1.09 [0.68, 1.74]
13.3 anxiety - between 3 - 7 days (oral phase) (only reported if occurred in $\geq 2\%$)	1	360	Risk Ratio (M-H, Fixed, 95% CI)	1.30 [0.54, 3.16]
13.4 headache - between 0 -1 day (IM phase) (only reported if occurred in $\geq 5\%$)	2	477	Risk Ratio (M-H, Fixed, 95% CI)	0.85 [0.45, 1.59]
13.5 headache - between 3 - 7 days (oral phase) (only reported if occurred in $\geq 2\%$)	1	360	Risk Ratio (M-H, Fixed, 95% CI)	1.06 [0.57, 1.97]
13.6 pain at injection site	1	117	Risk Ratio (M-H, Fixed, 95% CI)	0.32 [0.01, 7.62]
13.7 pain (1st injection)	1	360	Risk Ratio (M-H, Fixed, 95% CI)	0.32 [0.06, 1.54]
13.8 pain (2nd injection)	1	360	Risk Ratio (M-H, Fixed, 95% CI)	1.34 [1.03, 1.74]
14 Leaving the study early: 1. For specific reasons	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
14.1 any	2	480	Risk Ratio (M-H, Fixed, 95% CI)	2.07 [0.86, 4.98]
14.2 lack of efficacy	1	360	Risk Ratio (M-H, Fixed, 95% CI)	2.84 [0.12, 69.22]
14.3 adverse effects	2	480	Risk Ratio (M-H, Fixed, 95% CI)	0.97 [0.17, 5.59]
14.4 withdrew	1	360	Risk Ratio (M-H, Fixed, 95% CI)	1.42 [0.24, 8.39]
14.5 consent	2	480	Risk Ratio (M-H, Fixed, 95% CI)	6.0 [0.74, 48.34]
14.6 other	2	480	Risk Ratio (M-H, Fixed, 95% CI)	1.36 [0.27, 6.75]

Analysis 2.1. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 1 Repeated need for rapid tranquillisation: needing additional injection.

Analysis 2.2. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 2 Specific behaviour: 1a. Agitation - PANSS-EC response up to 2 hours (at least 40% change on PANSS-EC from baseline).

Analysis 2.3. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 3 Specific behaviour: 1b. Agitation - average scores - up to 2 hours.

Analysis 2.4. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 4 Global outcome: 1. Need for benzodiazepine.

Analysis 2.5. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 5 Global outcome: 2. Average scores - up to 2 hours.

Analysis 2.6. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIRAZOLE, Outcome 6 Mental state: 1. Average scores - up to 2 hours.

Analysis 2.7. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIRAZOLE, Outcome 7 Adverse effects: 1a. General.

Analysis 2.8. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 8 Adverse effects: 1b. General - serious.

Analysis 2.9. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 9 Adverse effects: 2a. Specific - arousal level.

**Analysis 2.10. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC:
a. ARIPIPRAZOLE, Outcome 10 Adverse effects: 2b. Specific - cardiac.**

**Analysis 2.11. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a.
ARIPIPRAZOLE, Outcome 11 Adverse effects: 2c. Specific - gastrointestinal.**

Analysis 2.12. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 12 Adverse effects: 2d. Specific - movement disorder.

Analysis 2.13. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIPRAZOLE, Outcome 13 Adverse effects: 2e. Specific - miscellaneous.

Analysis 2.14. Comparison 2 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: a. ARIPIRAZOLE, Outcome 14 Leaving the study early: 1. For specific reasons.

Comparison 3. HALOPERIDOL vs OTHER ANTIPSYCHOTIC: b. CHLORPROMAZINE

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Tranquillisation or asleep - asleep	1	39	Risk Ratio (M-H, Fixed, 95% CI)	1.93 [1.04, 3.60]
1.1 not asleep up to 2 hours	1	39	Risk Ratio (M-H, Fixed, 95% CI)	1.93 [1.04, 3.60]
2 Repeated need for rapid tranquillisation	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
2.1 more that 1 injection	1	30	Risk Ratio (M-H, Fixed, 95% CI)	1.07 [0.89, 1.28]
2.2 more than 3 injections	1	30	Risk Ratio (M-H, Fixed, 95% CI)	1.5 [0.53, 4.26]
3 Global outcome: 1. Not improved	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
3.1 not marked improvement	2	89	Risk Ratio (M-H, Fixed, 95% CI)	0.79 [0.61, 1.02]
3.2 not any improvement	2	89	Risk Ratio (M-H, Fixed, 95% CI)	0.15 [0.05, 0.49]
4 Adverse effects: any serious or specific adverse effects	3		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
4.1 allergy - haematological - leucopenia - mild	1	50	Risk Ratio (M-H, Fixed, 95% CI)	3.0 [0.13, 70.30]
4.2 allergy - hepatic - glutamic pyruvic transaminase elevated	1	50	Risk Ratio (M-H, Fixed, 95% CI)	0.33 [0.01, 7.81]
4.3 allergy - skin irritation - local	1	30	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
4.4 anticholinergic - dry mouth	1	39	Risk Ratio (M-H, Fixed, 95% CI)	1.38 [0.17, 10.93]
4.5 arousal - drowsy but asleep	1	39	Risk Ratio (M-H, Fixed, 95% CI)	0.06 [0.01, 0.42]
4.6 arousal - drowsy but awake	1	39	Risk Ratio (M-H, Fixed, 95% CI)	9.17 [0.59, 142.10]
4.7 cardiovascular - hypotension	2	69	Risk Ratio (M-H, Fixed, 95% CI)	0.51 [0.10, 2.60]
4.8 central nervous system - seizures	1	30	Risk Ratio (M-H, Fixed, 95% CI)	0.33 [0.01, 7.58]
4.9 movement disorders - extrapyramidal adverse effects	2	69	Risk Ratio (M-H, Fixed, 95% CI)	2.07 [0.28, 15.15]
5 Leaving the study early	4		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
5.1 any reason/general reasons	4	153	Risk Ratio (M-H, Fixed, 95% CI)	0.21 [0.07, 0.71]
5.2 could not be evaluated due to being asleep	1	39	Risk Ratio (M-H, Fixed, 95% CI)	0.07 [0.01, 0.52]
5.3 severe hypotensive reaction	1	30	Risk Ratio (M-H, Fixed, 95% CI)	0.2 [0.01, 3.85]
5.4 deviation from protocol	1	30	Risk Ratio (M-H, Fixed, 95% CI)	3.0 [0.13, 68.26]

Analysis 3.1. Comparison 3 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: b. CHLORPROMAZINE, Outcome 1 Tranquillisation or asleep - asleep.

Analysis 3.2. Comparison 3 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: b. CHLORPROMAZINE, Outcome 2 Repeated need for rapid tranquillisation.

Analysis 3.3. Comparison 3 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: b. CHLORPROMAZINE, Outcome 3 Global outcome: 1. Not improved.

Analysis 3.4. Comparison 3 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: b. CHLORPROMAZINE, Outcome 4 Adverse effects: any serious or specific adverse effects.

Analysis 3.5. Comparison 3 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: b. CHLORPROMAZINE, Outcome 5 Leaving the study early.

Comparison 4. HALOPERIDOL vs OTHER ANTIPSYCHOTIC: c. DROPERIDOL

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Tranquillisation or asleep: 1. Not asleep	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
1.1 up to 2 hours	1	228	Risk Ratio (M-H, Fixed, 95% CI)	1.07 [0.44, 2.60]
2 Tranquillisation or asleep: 2. Time to sleep	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
2.1 up to 2 hours	1	114	Mean Difference (IV, Fixed, 95% CI)	5.20 [-6.05, 16.45]
3 Repeated need for rapid tranquillisation	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
3.1 more than 1 injection	2	255	Risk Ratio (M-H, Fixed, 95% CI)	2.38 [1.27, 4.47]
3.2 more than 3 injections	1	27	Risk Ratio (M-H, Fixed, 95% CI)	2.12 [0.09, 47.68]
4 Global outcome: 1. Need for benzodiazepine	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
4.1 up to 2 hours	1	228	Risk Ratio (M-H, Fixed, 95% CI)	0.31 [0.07, 1.44]
5 Adverse effects: 1. General	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
5.1 overall adverse events up to 2 hours	1	228	Risk Ratio (M-H, Fixed, 95% CI)	0.15 [0.02, 1.23]
5.2 patients with one or more AEs	1	228	Risk Ratio (M-H, Fixed, 95% CI)	0.18 [0.02, 1.46]
6 Adverse effects: 2a. Specific - arousal level	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
6.1 "over" sedated	1	228	Risk Ratio (M-H, Fixed, 95% CI)	0.36 [0.01, 8.68]
7 Adverse effects: 2b. Specific - cardiovascular	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
7.1 hypotension up to 2 hours	1	228	Risk Ratio (M-H, Fixed, 95% CI)	0.27 [0.03, 2.36]
8 Adverse effects: 2c. Specific - movement disorders	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
8.1 dystonia	2	255	Risk Ratio (M-H, Fixed, 95% CI)	0.86 [0.11, 6.56]
9 Adverse effects: 2d. Specific - miscellaneous	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
9.1 desaturation	1	228	Risk Ratio (M-H, Fixed, 95% CI)	0.36 [0.01, 8.68]

Analysis 4.1. Comparison 4 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: c. DROPERIDOL, Outcome 1 Tranquillisation or asleep: 1. Not asleep.

Analysis 4.2. Comparison 4 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: c. DROPERIDOL, Outcome 2 Tranquillisation or asleep: 2. Time to sleep.

Analysis 4.3. Comparison 4 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: c. DROPERIDOL, Outcome 3 Repeated need for rapid tranquillisation.

Analysis 4.4. Comparison 4 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: c. DROPERIDOL, Outcome 4 Global outcome: 1. Need for benzodiazepine.

Analysis 4.5. Comparison 4 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: c. DROPERIDOL, Outcome 5 Adverse effects: 1. General.

**Analysis 4.6. Comparison 4 HALOPERIDOL vs OTHER ANTIPSYCHOTIC:
c. DROPERIDOL, Outcome 6 Adverse effects: 2a. Specific - arousal level.**

**Analysis 4.7. Comparison 4 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: c.
DROPERIDOL, Outcome 7 Adverse effects: 2b. Specific - cardiovascular.**

**Analysis 4.8. Comparison 4 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: c.
DROPERIDOL, Outcome 8 Adverse effects: 2c. Specific - movement disorders.**

Comparison 5. HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Tranquillisation or asleep - not asleep up to 24 hours	1	54	Risk Ratio (M-H, Fixed, 95% CI)	4.31 [0.54, 34.48]
2 Specific behaviour: 2. Aggression - various measures	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
2.1 no overall improvement	1	30	Risk Ratio (M-H, Fixed, 95% CI)	1.1 [0.69, 1.76]
3 Global outcome: 1. General - no change - i. over 24 hours	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
3.1 up to 48 hours (CGI-I, high=worse)	1	56	Risk Ratio (M-H, Fixed, 95% CI)	0.93 [0.14, 6.15]
3.2 up to 72 hours (CGI-I, high=worse)	1	54	Risk Ratio (M-H, Fixed, 95% CI)	0.39 [0.04, 3.49]
3.3 at endpoint (CGI-I, high=worse)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	0.40 [0.12, 1.32]
4 Global outcome: 2a. Specific - not sedated - i. by 30 minutes	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
4.1 at 30 minutes	1	30	Risk Ratio (M-H, Fixed, 95% CI)	1.29 [0.65, 2.54]
5 Global outcome: 2b. Specific - not sedated - ii. up to 2 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
5.1 at 1 hour	1	30	Risk Ratio (M-H, Fixed, 95% CI)	3.5 [0.86, 14.18]
5.2 at 2 hours	1	30	Risk Ratio (M-H, Fixed, 95% CI)	7.0 [0.98, 50.16]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
6 Mental state: 1. Average scores	1	52	Mean Difference (IV, Fixed, 95% CI)	6.10 [4.48, 7.72]
6.1 endpoint score (BPRS, high=worse)	1	52	Mean Difference (IV, Fixed, 95% CI)	6.10 [4.48, 7.72]
7 Adverse effects: 1. General	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
7.1 one or more drug related adverse effect	1	30	Risk Ratio (M-H, Fixed, 95% CI)	0.8 [0.44, 1.45]
8 Adverse effects: 2a. Specific - anti-cholinergic	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
8.1 constipation - between 3-28 days (Oral phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	0.32 [0.01, 7.26]
8.2 salivation - too little - between 0-3 days (IM phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	2.83 [0.33, 24.66]
8.3 salivation - too little - between 3-28 days (Oral phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	0.94 [0.22, 4.05]
8.4 salivation - too much - between 0-3 days (IM phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	0.94 [0.33, 2.69]
8.5 salivation - too much - between 0-10 days	1	54	Risk Ratio (M-H, Fixed, 95% CI)	0.43 [0.04, 4.48]
8.6 salivation - too much - between 3-28 days (Oral phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	1.05 [0.57, 1.93]
8.7 polyuria - between 3-28 days (Oral phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	2.84 [0.12, 65.34]
8.8 sweating - between 0-3 days (IM phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	0.94 [0.15, 5.97]
8.9 sweating - between 3-28 days (Oral phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	2.84 [0.12, 65.34]
9 Adverse effects: 2b. Specific - arousal level	3		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
9.1 asleep - by 12 hours	1	54	Risk Ratio (M-H, Fixed, 95% CI)	0.86 [0.72, 1.04]
9.2 drowsiness - between 0-3 days (IM phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	0.89 [0.70, 1.12]
9.3 drowsiness - between 3-28 days (Oral phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	33.16 [2.15, 511.57]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
9.4 drowsiness during 72 hours	1	30	Risk Ratio (M-H, Fixed, 95% CI)	0.75 [0.20, 2.79]
9.5 insomnia - between 0-3 days (IM phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	0.32 [0.01, 7.26]
9.6 insomnia - between 3-28 days (Oral phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	6.63 [0.37, 119.59]
10 Adverse effects: 2c. Specific - cardiovascular	3		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
10.1 blood pressure - increased - between 0-10 days	1	54	Risk Ratio (M-H, Fixed, 95% CI)	0.17 [0.01, 3.45]
10.2 dizziness - between 3-28 days	2	65	Risk Ratio (M-H, Fixed, 95% CI)	0.38 [0.09, 1.52]
10.3 dyspnoea - between 3-28 days (Oral phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	2.84 [0.12, 65.34]
10.4 palpitations during 72 hours	1	30	Risk Ratio (M-H, Fixed, 95% CI)	0.33 [0.01, 7.58]
10.5 pulse rate - increased - between 0-10 days	1	54	Risk Ratio (M-H, Fixed, 95% CI)	0.29 [0.01, 6.79]
11 Adverse effects: 2d. Specific - movement disorders	3		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
11.1 akathisia between 0-3 days (IM phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	0.57 [0.16, 2.02]
11.2 akathisia between 0-10 days	1	54	Risk Ratio (M-H, Fixed, 95% CI)	0.65 [0.26, 1.61]
11.3 akathisia between 3-28 days (Oral phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	1.18 [0.62, 2.27]
11.4 dysarthria between 3-28 days (Oral phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	0.14 [0.01, 2.44]
11.5 dyskinesia between 0-10 days	1	54	Risk Ratio (M-H, Fixed, 95% CI)	0.29 [0.01, 6.79]
11.6 dystonia between 0-3 days (IM phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	0.94 [0.06, 13.93]
11.7 dystonia between 0-10 days	1	54	Risk Ratio (M-H, Fixed, 95% CI)	0.65 [0.26, 1.61]
11.8 involuntary movement between 3-28 days (Oral phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	0.32 [0.01, 7.26]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
11.9 motor retardation between 3-28 days (Oral phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	0.94 [0.06, 13.93]
11.10 muscle spasms between 0-3 days (IM phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	1.26 [0.33, 4.82]
11.11 oculogyric crisis between 0-10 days	1	54	Risk Ratio (M-H, Fixed, 95% CI)	2.6 [0.11, 61.11]
11.12 rigidity between 0-3 days (IM phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	0.88 [0.65, 1.19]
11.13 rigidity between 0-10 days	1	54	Risk Ratio (M-H, Fixed, 95% CI)	0.86 [0.13, 5.68]
11.14 rigidity during between 3-28 days (Oral phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	1.13 [0.93, 1.38]
11.15 tremor between 0-3 days (IM phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	0.94 [0.42, 2.13]
11.16 tremor during between 3-28 days (Oral phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	1.12 [0.71, 1.76]
11.17 tremor between 0-10 days	1	54	Risk Ratio (M-H, Fixed, 95% CI)	2.6 [0.11, 61.11]
11.18 use of antiparkinson medication	1	35	Risk Ratio (M-H, Fixed, 95% CI)	2.83 [0.66, 12.16]
11.19 dystonia during 72 hours	1	30	Risk Ratio (M-H, Fixed, 95% CI)	5.0 [0.26, 96.13]
11.20 extrapyramidal effects - use of antiparkinson drugs during 72 hours	1	30	Risk Ratio (M-H, Fixed, 95% CI)	0.6 [0.17, 2.07]
11.21 EPS during 72 hours	1	30	Risk Ratio (M-H, Fixed, 95% CI)	7.0 [0.98, 50.16]
11.22 thick tongue - between 0-10 days	1	54	Risk Ratio (M-H, Fixed, 95% CI)	0.29 [0.01, 6.79]
12 Adverse effects: 2e. Specific - miscellaneous	3		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
12.1 allergy - itch - between 3-28 days (Oral phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	2.84 [0.12, 65.34]
12.2 allergy - tissue reaction at injection site - between 0-10 days	1	54	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
12.3 nervousness - between 0-10 days	1	54	Risk Ratio (M-H, Fixed, 95% CI)	0.29 [0.01, 6.79]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
12.4 pain - during 24 hours	1	30	Risk Ratio (M-H, Fixed, 95% CI)	0.67 [0.13, 3.44]
12.5 pain - headache - between 0-3 days (IM phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	0.32 [0.01, 7.26]
12.6 pain - myalgia - between 3-28 days (Oral phase)	1	35	Risk Ratio (M-H, Fixed, 95% CI)	2.84 [0.12, 65.34]
13 Leaving the study early: 1. For general reasons	2		Risk Ratio (M-H, Fixed, 90% CI)	Subtotals only
13.1 by 72 hours	1	35	Risk Ratio (M-H, Fixed, 90% CI)	0.63 [0.21, 1.85]
13.2 by 10 days	1	54	Risk Ratio (M-H, Fixed, 90% CI)	1.18 [0.67, 2.07]
13.3 by endpoint	1	35	Risk Ratio (M-H, Fixed, 90% CI)	0.94 [0.42, 2.13]
14 Leaving the study early: 2. Specific reasons	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
14.1 adverse effects by 72 hours	1	35	Risk Ratio (M-H, Fixed, 95% CI)	2.84 [0.12, 65.34]

Analysis 5.1. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 1 Tranquillisation or asleep - not asleep up to 24 hours.

Analysis 5.2. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 2 Specific behaviour: 2. Aggression - various measures.

Analysis 5.3. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 3 Global outcome: 1. General - no change - i. over 24 hours.

Analysis 5.4. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 4 Global outcome: 2a. Specific - not sedated - i. by 30 minutes.

Analysis 5.5. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 5 Global outcome: 2b. Specific - not sedated - ii. up to 2 hours.

Analysis 5.6. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 6 Mental state: 1. Average scores.

Analysis 5.7. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 7 Adverse effects: 1. General.

**Analysis 5.8. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC:
d. LOXAPINE, Outcome 8 Adverse effects: 2a. Specific - anticholinergic.**

Analysis 5.9. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 9 Adverse effects: 2b. Specific - arousal level.

Analysis 5.10. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 10 Adverse effects: 2c. Specific - cardiovascular.

Analysis 5.11. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 11 Adverse effects: 2d. Specific - movement disorders.

Analysis 5.12. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 12 Adverse effects: 2e. Specific - miscellaneous.

Analysis 5.13. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: d. LOXAPINE, Outcome 13 Leaving the study early: 1. For general reasons.

**Analysis 5.14. Comparison 5 HALOPERIDOL vs OTHER ANTIPSYCHOTIC:
d. LOXAPINE, Outcome 14 Leaving the study early: 2. Specific reasons.**

Comparison 6. HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Tranquillisation or asleep - asleep	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
1.1 not asleep up to 2 hours	1	257	Risk Ratio (M-H, Fixed, 95% CI)	1.16 [1.02, 1.32]
2 Repeated need for tranquillisation - needing additional injection	4		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
2.1 by 2 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	4.0 [0.47, 33.73]
2.2 by 12 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	3.29 [1.67, 6.47]
2.3 by 24 hours	3	392	Risk Ratio (M-H, Fixed, 95% CI)	1.06 [0.75, 1.51]
3 Specific behaviour: 1a. Agitation - various measures	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
3.1 response - up to 2 hours (≥40% reduction in PANSS-EC)	1	45	Risk Ratio (M-H, Fixed, 95% CI)	0.96 [0.58, 1.58]
3.2 agitation - reported as 'adverse effect'	1	49	Risk Ratio (M-H, Fixed, 95% CI)	1.04 [0.07, 15.73]
3.3 agitation during 21 days (if occurred in ≥10% at P < 0.05) - reported as 'adverse effect'	1	100	Risk Ratio (M-H, Fixed, 95% CI)	1.08 [0.33, 3.51]
4 Specific behaviour: 1b. Agitation - average scores - i. by 30 minutes	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
4.1 change score at 15 minutes (ACES, low=agitated, high=sedated)	1	46	Mean Difference (IV, Fixed, 95% CI)	0.5 [-0.35, 1.35]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
4.2 change scores at 15 minutes (PANSS-EC, high=worse)	1	46	Mean Difference (IV, Fixed, 95% CI)	0.30 [-1.64, 2.24]
4.3 change score at 30 minutes (ACES, low=agitated, high=sedated)	1	46	Mean Difference (IV, Fixed, 95% CI)	0.90 [-0.06, 1.86]
4.4 change scores at 30 minutes (PANSS-EC, high=worse)	1	46	Mean Difference (IV, Fixed, 95% CI)	1.30 [-1.16, 3.76]
5 Specific behaviour: 1c. Agitation - average scores - ii. up to 2 hours	5		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
5.1 change score (ABS, high=worse)	1	85	Mean Difference (IV, Fixed, 95% CI)	2.7 [0.38, 5.02]
5.2 change score at 1 hour (ACES, low=agitated, high=sedated)	1	46	Mean Difference (IV, Fixed, 95% CI)	1.2 [0.06, 2.34]
5.3 endpoint score at 1 hour (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	2.00 [1.18, 2.82]
5.4 change score at 1 hour (PANSS-EC, high=worse)	1	46	Mean Difference (IV, Fixed, 95% CI)	2.2 [-0.50, 4.90]
5.5 change score at 2 hours (ACES, low=agitated, high=sedated)	1	46	Mean Difference (IV, Fixed, 95% CI)	0.30 [-0.74, 1.34]
5.6 endpoint score at 2 hours (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	4.20 [3.54, 4.86]
5.7 score at 2 hours (PANSS-EC, high=worse)	3	332	Mean Difference (IV, Fixed, 95% CI)	-0.31 [-1.49, 0.86]
5.8 endpoint score at 2 hours (PANSS-PAS, high=worse)	1	29	Mean Difference (IV, Fixed, 95% CI)	2.10 [-2.56, 6.76]
6 Specific behaviour: 1d. Agitation - average scores - iii. up to 4 hours	1	60	Mean Difference (IV, Fixed, 95% CI)	0.20 [-0.05, 0.45]
6.1 endpoint score at 4 hours (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.20 [-0.05, 0.45]
7 Specific behaviour: 1e. Agitation - average scores - iv. up to 24 hours	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
7.1 change score (ABS, high=worse)	1	86	Mean Difference (IV, Fixed, 95% CI)	2.40 [0.01, 4.79]
7.2 change score (PANSS-EC, high=worse)	1	86	Mean Difference (IV, Fixed, 95% CI)	1.40 [-0.55, 3.35]
7.3 endpoint score at 6 hours (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.20 [0.05, 0.35]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
7.4 endpoint score at 12 hours (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.9 [0.82, 0.98]
8 Specific behaviour: 2a. Aggression - average scores - i. up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
8.1 endpoint score at 1 hour (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.90 [0.13, 1.67]
8.2 endpoint score at 2 hours (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.5 [-0.41, 1.41]
9 Specific behaviour: 2b. Aggression - average scores - ii. up to 4 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
9.1 endpoint score at 4 hours (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.10 [-0.11, 0.31]
10 Specific behaviour: 2c. Aggression - average scores - iii. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
10.1 endpoint score at 6 hours (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-0.10 [-0.26, 0.06]
10.2 endpoint score at 12 hours (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-0.20 [-0.39, -0.01]
11 Specific behaviour: 3. Hostility	1	49	Risk Ratio (M-H, Fixed, 95% CI)	0.35 [0.01, 8.12]
12 Global outcome: 1a. General - need for additional measures	4		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
12.1 need for benzodiazepine during 24 hours	2	343	Risk Ratio (M-H, Fixed, 95% CI)	1.05 [0.63, 1.74]
12.2 need for benzodiazepine during 7 days	1	100	Risk Ratio (M-H, Fixed, 95% CI)	1.00 [0.79, 1.27]
12.3 additional restraint, seclusion or special observation	2	160	Risk Ratio (M-H, Fixed, 95% CI)	1.49 [0.70, 3.17]
13 Global outcome: 1b. General - time and doses	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
13.1 time to discontinuation	1	100	Mean Difference (IV, Fixed, 95% CI)	-3.48 [-6.28, -0.68]
13.2 dose of adjunctive lorazepam	1	100	Mean Difference (IV, Fixed, 95% CI)	0.34 [-0.14, 0.82]
14 Global outcome: 1c. General - average scores - up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
14.1 endpoint score (CGI-I, high=worse)	1	42	Mean Difference (IV, Fixed, 95% CI)	-0.10 [-0.65, 0.45]
15 Global outcome: 1d. General - average scores - over 24 hours	3		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
15.1 change score (CGI-I, high=worse)	1	243	Mean Difference (IV, Fixed, 95% CI)	0.0 [-0.20, 0.20]
15.2 change score (CGI-S, high=worse)	1	86	Mean Difference (IV, Fixed, 95% CI)	0.0 [-0.24, 0.24]
15.3 change score by 21 days (CGI-I, high=worse)	1	100	Mean Difference (IV, Fixed, 95% CI)	0.36 [-0.18, 0.90]
16 Global outcome: 2a. Specific - alert - i. up to 2 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
16.1 alert at 1 hour	1	100	Risk Ratio (M-H, Fixed, 95% CI)	1.24 [0.99, 1.55]
17 Global outcome: 2b. Specific - alert - ii. up to 4 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
17.1 alert at 4 hours	1	100	Risk Ratio (M-H, Fixed, 95% CI)	0.96 [0.66, 1.42]
18 Global outcome: 2c. Specific - alert - iii. up to 24 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
18.1 alert at 8 hours	1	100	Risk Ratio (M-H, Fixed, 95% CI)	1.04 [0.67, 1.60]
18.2 alert at 16 hours	1	100	Risk Ratio (M-H, Fixed, 95% CI)	1.02 [0.58, 1.78]
18.3 alert at 24 hours	1	100	Risk Ratio (M-H, Fixed, 95% CI)	0.85 [0.60, 1.20]
19 Global outcome: 2d. Specific - tranquil - i. up to 2 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
19.1 tranquil at 1 hour	1	100	Risk Ratio (M-H, Fixed, 95% CI)	0.62 [0.29, 1.34]
20 Global outcome: 2e. Specific - tranquil - ii. up to 4 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
20.1 tranquil at 4 hours	1	100	Risk Ratio (M-H, Fixed, 95% CI)	1.84 [0.94, 3.62]
21 Global outcome: 2f. Specific - tranquil - iii. up to 24 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
21.1 tranquil at 8 hours	1	100	Risk Ratio (M-H, Fixed, 95% CI)	1.01 [0.55, 1.87]
21.2 tranquil at 16 hours	1	100	Risk Ratio (M-H, Fixed, 95% CI)	0.72 [0.36, 1.45]
21.3 tranquil at 24 hours	1	100	Risk Ratio (M-H, Fixed, 95% CI)	1.50 [0.83, 2.72]
22 Global outcome: 2g. Specific - sedated - i. up to 2 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
22.1 sedated at 1 hour	1	100	Risk Ratio (M-H, Fixed, 95% CI)	0.22 [0.01, 4.39]
23 Global outcome: 2h. Specific - sedated - ii. up to 4 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
23.1 sedated at 4 hours	1	100	Risk Ratio (M-H, Fixed, 95% CI)	0.43 [0.18, 1.03]
24 Global outcome: 2i. Specific - sedated - iii. up to 24 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
24.1 sedated at 8 hours	1	100	Risk Ratio (M-H, Fixed, 95% CI)	1.08 [0.49, 2.37]
24.2 sedated at 16 hours	1	100	Risk Ratio (M-H, Fixed, 95% CI)	0.84 [0.47, 1.50]
24.3 sedated at 24 hours	1	100	Risk Ratio (M-H, Fixed, 95% CI)	0.62 [0.19, 1.98]
25 Global outcome: 2j. Specific - asleep - i. up to 2 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
25.1 asleep at 1 hour	1	100	Risk Ratio (M-H, Fixed, 95% CI)	0.36 [0.02, 8.64]
26 Global outcome: 2k. Specific - asleep - ii. up to 4 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
26.1 asleep at 4 hours	1	100	Risk Ratio (M-H, Fixed, 95% CI)	1.08 [0.07, 16.84]
27 Global outcome: 2l. Specific - asleep - iii. up to 24 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
27.1 asleep at 8 hours	1	100	Risk Ratio (M-H, Fixed, 95% CI)	0.27 [0.03, 2.34]
27.2 asleep at 16 hours	1	100	Risk Ratio (M-H, Fixed, 95% CI)	4.33 [0.97, 19.40]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
27.3 asleep at 24 hours	1	100	Risk Ratio (M-H, Fixed, 95% CI)	3.24 [0.14, 77.79]
28 Service use: 1. Average days to discharge	1	100	Mean Difference (IV, Fixed, 95% CI)	-0.60 [-1.85, 0.65]
29 Service use: 2. Average patient-hours used during 24 hours (skewed data)			Other data	No numeric data
29.1 days 1-7			Other data	No numeric data
29.2 days 8-14			Other data	No numeric data
29.3 days 15-21			Other data	No numeric data
30 Mental state: 1. Various outcomes reported as 'adverse events'	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
30.1 anxiety	1	49	Risk Ratio (M-H, Fixed, 95% CI)	0.35 [0.01, 8.12]
30.2 anxiety during 21 days (if occurred in $\geq 10\%$ at $P < 0.05$)	1	100	Risk Ratio (M-H, Fixed, 95% CI)	0.36 [0.08, 1.70]
30.3 delusions	1	49	Risk Ratio (M-H, Fixed, 95% CI)	0.35 [0.01, 8.12]
30.4 nervousness during 21 days (if occurred in $\geq 10\%$ at $P < 0.05$)	1	100	Risk Ratio (M-H, Fixed, 95% CI)	2.17 [0.70, 6.74]
31 Mental state: 2a. Average scores - i. by 30 minutes	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
31.1 change score at 15 minutes (BPRS positive sub-scale, high=worse)	1	46	Mean Difference (IV, Fixed, 95% CI)	-0.30 [-2.11, 1.51]
31.2 change score at 30 minutes (BPRS positive sub-scale, high=worse)	1	46	Mean Difference (IV, Fixed, 95% CI)	-0.40 [-2.70, 1.90]
31.3 change score at 15 minutes (BPRS total, high=worse)	1	46	Mean Difference (IV, Fixed, 95% CI)	-0.70 [-5.42, 4.02]
31.4 change score at 30 minutes (BPRS total, high=worse)	1	46	Mean Difference (IV, Fixed, 95% CI)	-0.30 [-7.35, 6.75]
32 Mental state: 2b. Average scores - ii. up to 2 hours	3		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
32.1 change score at 1 hour (BPRS positive sub-scale, high=worse)	1	46	Mean Difference (IV, Fixed, 95% CI)	0.60 [-2.74, 3.94]
32.2 change score at 2 hours (BPRS positive sub-scale, high=worse)	3	385	Mean Difference (IV, Fixed, 95% CI)	0.28 [-0.34, 0.89]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
32.3 change score at 1 hour (BPRS total, high=worse)	1	46	Mean Difference (IV, Fixed, 95% CI)	4.70 [-5.03, 14.43]
32.4 change score at 2 hours (BPRS total, high=worse)	3	385	Mean Difference (IV, Fixed, 95% CI)	1.75 [-0.12, 3.62]
33 Mental state: 2c. Average scores - iii. up to 24 hours	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
33.1 change score at 24 hours (BPRS positive sub-scale, high=worse)	2	340	Mean Difference (IV, Fixed, 95% CI)	-0.30 [-0.97, 0.37]
33.2 change score at 24 hours (BPRS total, high=worse)	2	340	Mean Difference (IV, Fixed, 95% CI)	0.47 [-1.34, 2.29]
34 Mental state: 2d. Average scores - iv. over 24 hours	1	100	Mean Difference (IV, Fixed, 95% CI)	4.70 [-0.18, 9.58]
34.1 change score at 21 days (PANSS total, high=worse)	1	100	Mean Difference (IV, Fixed, 95% CI)	4.70 [-0.18, 9.58]
35 Adverse effects: 1a. General	3		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
35.1 one or more drug related adverse events	3	209	Risk Ratio (M-H, Fixed, 95% CI)	1.38 [1.09, 1.76]
36 Adverse effects: 1b. General - serious	3		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
36.1 one or more treatment emergent adverse events	1	49	Risk Ratio (M-H, Fixed, 95% CI)	0.81 [0.36, 1.83]
36.2 treatment emergent adverse events - all	1	49	Risk Ratio (M-H, Fixed, 95% CI)	1.23 [0.69, 2.19]
36.3 overall serious adverse effects	1	49	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
36.4 death	3	138	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
37 Adverse effects: 2a. Specific - anti-cholinergic	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
37.1 increased salivation during 21 days (if occurred in $\geq 10\%$ at $P < 0.05$)	1	100	Risk Ratio (M-H, Fixed, 95% CI)	9.73 [0.54, 176.18]
38 Adverse effects: 2b. Specific - arousal level - i. various	4		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
38.1 insomnia	1	49	Risk Ratio (M-H, Fixed, 95% CI)	1.04 [0.39, 2.78]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
38.2 insomnia during 21 days (if occurred in $\geq 10\%$ at $P < 0.05$)	1	100	Risk Ratio (M-H, Fixed, 95% CI)	2.17 [0.57, 8.19]
38.3 somnolence at 24 hours	1	257	Risk Ratio (M-H, Fixed, 95% CI)	1.04 [0.45, 2.41]
38.4 somnolence during 21 days (if occurred in $\geq 10\%$ at $P < 0.05$)	1	100	Risk Ratio (M-H, Fixed, 95% CI)	1.44 [0.67, 3.12]
38.5 excessive sedation during 24 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	3.0 [0.33, 27.23]
39 Adverse effects: 2c. Specific - arousal level - ii. average scores - i. up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
39.1 endpoint score at 1 hour (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.30 [-0.13, 0.73]
39.2 endpoint score at 2 hours (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.0 [-0.36, 0.36]
40 Adverse effects: 2d. Specific - arousal level - ii. average scores - ii. up to 4 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
40.1 endpoint score at 4 hours (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-0.20 [-0.61, 0.21]
41 Adverse effects: 2e. Specific - arousal level - ii. average scores - iii. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
41.1 endpoint score at 6 hours (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-0.10 [-0.38, 0.18]
41.2 endpoint score at 12 hours (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.0 [-0.26, 0.26]
42 Adverse effects: 2f. Specific - cardiovascular i. binary	3	195	Risk Ratio (M-H, Fixed, 95% CI)	0.27 [0.03, 2.38]
42.1 clinically significant ECG change	1	49	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
42.2 hypotension during 24 hours	2	146	Risk Ratio (M-H, Fixed, 95% CI)	0.27 [0.03, 2.38]
43 Adverse effects: 2g. Specific - cardiovascular ii. continuous	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
43.1 QT interval: average change at 24 hours	1	257	Mean Difference (IV, Fixed, 95% CI)	1.8 [-3.81, 7.41]
43.2 QT interval: average endpoint score at 24 hours	1	86	Mean Difference (IV, Fixed, 95% CI)	8.5 [-3.28, 20.28]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
44 Adverse effects: 2h. Specific - gas-trointestinal	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
44.1 abdominal pain	1	49	Risk Ratio (M-H, Fixed, 95% CI)	3.12 [0.13, 73.04]
44.2 vomiting	1	49	Risk Ratio (M-H, Fixed, 95% CI)	5.2 [0.26, 103.03]
45 Adverse effects: 2i. Specific - movement disorder - i. various	5		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
45.1 dystonia during 24 hours	2	343	Risk Ratio (M-H, Fixed, 95% CI)	12.92 [1.67, 99.78]
45.2 dystonia during 21 days (if occurred in $\geq 10\%$ at $P < 0.05$)	1	100	Risk Ratio (M-H, Fixed, 95% CI)	9.73 [0.54, 176.18]
45.3 hypertonia during 21 days (if occurred in $\geq 10\%$ at $P < 0.05$)	1	100	Risk Ratio (M-H, Fixed, 95% CI)	9.73 [0.54, 176.18]
45.4 EPS during 24 hours	3	403	Risk Ratio (M-H, Fixed, 95% CI)	8.35 [2.27, 30.63]
45.5 EPS during study	1	49	Risk Ratio (M-H, Fixed, 95% CI)	1.04 [0.07, 15.73]
45.6 extrapyramidal effects - use of antiparkinson drugs during 24 hours	1	257	Risk Ratio (M-H, Fixed, 95% CI)	4.51 [1.92, 10.58]
45.7 extrapyramidal effects - use of antiparkinson drugs during 21 days	1	100	Risk Ratio (M-H, Fixed, 95% CI)	9.73 [0.54, 176.18]
45.8 extrapyramidal effects - use of antiparkinson drugs during study	1	49	Risk Ratio (M-H, Fixed, 95% CI)	1.04 [0.07, 15.73]
46 Adverse effects: 2j. Specific - movement disorder - ii. average scores - i. up to 24 hours	1	483	Mean Difference (IV, Fixed, 95% CI)	0.34 [0.16, 0.53]
46.1 change score at 24 hours (SAS, high=worse)	1	242	Mean Difference (IV, Fixed, 95% CI)	1.31 [0.56, 2.06]
46.2 change score at 24 hours (BAS, high=worse)	1	241	Mean Difference (IV, Fixed, 95% CI)	0.28 [0.09, 0.47]
47 Adverse effects: 2k. Specific - movement disorder - iii. average scores - i. up to 24 hours			Other data	No numeric data
47.1 endpoint score at 24 hours (SAS, high=worse) - (skewed data)			Other data	No numeric data
47.2 endpoint score at 24 hours (BAS, high=worse) - (skewed data)			Other data	No numeric data

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
48 Adverse effects: 2l. Specific - movement disorder - vi. average scores - i. over 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
48.1 endpoint score at 96 hours (AIMS, high=worse)	1	29	Mean Difference (IV, Fixed, 95% CI)	0.20 [-0.87, 1.27]
48.2 endpoint score at 96 hours (BARS, high=worse)	1	29	Mean Difference (IV, Fixed, 95% CI)	0.8 [-0.01, 1.61]
48.3 endpoint score at 96 hours (SAS, high=worse)	1	29	Mean Difference (IV, Fixed, 95% CI)	2.2 [-0.07, 4.47]
49 Adverse effects: 2m. Specific - miscellaneous	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
49.1 pain	1	49	Risk Ratio (M-H, Fixed, 95% CI)	3.12 [0.13, 73.04]
49.2 pain during 21 days (if occurred in $\geq 10\%$ at $P < 0.05$)	1	100	Risk Ratio (M-H, Fixed, 95% CI)	1.08 [0.33, 3.51]
49.3 pain - headache during 21 days (if occurred in $\geq 10\%$ at $P < 0.05$)	1	100	Risk Ratio (M-H, Fixed, 95% CI)	2.17 [0.88, 5.32]
49.4 other - nose bleed	1	49	Risk Ratio (M-H, Fixed, 95% CI)	3.12 [0.13, 73.04]
49.5 other - clinically relevant laboratory changes	2	149	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
50 Leaving the study early	3		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
50.1 any reason	2	148	Risk Ratio (M-H, Fixed, 95% CI)	1.66 [1.04, 2.65]
50.2 adverse effects	1	100	Risk Ratio (M-H, Fixed, 95% CI)	8.67 [1.13, 66.75]
50.3 lack of efficacy	2	129	Risk Ratio (M-H, Fixed, 95% CI)	0.36 [0.09, 1.44]
50.4 lost at follow-up	1	100	Risk Ratio (M-H, Fixed, 95% CI)	0.54 [0.10, 2.82]
50.5 participants decision	1	100	Risk Ratio (M-H, Fixed, 95% CI)	2.17 [0.57, 8.19]
50.6 non compliance	1	100	Risk Ratio (M-H, Fixed, 95% CI)	2.17 [0.42, 11.30]
50.7 physician decision	1	100	Risk Ratio (M-H, Fixed, 95% CI)	4.33 [0.50, 37.42]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
50.8 sponsor decision	1	100	Risk Ratio (M-H, Fixed, 95% CI)	0.36 [0.02, 8.64]

Analysis 6.1. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 1 Tranquillisation or asleep - asleep.

Analysis 6.2. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 2 Repeated need for tranquillisation - needing additional injection.

Analysis 6.3. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 3 Specific behaviour: 1a. Agitation - various measures.

Analysis 6.4. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 4 Specific behaviour: 1b. Agitation - average scores - i. by 30 minutes.

Analysis 6.5. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 5 Specific behaviour: 1c. Agitation - average scores - ii. up to 2 hours.

Analysis 6.6. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 6 Specific behaviour: 1d. Agitation - average scores - iii. up to 4 hours.

Analysis 6.7. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 7 Specific behaviour: 1e. Agitation - average scores - iv. up to 24 hours.

**Analysis 6.8. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE,
Outcome 8 Specific behaviour: 2a. Aggression - average scores - i. up to 2 hours.**

**Analysis 6.9. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE,
Outcome 9 Specific behaviour: 2b. Aggression - average scores - ii. up to 4 hours.**

Analysis 6.10. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 10 Specific behaviour: 2c. Aggression - average scores - iii. up to 24 hours.

Analysis 6.11. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 11 Specific behaviour: 3. Hostility.

Analysis 6.12. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 12 Global outcome: 1a. General - need for additional measures.

Analysis 6.13. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 13 Global outcome: 1b. General - time and doses.

Analysis 6.14. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 14 Global outcome: 1c. General - average scores - up to 2 hours.

**Analysis 6.15. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e.
OLANZAPINE, Outcome 15 Global outcome: 1d. General - average scores - over 24 hours.**

**Analysis 6.16. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e.
OLANZAPINE, Outcome 16 Global outcome: 2a. Specific - alert - i. up to 2 hours.**

**Analysis 6.17. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e.
OLANZAPINE, Outcome 17 Global outcome: 2b. Specific - alert - ii. up to 4 hours.**

**Analysis 6.18. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e.
OLANZAPINE, Outcome 18 Global outcome: 2c. Specific - alert - iii. up to 24 hours.**

**Analysis 6.19. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e.
OLANZAPINE, Outcome 19 Global outcome: 2d. Specific - tranquil - i. up to 2 hours.**

**Analysis 6.20. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e.
OLANZAPINE, Outcome 20 Global outcome: 2e. Specific - tranquil - ii. up to 4 hours.**

**Analysis 6.21. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e.
OLANZAPINE, Outcome 21 Global outcome: 2f. Specific - tranquil - iii. up to 24 hours.**

**Analysis 6.22. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e.
OLANZAPINE, Outcome 22 Global outcome: 2g. Specific - sedated - i. up to 2 hours.**

Analysis 6.23. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 23 Global outcome: 2h. Specific - sedated - ii. up to 4 hours.

Analysis 6.24. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 24 Global outcome: 2i. Specific - sedated - iii. up to 24 hours.

Analysis 6.25. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 25 Global outcome: 2j. Specific - asleep - i. up to 2 hours.

**Analysis 6.26. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e.
OLANZAPINE, Outcome 26 Global outcome: 2k. Specific - asleep - ii. up to 4 hours.**

**Analysis 6.27. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e.
OLANZAPINE, Outcome 27 Global outcome: 2l. Specific - asleep - iii. up to 24 hours.**

Analysis 6.28. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 28 Service use: 1. Average days to discharge.

Analysis 6.29. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 29 Service use: 2. Average patient-hours used during 24 hours (skewed data).

Service use: 2. Average patient-hours used during 24 hours (skewed data)				
Study	Intervention	Mean	SD	N
days 1-7				
Kinon 2001d	Haloperidol	2.59	6.79	48
Kinon 2001d	Olanzapine	1.57	5.52	52
days 8-14				
Kinon 2001d	Haloperidol	0.92	4.05	48
Kinon 2001d	Olanzapine	0.33	2.23	52
days 15-21				
Kinon 2001d	Haloperidol	0.55	2.74	48
Kinon 2001d	Olanzapine	0	0	52

Analysis 6.30. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 30 Mental state: 1. Various outcomes reported as 'adverse events'.

**Analysis 6.31. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e.
OLANZAPINE, Outcome 31 Mental state: 2a. Average scores - i. by 30 minutes.**

**Analysis 6.32. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e.
OLANZAPINE, Outcome 32 Mental state: 2b. Average scores - ii. up to 2 hours.** **Analysis 6.33. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e.
OLANZAPINE, Outcome 33 Mental state: 2c. Average scores - iii. up to 24 hours.**

Study or subgroup	haloperidol		olanzapine		Mean Difference Fixed, 95% CI	Weight	Mean Difference Fixed, 95% CI				
	N	Mean(SD)	N	Mean(SD)							
Test for overall effect: Z=0.51(P=0.61)											
Test for subgroup differences: Chi²=0.62, df=1 (P=0.43), I²=0%											
					haloperidol	-100	-50	0	50	100	olanzapine

Analysis 6.34. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 34 Mental state: 2d. Average scores - iv. over 24 hours.

Study or subgroup	haloperidol		olanzapine		Mean Difference Fixed, 95% CI	Weight	Mean Difference Fixed, 95% CI				
	N	Mean(SD)	N	Mean(SD)							
6.34.1 change score at 21 days (PANSS total, high=worse)											
Kinon 2001d	48	-16 (13.8)	52	-20.7 (10.8)		100%	4.7[-0.18,9.58]				
Subtotal ***	48		52			100%	4.7[-0.18,9.58]				
Heterogeneity: Not applicable											
Test for overall effect: Z=1.89(P=0.06)											
Total ***	48		52			100%	4.7[-0.18,9.58]				
Heterogeneity: Not applicable											
Test for overall effect: Z=1.89(P=0.06)											
					haloperidol	-10	-5	0	5	10	olanzapine

Analysis 6.35. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 35 Adverse effects: 1a. General.

Study or subgroup	haloperidol	olanzapine	Risk Ratio		Weight	Risk Ratio
	n/N	n/N	M-H, Fixed, 95% CI			M-H, Fixed, 95% CI
6.35.1 one or more drug related adverse events						
Baldacara 2011	11/30	5/30			12.63%	2.2[0.87,5.57]
Eli Lilly 2004	1/24	1/25			2.47%	1.04[0.07,15.73]
Kinon 2001d	41/48	35/52			84.89%	1.27[1.02,1.59]
Subtotal (95% CI)	102	107			100%	1.38[1.09,1.76]
Total events: 53 (haloperidol), 41 (olanzapine)						
Heterogeneity: Tau ² =0; Chi ² =1.56, df=2(P=0.46); I ² =0%						
Test for overall effect: Z=2.62(P=0.01)						
			haloperidol	0.01 0.1 1 10 100	olanzapine	

Analysis 6.36. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 36 Adverse effects: 1b. General - serious.

Study or subgroup	haloperidol	olanzapine	Risk Ratio		Weight	Risk Ratio
	n/N	n/N	M-H, Fixed, 95% CI			M-H, Fixed, 95% CI
6.36.1 one or more treatment emergent adverse events						
Eli Lilly 2004	7/24	9/25			100%	0.81[0.36,1.83]
Subtotal (95% CI)	24	25			100%	0.81[0.36,1.83]
Total events: 7 (haloperidol), 9 (olanzapine)						
			haloperidol	0.5 0.7 1 1.5 2	olanzapine

Analysis 6.37. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 37 Adverse effects: 2a. Specific - anticholinergic.

Analysis 6.38. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 38 Adverse effects: 2b. Specific - arousal level - i. various.

Analysis 6.39. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 39 Adverse effects: 2c. Specific - arousal level - ii. average scores - i. up to 2 hours.

**Analysis 6.40. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE,
Outcome 40 Adverse effects: 2d. Specific - arousal level - ii. average scores - ii. up to 4 hours.**

**Analysis 6.41. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE,
Outcome 41 Adverse effects: 2e. Specific - arousal level - ii. average scores - iii. up to 24 hours.**

**Analysis 6.42. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e.
OLANZAPINE, Outcome 42 Adverse effects: 2f. Specific - cardiovascular i. binary.**

**Analysis 6.43. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e.
OLANZAPINE, Outcome 43 Adverse effects: 2g. Specific - cardiovascular ii. continuous.**

Analysis 6.44. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 44 Adverse effects: 2h. Specific - gastrointestinal.

Analysis 6.45. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 45 Adverse effects: 2i. Specific - movement disorder - i. various.

Analysis 6.46. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 46 Adverse effects: 2j. Specific - movement disorder - ii. average scores - i. up to 24 hours.

Analysis 6.47. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 47 Adverse effects: 2k. Specific - movement disorder - iii. average scores - i. up to 24 hours.

Adverse effects: 2k. Specific - movement disorder - iii. average scores - i. up to 24 hours					
Study	Intervention	Mean	SD	N	Notes
endpoint score at 24 hours (SAS, high=worse) - (skewed data)					
Eli Lilly 2004	Haloperidol	1.4	2.8	22	
Eli Lilly 2004	Olanzapine	2.2	3.5	20	
endpoint score at 24 hours (BAS, high=worse) - (skewed data)					
Eli Lilly 2004	Haloperidol	1.5	1.7	22	
Eli Lilly 2004	Olanzapine	1.7	2.6	20	

Analysis 6.48. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 48 Adverse effects: 2l. Specific - movement disorder - vi. average scores - i. over 24 hours.

Analysis 6.49. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 49 Adverse effects: 2m. Specific - miscellaneous.

Analysis 6.50. Comparison 6 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: e. OLANZAPINE, Outcome 50 Leaving the study early.

Comparison 7. HALOPERIDOL vs OTHER ANTIPSYCHOTIC: f. PERPHENAZINE

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Global outcome: No improvement	1	44	Risk Ratio (M-H, Fixed, 95% CI)	0.46 [0.04, 4.68]
2 Adverse effects: 1. General	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
2.1 one or more adverse effect	1	44	Risk Ratio (M-H, Fixed, 95% CI)	1.30 [0.61, 2.80]
2.2 clinically significant laboratory changes	1	44	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
3 Adverse effects: 2. Specific	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
3.1 hypotensive episode	1	44	Risk Ratio (M-H, Fixed, 95% CI)	0.31 [0.01, 7.12]
3.2 require antiparkinson medication	1	44	Risk Ratio (M-H, Fixed, 95% CI)	2.74 [0.62, 12.12]
3.3 discontinued due to EPS	1	44	Risk Ratio (M-H, Fixed, 95% CI)	1.83 [0.18, 18.70]
3.4 discontinued due to drowsiness/tension and no therapeutic effect	1	44	Risk Ratio (M-H, Fixed, 95% CI)	2.75 [0.12, 64.04]
4 Leaving the study early: 1. Specific reasons	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
4.1 discontinued due to EPS	1	44	Risk Ratio (M-H, Fixed, 95% CI)	1.83 [0.18, 18.70]
4.2 discontinued due to drowsiness/tension and no therapeutic effect	1	44	Risk Ratio (M-H, Fixed, 95% CI)	2.75 [0.12, 64.04]

**Analysis 7.1. Comparison 7 HALOPERIDOL vs OTHER ANTIPSYCHOTIC:
f. PERPHENAZINE, Outcome 1 Global outcome: No improvement.**

**Analysis 7.2. Comparison 7 HALOPERIDOL vs OTHER ANTIPSYCHOTIC:
f. PERPHENAZINE, Outcome 2 Adverse effects: 1. General.**

**Analysis 7.3. Comparison 7 HALOPERIDOL vs OTHER ANTIPSYCHOTIC:
f. PERPHENAZINE, Outcome 3 Adverse effects: 2. Specific.**

Analysis 7.4. Comparison 7 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: f. PERPHENAZINE, Outcome 4 Leaving the study early: 1. Specific reasons.

Comparison 8. HALOPERIDOL vs OTHER ANTIPSYCHOTIC: g. QUETIAPINE

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Specific behaviour: 1a. Agitation - average scores - i. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
1.1 change score at 24 hours (PANSS-EC, high=worse)	1	80	Mean Difference (IV, Fixed, 95% CI)	0.10 [-0.56, 0.76]
2 Specific behaviour: 1b. Agitation - average scores - ii. over 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
2.1 change score at 72 hours (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2.2 change score at 7 days (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2.3 change score at 10 days (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2.4 change score at 14 days (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2.5 change score at 28 days (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
3 Mental state: Average scores - i. over 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
3.1 change score at 28 days (PANSS total, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
3.2 change score at 28 days (PANSS positive sub-scale, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
3.3 change score at 28 days (PANSS negative sub-scale, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
4 Adverse effects: 1. General	1		Risk Ratio (M-H, Fixed, 95% CI)	Totals not selected
4.1 one or more drug related adverse events by 28 days	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
5 Adverse effects: 2. Specific - movement disorder - i. average scores	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
5.1 endpoint score at 28 days (SAS, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
6 Leaving the study early	1		Risk Ratio (M-H, Fixed, 95% CI)	Totals not selected
6.1 any reason	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
6.2 adverse event	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
6.3 lost at follow-up	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]

Analysis 8.1. Comparison 8 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: g. QUETIAPINE, Outcome 1 Specific behaviour: 1a. Agitation - average scores - i. up to 24 hours.

Analysis 8.2. Comparison 8 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: g. QUETIAPINE, Outcome 2 Specific behaviour: 1b. Agitation - average scores - ii. over 24 hours.

Analysis 8.3. Comparison 8 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: g. QUETIAPINE, Outcome 3 Mental state: Average scores - i. over 24 hours.

Analysis 8.4. Comparison 8 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: g. QUETIAPINE, Outcome 4 Adverse effects: 1. General.

Analysis 8.5. Comparison 8 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: g. QUETIAPINE, Outcome 5 Adverse effects: 2. Specific - movement disorder - i. average scores.

Analysis 8.6. Comparison 8 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: g. QUETIAPINE, Outcome 6 Leaving the study early.

Comparison 9. HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Tranquillisation or asleep - i. by 30 minutes	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
1.1 not asleep at 30 minutes	1	162	Risk Ratio (M-H, Fixed, 95% CI)	0.84 [0.74, 0.95]
2 Tranquillisation or asleep - ii. up to 2 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
2.1 not asleep at 1 hour	1	162	Risk Ratio (M-H, Fixed, 95% CI)	0.74 [0.59, 0.92]
2.2 not asleep at 2 hours	1	162	Risk Ratio (M-H, Fixed, 95% CI)	0.71 [0.51, 0.99]
3 Specific behaviour: 1a. Agitation - various measures	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
3.1 $\geq 50\%$ reduction in PANSS-EC score up to 24 hours	1	124	Risk Ratio (M-H, Fixed, 95% CI)	0.96 [0.79, 1.16]
3.2 discontinued due to severe agitation up to 24 hours	1	124	Risk Ratio (M-H, Fixed, 95% CI)	0.33 [0.01, 8.03]
4 Specific behaviour: 1b. Agitation - average scores - i. up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
4.1 endpoint score at 2 hours (PANSS-PAS, high=worse)	1	28	Mean Difference (IV, Fixed, 95% CI)	-0.40 [-5.22, 4.42]
5 Specific behaviour: 2a. Aggression - average scores - i. by 30 minutes	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
5.1 change score at 30 minutes (OAS total aggression, high=worse)	1	147	Mean Difference (IV, Fixed, 95% CI)	-0.5 [-0.58, -0.42]
6 Specific behaviour: 2b. Aggression - average scores - ii. up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
6.1 change score at 1 hour (OAS total aggression, high=worse)	1	146	Mean Difference (IV, Fixed, 95% CI)	-0.20 [-0.27, -0.13]
6.2 change score at 2 hours (OAS total aggression, high=worse)	1	145	Mean Difference (IV, Fixed, 95% CI)	-0.10 [-0.13, -0.07]
7 Global outcome: 1. Binary measures	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
7.1 needing additional benzodiazepine	2	286	Risk Ratio (M-H, Fixed, 95% CI)	0.98 [0.65, 1.47]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
7.2 rated as severe at 24 hours (CGI-S)	1	162	Risk Ratio (M-H, Fixed, 95% CI)	0.89 [0.51, 1.58]
8 Global outcome: 2. Continuous measures	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
8.1 dose of lorazepam	1	147	Mean Difference (IV, Fixed, 95% CI)	-0.10 [-0.49, 0.29]
8.2 time to additional dose	1	147	Mean Difference (IV, Fixed, 95% CI)	0.20 [-2.17, 2.57]
9 Adverse effects: 1a. General - one or more adverse effects	2	286	Risk Ratio (M-H, Fixed, 95% CI)	1.01 [0.84, 1.23]
9.1 up to 24 hours	2	286	Risk Ratio (M-H, Fixed, 95% CI)	1.01 [0.84, 1.23]
10 Adverse effects: 2a. Specific - arousal	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
10.1 sedated at 30 minutes	1	162	Risk Ratio (M-H, Fixed, 95% CI)	1.90 [1.39, 2.59]
10.2 sedated at 60 minutes	1	162	Risk Ratio (M-H, Fixed, 95% CI)	1.41 [1.15, 1.72]
10.3 sedated at 120 minutes	1	162	Risk Ratio (M-H, Fixed, 95% CI)	1.10 [0.98, 1.23]
10.4 somnolence during 24 hours (only reported if occurred in ≥5%)	1	162	Risk Ratio (M-H, Fixed, 95% CI)	0.96 [0.43, 2.12]
10.5 somnolence during 24 hours	1	124	Risk Ratio (M-H, Fixed, 95% CI)	0.78 [0.31, 1.96]
10.6 insomnia - severe	1	162	Risk Ratio (M-H, Fixed, 95% CI)	0.35 [0.01, 8.47]
10.7 somnolence - severe	1	162	Risk Ratio (M-H, Fixed, 95% CI)	3.15 [0.13, 76.20]
11 Adverse effects: 2b. Specific - cardiovascular - i. binary outcomes	2	286	Risk Ratio (M-H, Fixed, 95% CI)	1.46 [0.45, 4.70]
11.1 dizziness during 24 hours	1	124	Risk Ratio (M-H, Fixed, 95% CI)	1.0 [0.26, 3.82]
11.2 orthostatic hypertension	1	162	Risk Ratio (M-H, Fixed, 95% CI)	5.25 [0.26, 107.67]
12 Adverse effects: 2c. Specific - cardiovascular - ii. pulse rate - i. up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
12.1 heartbeat change at 60 minutes	1	162	Mean Difference (IV, Fixed, 95% CI)	-9.4 [-9.99, -8.81]
13 Adverse effects: 2d. Specific - cardiovascular - ii. pulse rate - ii. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
13.1 heartbeat change at 8 hours	1	162	Mean Difference (IV, Fixed, 95% CI)	-8.8 [-9.75, -7.85]
14 Adverse effects: 2e. Specific - movement disorder - i. binary outcomes	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
14.1 dystonia during 24 hours	2	286	Risk Ratio (M-H, Fixed, 95% CI)	1.44 [0.28, 7.28]
14.2 dyskinesia during 24 hours	1	162	Risk Ratio (M-H, Fixed, 95% CI)	3.15 [0.13, 76.20]
14.3 hyperkinesia during 24 hours (only reported if occurred in ≥5%)	1	162	Risk Ratio (M-H, Fixed, 95% CI)	4.20 [0.48, 36.79]
14.4 hypertonia/rigidity during 24 hours	1	162	Risk Ratio (M-H, Fixed, 95% CI)	0.35 [0.01, 8.47]
14.5 tremor during 24 hours	1	162	Risk Ratio (M-H, Fixed, 95% CI)	5.25 [0.26, 107.67]
14.6 movement disorder	1	124	Risk Ratio (M-H, Fixed, 95% CI)	1.6 [0.55, 4.62]
15 Adverse effects: 2f. Specific - movement disorders - ii. average scores - i. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
15.1 change score at 24 hours (BAS akathisia, high=worse)	1	162	Mean Difference (IV, Fixed, 95% CI)	0.3 [0.24, 0.36]
15.2 change score at 24 hours (SAS movement disorders, high=worse)	1	162	Mean Difference (IV, Fixed, 95% CI)	0.4 [0.34, 0.46]
16 Adverse effects: 2g. Specific - movement disorder - ii. average scores - ii. over 24 hours	1	84	Mean Difference (IV, Fixed, 95% CI)	0.58 [-0.32, 1.47]
16.1 endpoint score at 96 hours (AIMS, high=worse)	1	28	Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
16.2 endpoint score at 96 hours (BARS, high=worse)	1	28	Mean Difference (IV, Fixed, 95% CI)	0.60 [-0.36, 1.56]
16.3 endpoint score at 96 hours (SAS, high=worse)	1	28	Mean Difference (IV, Fixed, 95% CI)	0.40 [-2.20, 3.00]
17 Adverse effects: 2h. Specific - miscellaneous	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
17.1 agitation during 24 hours (only reported if occurred in $\geq 5\%$)	1	162	Risk Ratio (M-H, Fixed, 95% CI)	1.05 [0.27, 4.06]
17.2 agitation - severe	1	162	Risk Ratio (M-H, Fixed, 95% CI)	2.10 [0.19, 22.72]
17.3 anxiety - severe	1	162	Risk Ratio (M-H, Fixed, 95% CI)	3.15 [0.13, 76.20]
17.4 headache during 24 hours (only reported if occurred in $\geq 5\%$)	1	162	Risk Ratio (M-H, Fixed, 95% CI)	1.31 [0.37, 4.71]
17.5 headache during 24 hours	1	124	Risk Ratio (M-H, Fixed, 95% CI)	1.33 [0.31, 5.71]
18 Leaving the study early - i. up to 24 hours	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
18.1 due to severe agitation	1	124	Risk Ratio (M-H, Fixed, 95% CI)	0.33 [0.01, 8.03]
18.2 due to acute dystonia	1	124	Risk Ratio (M-H, Fixed, 95% CI)	5.0 [0.24, 102.07]
18.3 due to adverse effects	1	162	Risk Ratio (M-H, Fixed, 95% CI)	1.05 [0.07, 16.51]
18.4 unspecified general reasons	1	162	Risk Ratio (M-H, Fixed, 95% CI)	0.77 [0.42, 1.44]
19 Leaving the study early - ii. over 24 hours	1	28	Risk Ratio (M-H, Fixed, 95% CI)	0.11 [0.01, 1.89]
19.1 Lack of efficacy at 96 hours	1	28	Risk Ratio (M-H, Fixed, 95% CI)	0.11 [0.01, 1.89]

Analysis 9.1. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 1 Tranquillisation or asleep - i. by 30 minutes.

Analysis 9.2. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 2 Tranquillisation or asleep - ii. up to 2 hours.

Analysis 9.3. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 3 Specific behaviour: 1a. Agitation - various measures.

Analysis 9.4. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 4 Specific behaviour: 1b. Agitation - average scores - i. up to 2 hours.

Analysis 9.5. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 5 Specific behaviour: 2a. Aggression - average scores - i. by 30 minutes.

Analysis 9.6. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 6 Specific behaviour: 2b. Aggression - average scores - ii. up to 2 hours.

Analysis 9.7. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 7 Global outcome: 1. Binary measures.

Analysis 9.8. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 8 Global outcome: 2. Continuous measures.

Analysis 9.9. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 9 Adverse effects: 1a. General - one or more adverse effects.

Analysis 9.10. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 10 Adverse effects: 2a. Specific - arousal.

Analysis 9.11. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 11 Adverse effects: 2b. Specific - cardiovascular - i. binary outcomes.

Analysis 9.12. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 12 Adverse effects: 2c. Specific - cardiovascular - ii. pulse rate - i. up to 2 hours.

Analysis 9.13. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 13 Adverse effects: 2d. Specific - cardiovascular - ii. pulse rate - ii. up to 24 hours.

Analysis 9.14. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 14 Adverse effects: 2e. Specific - movement disorder - i. binary outcomes.

Analysis 9.15. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 15 Adverse effects: 2f. Specific - movement disorders - ii. average scores - i. up to 24 hours.

Analysis 9.16. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 16 Adverse effects: 2g. Specific - movement disorder - ii. average scores - ii. over 24 hours.

Analysis 9.17. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 17 Adverse effects: 2h. Specific - miscellaneous.

Analysis 9.18. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 18 Leaving the study early - i. up to 24 hours.

Analysis 9.19. Comparison 9 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: h. RISPERIDONE, Outcome 19 Leaving the study early - ii. over 24 hours.

Comparison 10. HALOPERIDOL vs OTHER ANTIPSYCHOTIC: i. THIOTHIXENE

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Repeated need for rapid tranquillisation	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
1.1 more than 1 injection	1	30	Risk Ratio (M-H, Fixed, 95% CI)	1.07 [0.89, 1.28]
1.2 more than 3 injections	1	30	Risk Ratio (M-H, Fixed, 95% CI)	2.5 [0.57, 10.93]
2 Specific behaviour: 1. Agitation - rated as 'adverse effect'	1	44	Risk Ratio (M-H, Fixed, 95% CI)	0.28 [0.01, 6.52]
3 Global outcome	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
3.1 no response to 1st injection	1	44	Risk Ratio (M-H, Fixed, 95% CI)	2.5 [0.57, 11.05]
3.2 no improvement at 1 hour	1	44	Risk Ratio (M-H, Fixed, 95% CI)	0.5 [0.14, 1.84]
3.3 no improvement at 2 hours	1	44	Risk Ratio (M-H, Fixed, 95% CI)	0.17 [0.01, 3.31]
3.4 no improvement at 3 hours	1	44	Risk Ratio (M-H, Fixed, 95% CI)	0.83 [0.06, 12.49]
3.5 no improvement at 4 hours	1	44	Risk Ratio (M-H, Fixed, 95% CI)	0.83 [0.06, 12.49]
3.6 no improvement at 5 hours	1	44	Risk Ratio (M-H, Fixed, 95% CI)	4.2 [0.21, 82.72]
4 Adverse effects: 1. General - one or more adverse effects	2	74	Risk Ratio (M-H, Fixed, 95% CI)	1.47 [0.97, 2.22]
5 Adverse effects: 2a. Specific - movement disorders	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
5.1 ataxia	1	44	Risk Ratio (M-H, Fixed, 95% CI)	0.83 [0.06, 12.49]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
5.2 thick tongue	1	44	Risk Ratio (M-H, Fixed, 95% CI)	2.52 [0.11, 58.67]
5.3 use of antiparkinson drugs	1	30	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
6 Adverse effects: 2b. Specific - others	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
6.1 anticholinergic - blurred vision	1	44	Risk Ratio (M-H, Fixed, 95% CI)	4.2 [0.21, 82.72]
6.2 anticholinergic - dry mouth	1	44	Risk Ratio (M-H, Fixed, 95% CI)	0.17 [0.01, 3.31]
6.3 arousal - drowsiness	2	74	Risk Ratio (M-H, Fixed, 95% CI)	1.72 [1.02, 2.90]
6.4 arousal - lightheadedness	1	44	Risk Ratio (M-H, Fixed, 95% CI)	4.2 [0.21, 82.72]
6.5 cardiac - chest pain	1	44	Risk Ratio (M-H, Fixed, 95% CI)	0.28 [0.01, 6.52]
6.6 cardiac - dizziness	1	44	Risk Ratio (M-H, Fixed, 95% CI)	2.5 [0.28, 22.20]
6.7 cardiac - hypotensive crisis	1	30	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
6.8 cardiac - orthostatic hypotension	1	44	Risk Ratio (M-H, Fixed, 95% CI)	0.83 [0.06, 12.49]
6.9 cardiac - tachycardia	1	44	Risk Ratio (M-H, Fixed, 95% CI)	0.28 [0.01, 6.52]
6.10 other - depressed mood	1	44	Risk Ratio (M-H, Fixed, 95% CI)	2.52 [0.11, 58.67]
6.11 other - diaphoresis	1	44	Risk Ratio (M-H, Fixed, 95% CI)	0.28 [0.01, 6.52]
6.12 other - weakness	2	74	Risk Ratio (M-H, Fixed, 95% CI)	2.75 [0.30, 25.21]
6.13 other - unsteadiness	1	30	Risk Ratio (M-H, Fixed, 95% CI)	3.0 [0.13, 68.26]

Analysis 10.1. Comparison 10 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: i. THIOTHIXENE, Outcome 1 Repeated need for rapid tranquillisation.

Analysis 10.2. Comparison 10 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: i. THIOTHIXENE, Outcome 2 Specific behaviour: 1. Agitation - rated as 'adverse effect'.

Analysis 10.3. Comparison 10 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: i. THIOTHIXENE, Outcome 3 Global outcome.

Analysis 10.4. Comparison 10 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: i. THIOTHIXENE, Outcome 4 Adverse effects: 1. General - one or more adverse effects.

Analysis 10.5. Comparison 10 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: i. THIOTHIXENE, Outcome 5 Adverse effects: 2a. Specific - movement disorders.

Analysis 10.6. Comparison 10 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: i. THIOTHIXENE, Outcome 6 Adverse effects: 2b. Specific - others.

Comparison 11. HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Repeated need for tranquillisation - need for additional drugs for tranquillisation	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
1.1 up to 2 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	0.36 [0.13, 1.01]
1.2 up to 24 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	1.64 [1.07, 2.53]
2 Specific behaviour: 1a. Agitation - average scores - i. up to 2 hours	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
2.1 endpoint score at 1 hour (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-7.70 [-9.41, -5.99]
2.2 endpoint score at 2 hours (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-3.90 [-6.38, -1.42]
2.3 endpoint score at 2 hours (PANSS-EC, high=worse)	1	231	Mean Difference (IV, Fixed, 95% CI)	0.06 [-1.13, 1.25]
3 Specific behaviour: 1b. Agitation - average scores - ii. up to 4 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
3.1 endpoint score at 4 hours (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.20 [-0.05, 0.45]
4 Specific behaviour: 1c. Agitation - average scores - iii. up to 24 hours	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
4.1 endpoint score at 6 hours (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.20 [0.05, 0.35]
4.2 endpoint score at 6 hours (PANSS-EC, high=worse)	1	231	Mean Difference (IV, Fixed, 95% CI)	-0.47 [-1.68, 0.74]
4.3 endpoint score at 12 hours (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.9 [0.82, 0.98]
4.4 endpoint score at 24 hours (PANSS-EC, high=worse)	1	231	Mean Difference (IV, Fixed, 95% CI)	-0.21 [-1.36, 0.94]
5 Specific behaviour: 1d. Agitation - average scores - iv. over 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
5.1 endpoint scores at 48 hours (PANSS-EC, high=worse)	1	231	Mean Difference (IV, Fixed, 95% CI)	0.06 [-1.03, 1.15]
5.2 endpoint scores at 72 hours (PANSS-EC, high=worse)	1	231	Mean Difference (IV, Fixed, 95% CI)	0.62 [-0.45, 1.69]
6 Specific behaviour: 2a. Aggression - average scores - i. up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
6.1 endpoint score at 1 hour (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.0 [-0.77, 0.77]
6.2 endpoint score at 2 hours (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.70 [-0.25, 1.65]
7 Specific behaviour: 2b. Aggression - average scores - ii. up to 4 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
7.1 endpoint score at 4 hours (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.30 [-0.04, 0.64]
8 Specific behaviour: 2c. Aggression - average scores - iii. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
8.1 endpoint score at 6 hours (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.70 [0.60, 0.80]
8.2 endpoint score at 12 hours (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.20 [0.07, 0.33]
9 Global outcome: 1. General - need for additional measures	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
9.1 need for anxiolytic during 7 days	1	132	Risk Ratio (M-H, Fixed, 95% CI)	1.11 [0.84, 1.48]
9.2 need for hypnotics for night time sedation during 7 days	1	132	Risk Ratio (M-H, Fixed, 95% CI)	0.71 [0.20, 2.50]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
9.3 additional restraint, seclusion or special observation during 12 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	0.6 [0.25, 1.44]
10 Global outcome: 2. Average scores - i. over 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
10.1 change score at 72 hours (CGI-S, high=worse)	1	132	Mean Difference (IV, Fixed, 95% CI)	0.34 [0.13, 0.55]
10.2 change score at 7 days (CGI-S, high=worse)	1	132	Mean Difference (IV, Fixed, 95% CI)	0.51 [0.07, 0.95]
11 Mental state: 1a. Average scores - i. up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
11.1 endpoint score at 2 hours (BPRS, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
12 Mental state: 1b. Average scores - ii. up to 4 hours	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
12.1 endpoint score at 4 hours (BPRS, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
13 Mental state: 1c. Average scores - iii. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
13.1 endpoint score at 24 hours (BPRS, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
14 Mental state: 1d. Average scores - iv. over 24 hours	4		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
14.1 endpoint score at 48 hours (BPRS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-2.05 [-6.45, 2.35]
14.2 endpoint score at 72 hours (PANSS total, high=worse)	1	231	Mean Difference (IV, Fixed, 95% CI)	2.45 [-2.19, 7.09]
14.3 endpoint score at 72 hours (PANSS positive sub-scale, high=worse)	1	231	Mean Difference (IV, Fixed, 95% CI)	1.11 [-0.45, 2.67]
14.4 endpoint score at 72 hours (PANSS negative sub-scale, high=worse)	1	231	Mean Difference (IV, Fixed, 95% CI)	-4.19 [-5.71, -2.67]
14.5 endpoint score at 72 hours (PANSS general pathology score, high=worse)	1	231	Mean Difference (IV, Fixed, 95% CI)	0.0 [-2.27, 2.27]
14.6 score at 72 hours (BPRS, high=worse)	3	511	Mean Difference (IV, Fixed, 95% CI)	-0.43 [-1.93, 1.07]
14.7 change score at 7 days (BPRS, high=worse)	1	132	Mean Difference (IV, Fixed, 95% CI)	2.93 [-0.81, 6.67]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
15 Adverse effects: 1a. General - i. up to 24 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
15.1 one or more drug related adverse effects - by 12 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	1.38 [0.64, 2.93]
16 Adverse effects: 1b. General - ii. over 24 hours	4		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
16.1 one or more drug related adverse effects - by 72 hours	4	799	Risk Ratio (M-H, Fixed, 95% CI)	1.74 [1.47, 2.06]
16.2 one or more drug related adverse effects - by 7 days	1	132	Risk Ratio (M-H, Fixed, 95% CI)	1.31 [0.93, 1.83]
17 Adverse effects: 1c. General - serious	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
17.1 severe adverse effect - by 72 hours	1	376	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
17.2 death	2	436	Risk Ratio (M-H, Fixed, 95% CI)	0.14 [0.01, 2.78]
18 Adverse effects: 2a. Specific - anti-cholinergic	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
18.1 blurred vision by 72 hours	1	231	Risk Ratio (M-H, Fixed, 95% CI)	3.97 [1.15, 13.68]
18.2 constipation by 72 hours	1	376	Risk Ratio (M-H, Fixed, 95% CI)	0.40 [0.08, 2.06]
18.3 dry mouth by 72 hours	1	231	Risk Ratio (M-H, Fixed, 95% CI)	2.97 [1.22, 7.22]
18.4 hypersalivation by 72 hours	1	231	Risk Ratio (M-H, Fixed, 95% CI)	2.55 [1.11, 5.87]
19 Adverse effects: 2b. Specific - arousal level - i. various	3		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
19.1 excessive sedation at 12 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	1.0 [0.22, 4.56]
19.2 insomnia - by 72 hours	1	376	Risk Ratio (M-H, Fixed, 95% CI)	0.51 [0.05, 5.53]
19.3 lethargy - by 72 hours	1	231	Risk Ratio (M-H, Fixed, 95% CI)	1.26 [0.67, 2.35]
19.4 somnolence - by 72 hours	1	376	Risk Ratio (M-H, Fixed, 95% CI)	1.16 [0.43, 3.12]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
20 Adverse effects: 2c. Specific - arousal level - ii. average scores - i. up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
20.1 endpoint score at 1 hour (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.0 [-0.46, 0.46]
20.2 endpoint score at 2 hours (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.0 [-0.46, 0.46]
21 Adverse effects: 2d. Specific - arousal level - ii. average scores - ii. up to 4 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
21.1 endpoint score at 4 hours (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-0.40 [-0.83, 0.03]
22 Adverse effects: 2e. Specific - arousal level - ii. average scores - iii. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
22.1 endpoint score at 6 hours (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-0.10 [-0.40, 0.20]
22.2 endpoint score at 12 hours (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-0.10 [-0.40, 0.20]
23 Adverse effects: 2f. Specific - cardiovascular	4		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
23.1 hypotension at 12 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	0.08 [0.00, 1.31]
23.2 dizziness - by 72 hours	2	607	Risk Ratio (M-H, Fixed, 95% CI)	0.58 [0.28, 1.19]
23.3 ECG - abnormal - by 72 hours	2	607	Risk Ratio (M-H, Fixed, 95% CI)	0.38 [0.17, 0.84]
23.4 ECG change - clinically significant - by 72 hours	1	376	Risk Ratio (M-H, Fixed, 95% CI)	1.01 [0.60, 1.71]
23.5 tachycardia - by 72 hours	3	739	Risk Ratio (M-H, Fixed, 95% CI)	0.49 [0.23, 1.04]
24 Adverse effects: 2g. Specific - gastrointestinal	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
24.1 nausea by 72 hours	1	376	Risk Ratio (M-H, Fixed, 95% CI)	0.40 [0.08, 2.06]
24.2 vomiting by 72 hours	1	132	Risk Ratio (M-H, Fixed, 95% CI)	0.30 [0.02, 5.72]
24.3 vomiting by 7 days	1	132	Risk Ratio (M-H, Fixed, 95% CI)	0.16 [0.01, 2.82]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
25 Adverse effects: 2h. Specific - hematological tests	3		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
25.1 abnormal lab results - by 72 hours	2	508	Risk Ratio (M-H, Fixed, 95% CI)	0.90 [0.69, 1.17]
25.2 abnormal lab results - by 7 days	1	132	Risk Ratio (M-H, Fixed, 95% CI)	1.01 [0.47, 2.15]
25.3 aspartate aminotransference - by 72 hours	1	376	Risk Ratio (M-H, Fixed, 95% CI)	11.12 [0.62, 199.64]
25.4 glucose increased - by 24 hours	1	132	Risk Ratio (M-H, Fixed, 95% CI)	0.95 [0.31, 2.92]
25.5 haemogram abnormal - by 72 hours	1	231	Risk Ratio (M-H, Fixed, 95% CI)	0.17 [0.02, 1.35]
25.6 lactate dehydrogenase increase - by 72 hours	1	231	Risk Ratio (M-H, Fixed, 95% CI)	0.50 [0.05, 5.39]
25.7 liver function abnormal - by 72 hours	1	231	Risk Ratio (M-H, Fixed, 95% CI)	0.50 [0.05, 5.39]
25.8 triglyceride increase - by 72 hours	1	231	Risk Ratio (M-H, Fixed, 95% CI)	4.96 [0.24, 102.14]
26 Adverse effects: 2i. Specific - movement disorders - i. binary measures	4		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
26.1 akathisia - by 72 hours	3	739	Risk Ratio (M-H, Fixed, 95% CI)	2.32 [1.34, 4.01]
26.2 akathisia - by 7 days	1	132	Risk Ratio (M-H, Fixed, 95% CI)	4.29 [1.13, 16.31]
26.3 dystonia - by 72 hours	2	508	Risk Ratio (M-H, Fixed, 95% CI)	10.26 [1.67, 63.17]
26.4 dystonia - by 7 days	1	132	Risk Ratio (M-H, Fixed, 95% CI)	2.68 [0.76, 9.47]
26.5 EPS - by 12 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	11.0 [0.64, 190.53]
26.6 EPS - by 72 hours	2	508	Risk Ratio (M-H, Fixed, 95% CI)	19.13 [7.59, 48.21]
26.7 EPS - by 7 days	1	132	Risk Ratio (M-H, Fixed, 95% CI)	34.29 [4.70, 250.02]
26.8 extrapyramidal effects - use of antiparkinson drugs - by 7 days	1	132	Risk Ratio (M-H, Fixed, 95% CI)	3.30 [1.82, 5.97]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
26.9 hypertonia/rigidity - by 72 hours	1	132	Risk Ratio (M-H, Fixed, 95% CI)	14.81 [0.78, 280.47]
26.10 hypertonia - by 7 days	1	132	Risk Ratio (M-H, Fixed, 95% CI)	3.57 [0.90, 14.25]
26.11 myotonia - by 72 hours	1	231	Risk Ratio (M-H, Fixed, 95% CI)	3.84 [1.85, 8.00]
26.12 pyramidal tract syndrome - by 72 hours	1	376	Risk Ratio (M-H, Fixed, 95% CI)	17.18 [1.00, 295.55]
26.13 reduced movement - by 72 hours	1	231	Risk Ratio (M-H, Fixed, 95% CI)	2.14 [1.17, 3.91]
26.14 torsional spasm - by 72 hours	1	231	Risk Ratio (M-H, Fixed, 95% CI)	3.30 [0.93, 11.70]
26.15 tremor - by 72 hours	2	363	Risk Ratio (M-H, Fixed, 95% CI)	2.64 [1.37, 5.11]
26.16 tremor - by 7 days	1	132	Risk Ratio (M-H, Fixed, 95% CI)	4.29 [0.82, 22.48]
27 Adverse effects: 2j. Specific - movement disorders - ii. average scores - i. over 24 hours	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
27.1 change score at 72 hours (BAS, high=worse)	1	131	Mean Difference (IV, Fixed, 95% CI)	0.47 [0.18, 0.76]
27.2 score at 72 hours (SAS, high=worse)	2	191	Mean Difference (IV, Fixed, 95% CI)	3.59 [2.15, 5.03]
27.3 change score at 7 days (BAS, high=worse)	1	131	Mean Difference (IV, Fixed, 95% CI)	0.9 [0.51, 1.29]
27.4 change score at 7 days (SAS, high=worse)	1	131	Mean Difference (IV, Fixed, 95% CI)	6.1 [3.91, 8.29]
28 Adverse effects: 2k. Specific - miscellaneous	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
28.1 physical examination significant changes - by 72 hours	1	376	Risk Ratio (M-H, Fixed, 95% CI)	27.29 [1.63, 455.72]
29 Leaving the study early: 1. For any reason - i. over 24 hours	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
29.1 by 72 hours	1	376	Risk Ratio (M-H, Fixed, 95% CI)	1.77 [0.53, 5.94]
29.2 by 7 days	1	132	Risk Ratio (M-H, Fixed, 95% CI)	2.14 [0.86, 5.32]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
30 Leaving the study early: 2. For specific reasons - i. over 24 hours	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
30.1 adverse events - by 72 hours	2	508	Risk Ratio (M-H, Fixed, 95% CI)	2.40 [0.45, 12.75]
30.2 adverse events - by 7 days	1	132	Risk Ratio (M-H, Fixed, 95% CI)	0.54 [0.06, 4.65]
30.3 withdrew consent - by 72 hours	1	376	Risk Ratio (M-H, Fixed, 95% CI)	5.05 [0.24, 104.55]
30.4 other - by 72 hours	1	376	Risk Ratio (M-H, Fixed, 95% CI)	0.20 [0.01, 4.18]

Analysis 11.1. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 1 Repeated need for tranquillisation - need for additional drugs for tranquillisation.

Analysis 11.2. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 2 Specific behaviour: 1a. Agitation - average scores - i. up to 2 hours.

Analysis 11.3. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 3 Specific behaviour: 1b. Agitation - average scores - ii. up to 4 hours.

Analysis 11.4. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 4 Specific behaviour: 1c. Agitation - average scores - iii. up to 24 hours.

**Analysis 11.5. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE,
Outcome 5 Specific behaviour: 1d. Agitation - average scores - iv. over 24 hours.**

**Analysis 11.6. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE,
Outcome 6 Specific behaviour: 2a. Aggression - average scores - i. up to 2 hours.**

Analysis 11.7. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 7 Specific behaviour: 2b. Aggression - average scores - ii. up to 4 hours.

Analysis 11.8. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 8 Specific behaviour: 2c. Aggression - average scores - iii. up to 24 hours.

Analysis 11.9. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 9 Global outcome: 1. General - need for additional measures.

Analysis 11.10. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 10 Global outcome: 2. Average scores - i. over 24 hours.

Analysis 11.11. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 11 Mental state: 1a. Average scores - i. up to 2 hours.

Analysis 11.12. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 12 Mental state: 1b. Average scores - ii. up to 4 hours.

Study or subgroup	haloperidol		ziprasidone		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
11.12.1 endpoint score at 4 hours (BPRS, high=worse)						
Yin 2012	30	34.8 (8.3)	30	39.4 (11.2)	+	-4.6[-9.6,0.4]
					haloperidol -100 -50 0 50 100 ziprasidone	

Analysis 11.13. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 13 Mental state: 1c. Average scores - iii. up to 24 hours.

Study or subgroup	haloperidol		ziprasidone		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
11.13.1 endpoint score at 24 hours (BPRS, high=worse)						
Yin 2012	30	33.9 (9)	30	37.9 (11)	+	-4[-9.09,1.09]
					haloperidol -100 -50 0 50 100 ziprasidone	

Analysis 11.14. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 14 Mental state: 1d. Average scores - iv. over 24 hours.

Study or subgroup	haloperidol		ziprasidone		Mean Difference	Weight	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
11.14.1 endpoint score at 48 hours (BPRS, high=worse)							
Yin 2012	30	31.4 (7.1)	30	33.4 (10.1)		100%	-2.05[-6.45,2.35]
Subtotal ***	30		30			100%	-2.05[-6.45,2.35]
Heterogeneity: Not applicable Test for overall effect: Z=0.91(P=0.36)							
11.14.2 endpoint score at 72 hours (PANSS total, high=worse)							
Li 2006	116	70.9 (17.4)	115	68.4 (18.5)		100%	2.45[-2.19,7.09]
Subtotal ***	116		115			100%	2.45[-2.19,7.09]
Heterogeneity: Not applicable Test for overall effect: Z=1.04(P=0.3)							
11.14.3 endpoint score at 72 hours (PANSS positive sub-scale, high=worse)							
Li 2006	116	19.2 (6)	115	18.1 (6.1)		100%	1.11[-0.45,2.67]
Subtotal ***	116		115			100%	1.11[-0.45,2.67]
Heterogeneity: Not applicable Test for overall effect: Z=1.39(P=0.16)							
11.14.4 endpoint score at 72 hours (PANSS negative sub-scale, high=worse)							
Li 2006	116	16.3 (5.6)	115	20.5 (6.2)		100%	-4.19[-5.71,-2.67]
Subtotal ***	116		115			100%	-4.19[-5.71,-2.67]
Heterogeneity: Not applicable Test for overall effect: Z=5.39(P<0.0001)							
11.14.5 endpoint score at 72 hours (PANSS general pathology score, high=worse)							
					haloperidol -5 -2.5 0 2.5 5 ziprasidone

Analysis 11.15. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 15 Adverse effects: 1a. General - i. up to 24 hours.

Analysis 11.16. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 16 Adverse effects: 1b. General - ii. over 24 hours.

Analysis 11.17. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 17 Adverse effects: 1c. General - serious.

Analysis 11.18. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 18 Adverse effects: 2a. Specific - anticholinergic.

**Analysis 11.19. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j.
ZIPRASIDONE, Outcome 19 Adverse effects: 2b. Specific - arousal level - i. various.**

Analysis 11.20. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 20 Adverse effects: 2c. Specific - arousal level - ii. average scores - i. up to 2 hours.

Analysis 11.21. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 21 Adverse effects: 2d. Specific - arousal level - ii. average scores - ii. up to 4 hours.

Analysis 11.22. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 22 Adverse effects: 2e. Specific - arousal level - ii. average scores - iii. up to 24 hours.

Analysis 11.23. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 23 Adverse effects: 2f. Specific - cardiovascular.

Analysis 11.24. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 24 Adverse effects: 2g. Specific - gastrointestinal.

Analysis 11.25. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 25 Adverse effects: 2h. Specific - hematological tests.

Study or subgroup	haloperidol n/N	ziprasidone n/N	Risk Ratio M-H, Fixed, 95% CI	Weight	Risk Ratio M-H, Fixed, 95% CI
Test for subgroup differences: $\chi^2=7.2$, $df=1$ ($P=0.41$), $I^2=2.84\%$					
	haloperidol		0.05 0.2 1 5 20		ziprasidone

Analysis 11.26. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 26 Adverse effects: 2i. Specific - movement disorders - i. binary measures.

Study or subgroup	haloperidol n/N	ziprasidone n/N	Risk Ratio M-H, Fixed, 95% CI	Weight	Risk Ratio M-H, Fixed, 95% CI
11.26.1 akathisia - by 72 hours					
Brook 2000	0/42	2/90		9.65%	0.42[0.02,8.63]
Li 2006	31/116	10/115		60.42%	3.07[1.58,5.97]
Shu 2010	7/187	5/189		29.92%	1.41[0.46,4.38]
Subtotal (95% CI)	345	394		100%	2.32[1.34,4.01]
Total events: 38 (haloperidol), 17 (ziprasidone)					
Heterogeneity: $\tau^2=0$; $\chi^2=2.65$, $df=2$ ($P=0.27$); $I^2=24.44\%$					
Test for overall effect: $Z=3.02$ ($P=0$)					
11.26.2 akathisia - by 7 days					
Brook 2000	6/42	3/90		100%	4.29[1.13,16.31]
Subtotal (95% CI)	42	90		100%	4.29[1.13,16.31]
Total events: 6 (haloperidol), 3 (ziprasidone)					
Heterogeneity: Not applicable					
Test for overall effect: $Z=2.13$ ($P=0.03$)					
11.26.3 dystonia - by 72 hours					
Brook 2000	3/42	1/90		56.13%	6.43[0.69,59.98]
Shu 2010	7/187	0/189		43.87%	15.16[0.87,263.55]
Subtotal (95% CI)	229	279		100%	10.26[1.67,63.17]
Total events: 10 (haloperidol), 1 (ziprasidone)					
Heterogeneity: $\tau^2=0$; $\chi^2=0.24$, $df=1$ ($P=0.62$); $I^2=0\%$					
Test for overall effect: $Z=2.51$ ($P=0.01$)					
11.26.4 dystonia - by 7 days					
Brook 2000	5/42	4/90		100%	2.68[0.76,9.47]
Subtotal (95% CI)	42	90		100%	2.68[0.76,9.47]
Total events: 5 (haloperidol), 4 (ziprasidone)					
Heterogeneity: Not applicable					
Test for overall effect: $Z=1.53$ ($P=0.13$)					
11.26.5 EPS - by 12 hours					
Baldacara 2011	5/30	0/30		100%	11[0.64,190.53]
Subtotal (95% CI)	30	30		100%	11[0.64,190.53]
Total events: 5 (haloperidol), 0 (ziprasidone)					
Heterogeneity: Not applicable					
Test for overall effect: $Z=1.65$ ($P=0.1$)					
11.26.6 EPS - by 72 hours					
Brook 2000	9/42	0/90		7.46%	40.21[2.4,674.98]
Shu 2010	69/187	4/189		92.54%	17.43[6.49,46.8]
	haloperidol		0.002 0.1 1 10 500		ziprasidone

Analysis 11.27. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 27 Adverse effects: 2j. Specific - movement disorders - ii. average scores - i. over 24 hours.

Analysis 11.28. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 28 Adverse effects: 2k. Specific - miscellaneous.

Analysis 11.29. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 29 Leaving the study early: 1. For any reason - i. over 24 hours.

Analysis 11.30. Comparison 11 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: j. ZIPRASIDONE, Outcome 30 Leaving the study early: 2. For specific reasons - i. over 24 hours.

Comparison 12. HALOPERIDOL vs OTHER ANTIPSYCHOTIC: k. ZUCLOPENTHIXOL ACETATE

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Repeated need for tranquillisation: more than 3 injections	1	70	Risk Ratio (M-H, Fixed, 95% CI)	2.54 [1.19, 5.46]
2 Adverse effects: 1a. Specific - movement disorders	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
2.1 movement disorder - tremor by 7 days	1	70	Risk Ratio (M-H, Fixed, 95% CI)	4.16 [0.93, 18.62]
3 Adverse effects: 1b. Specific - miscellaneous	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
3.1 pain/allergy - reaction at injection site	1	70	Risk Ratio (M-H, Fixed, 95% CI)	3.55 [0.15, 84.14]
4 Leaving the study early	1	70	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]

Analysis 12.1. Comparison 12 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: k. ZUCLOPENTHIXOL ACETATE, Outcome 1 Repeated need for tranquillisation: more than 3 injections.

Analysis 12.2. Comparison 12 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: k. ZUCLOPENTHIXOL ACETATE, Outcome 2 Adverse effects: 1a. Specific - movement disorders.

Analysis 12.3. Comparison 12 HALOPERIDOL vs OTHER ANTIPSYCHOTIC: k. ZUCLOPENTHIXOL ACETATE, Outcome 3 Adverse effects: 1b. Specific - miscellaneous.

**Analysis 12.4. Comparison 12 HALOPERIDOL vs OTHER ANTIPSYCHOTIC:
k. ZUCLOPENTHIXOL ACETATE, Outcome 4 Leaving the study early.**

Comparison 13. HALOPERIDOL vs BENZODIAZEPINE: a. FLUNITRAZEPAM

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Specific behaviour: 1. Aggression - binary measures - i. up to 2 hours	1	28	Risk Ratio (M-H, Fixed, 95% CI)	1.15 [0.86, 1.55]
1.1 ≥50% reduction at 90 minutes (OAS)	1	28	Risk Ratio (M-H, Fixed, 95% CI)	1.15 [0.86, 1.55]
2 Global outcome: 1. Need for seclusion or restraint	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
3 Adverse effects: 1. Specific - movement disorders - i. binary measures - i. by 30 minutes	1	28	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
3.1 EPS by 30 minutes	1	28	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]

**Analysis 13.1. Comparison 13 HALOPERIDOL vs BENZODIAZEPINE: a. FLUNITRAZEPAM,
Outcome 1 Specific behaviour: 1. Aggression - binary measures - i. up to 2 hours.**

Analysis 13.2. Comparison 13 HALOPERIDOL vs BENZODIAZEPINE: a. FLUNITRAZEPAM, Outcome 2 Global outcome: 1. Need for seclusion or restraint.

Analysis 13.3. Comparison 13 HALOPERIDOL vs BENZODIAZEPINE: a. FLUNITRAZEPAM, Outcome 3 Adverse effects: 1. Specific - movement disorders - i. binary measures - i. by 30 minutes.

Comparison 14. HALOPERIDOL vs BENZODIAZEPINE: b. LORAZEPAM

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Tranquillisation or asleep - not asleep	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1.1 at 1 hour	1	60	Risk Ratio (M-H, Fixed, 95% CI)	1.05 [0.76, 1.44]
1.2 by 3 hours	1	66	Risk Ratio (M-H, Fixed, 95% CI)	1.93 [1.14, 3.27]
2 Repeated need for rapid tranquillisation	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
2.1 more than 1 injection	1	66	Risk Ratio (M-H, Fixed, 95% CI)	1.14 [0.91, 1.43]
2.2 more than 3 injections	1	66	Risk Ratio (M-H, Fixed, 95% CI)	1.11 [0.50, 2.45]
3 Global outcome: 1. No overall improvement	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
3.1 by 30 minutes	1	44	Risk Ratio (M-H, Fixed, 95% CI)	1.10 [0.70, 1.71]
3.2 at 1 hour	1	44	Risk Ratio (M-H, Fixed, 95% CI)	1.64 [0.54, 5.03]
3.3 at > 1 hour	1	44	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
4 Mental state: 1a. Average scores - i. up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
4.1 endpoint score at 1 hour (BPRS, high=worse)	1	37	Mean Difference (IV, Fixed, 95% CI)	3.26 [-4.13, 10.65]
4.2 endpoint score at 2 hours (BPRS, high=worse)	1	37	Mean Difference (IV, Fixed, 95% CI)	4.07 [-2.62, 10.76]
5 Mental state: 1b. Average scores - ii. up to 4 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
5.1 endpoint score at 3 hours (BPRS, high=worse)	1	37	Mean Difference (IV, Fixed, 95% CI)	5.03 [-2.98, 13.04]
5.2 endpoint score at 4 hours (BPRS, high=worse)	1	37	Mean Difference (IV, Fixed, 95% CI)	1.73 [-6.14, 9.60]
6 Adverse effects: 1a. General	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
6.1 one or more drug-related adverse effect up to 24 hours	1	66	Risk Ratio (M-H, Fixed, 95% CI)	1.13 [0.60, 2.10]
7 Adverse effects: 1b. General - serious	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
8 Adverse effects: 2a. Specific - anti-cholinergic	1	66	Risk Ratio (M-H, Fixed, 95% CI)	0.53 [0.14, 2.04]
8.1 dry mouth up to 24 hours (only reported if occurred $\geq 9\%$)	1	66	Risk Ratio (M-H, Fixed, 95% CI)	0.53 [0.14, 2.04]
9 Adverse effects: 2b. Specific - arousal	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
9.1 asleep - at 1 hour	1	60	Risk Ratio (M-H, Fixed, 95% CI)	0.89 [0.40, 1.99]
10 Adverse effects: 2c. Specific - cardiovascular	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
10.1 dizziness (only reported if occurred in $\geq 9\%$) - during 24 hours	1	66	Risk Ratio (M-H, Fixed, 95% CI)	0.89 [0.19, 4.07]
10.2 hypertension - use of additional clonidine - during 24 hours	1	66	Risk Ratio (M-H, Fixed, 95% CI)	2.67 [0.11, 63.17]
11 Adverse effects: 2d. Specific - movement disorder	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
11.1 ataxia (only reported if occurred in $\geq 9\%$) - during 24 hours	1	66	Risk Ratio (M-H, Fixed, 95% CI)	0.44 [0.04, 4.65]
11.2 dystonia (only reported if occurred in $\geq 9\%$) - during 24 hours	1	66	Risk Ratio (M-H, Fixed, 95% CI)	3.54 [0.42, 30.03]
11.3 EPS	1	60	Risk Ratio (M-H, Fixed, 95% CI)	15.0 [2.11, 106.49]
11.4 hypertonia/rigidity (only reported if occurred in $\geq 9\%$) - during 24 hours	1	66	Risk Ratio (M-H, Fixed, 95% CI)	6.22 [0.33, 115.91]
11.5 speech disorder (only reported if occurred in $\geq 9\%$) - during 24 hours	1	66	Risk Ratio (M-H, Fixed, 95% CI)	1.77 [0.35, 9.01]
11.6 tremor (only reported if occurred in $\geq 9\%$) - during 24 hours	1	66	Risk Ratio (M-H, Fixed, 95% CI)	1.77 [0.17, 18.60]
11.7 use of additional benzotropine - during 24 hours	1	66	Risk Ratio (M-H, Fixed, 95% CI)	1.99 [0.68, 5.83]

**Analysis 14.1. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE:
b. LORAZEPAM, Outcome 1 Tranquillisation or asleep - not asleep.**

Study or subgroup	Haloperidol n/N	Lorazepam n/N	Risk Ratio M-H, Fixed, 95% CI	Weight	Risk Ratio M-H, Fixed, 95% CI
14.1.1 at 1 hour					
		haloperidol	0.2 0.5 1 2 5		lorazepam

**Analysis 14.2. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE:
b. LORAZEPAM, Outcome 2 Repeated need for rapid tranquillisation.**

**Analysis 14.3. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b.
LORAZEPAM, Outcome 3 Global outcome: 1. No overall improvement.**

Analysis 14.4. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b. LORAZEPAM, Outcome 4 Mental state: 1a. Average scores - i. up to 2 hours.

Analysis 14.5. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b. LORAZEPAM, Outcome 5 Mental state: 1b. Average scores - ii. up to 4 hours.

Analysis 14.6. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b. LORAZEPAM, Outcome 6 Adverse effects: 1a. General.

Analysis 14.7. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b. LORAZEPAM, Outcome 7 Adverse effects: 1b. General - serious.

Analysis 14.8. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: LORAZEPAM, Outcome 8 Adverse effects: 2a. Specific - anticholinergic.

**Analysis 14.9. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE:
b. LORAZEPAM, Outcome 9 Adverse effects: 2b. Specific - arousal.**

**Analysis 14.10. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b.
LORAZEPAM, Outcome 10 Adverse effects: 2c. Specific - cardiovascular.**

**Analysis 14.11. Comparison 14 HALOPERIDOL vs BENZODIAZEPINE: b.
LORAZEPAM, Outcome 11 Adverse effects: 2d. Specific - movement disorder.**

Comparison 15. HALOPERIDOL vs BENZODIAZEPINE: c. MIDAZOLAM

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Tranquillisation or asleep: average times in minutes - (skewed data)			Other data	No numeric data
1.1 average time to sedation			Other data	No numeric data
1.2 average time to arousal			Other data	No numeric data
2 Global outcome: 1. Need for rescue drug	1	84	Risk Ratio (M-H, Fixed, 95% CI)	1.14 [0.46, 2.87]
3 Adverse effects: 1. General - one or more drug related adverse effect	1	84	Risk Ratio (M-H, Fixed, 95% CI)	5.0 [0.25, 101.11]
4 Adverse effects: 2. Specific	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
4.1 hypotensive	1	84	Risk Ratio (M-H, Fixed, 95% CI)	3.0 [0.13, 71.61]
4.2 apnoea	1	84	Risk Ratio (M-H, Fixed, 95% CI)	3.0 [0.13, 71.61]

Analysis 15.1. Comparison 15 HALOPERIDOL vs BENZODIAZEPINE: c. MIDAZOLAM, Outcome 1 Tranquillisation or asleep: average times in minutes - (skewed data).

Tranquillisation or asleep: average times in minutes - (skewed data)					
Study	Intervention	Average	SD	N	Notes
average time to sedation					
Nobay 2004	Haloperidol	28.3	25	42	
Nobay 2004	Midazolam	18.3	14	42	
average time to arousal					
Nobay 2004	Haloperidol	126.6	85	42	
Nobay 2004	Midazolam	81.9	66	42	

Analysis 15.2. Comparison 15 HALOPERIDOL vs BENZODIAZEPINE: c. MIDAZOLAM, Outcome 2 Global outcome: 1. Need for rescue drug.

Study or subgroup	haloperidol n/N	midazolam n/N	Risk Ratio M-H, Fixed, 95% CI	Weight	Risk Ratio M-H, Fixed, 95% CI
Nobay 2004	8/42	7/42		100%	1.14[0.46,2.87]
Total (95% CI)	42	42		100%	1.14[0.46,2.87]
Total events: 8 (haloperidol), 7 (midazolam)					
Heterogeneity: Not applicable					
Test for overall effect: Z=0.28(P=0.78)					
haloperidol 0.01 0.1 1 10 100 midazolam					

Analysis 15.3. Comparison 15 HALOPERIDOL vs BENZODIAZEPINE: c. MIDAZOLAM, Outcome 3 Adverse effects: 1. General - one or more drug related adverse effect.

Analysis 15.4. Comparison 15 HALOPERIDOL vs BENZODIAZEPINE: c. MIDAZOLAM, Outcome 4 Adverse effects: 2. Specific.

Comparison 16. HALOPERIDOL vs COMBINATIONS: a. HALOPERIDOL + LEVOMEPRMAZINE

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Global outcome: 1. No overall improvement	1	19	Risk Ratio (M-H, Fixed, 95% CI)	8.18 [0.50, 133.66]

Analysis 16.1. Comparison 16 HALOPERIDOL vs COMBINATIONS: a. HALOPERIDOL + LEVOMEPRMAZINE, Outcome 1 Global outcome: 1. No overall improvement.

Comparison 17. HALOPERIDOL vs COMBINATIONS: b. HALOPERIDOL + LORAZEPAM

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Tranquillisation or asleep - asleep	1	67	Risk Ratio (M-H, Fixed, 95% CI)	1.83 [1.11, 3.02]
1.1 not asleep by 3 hours	1	67	Risk Ratio (M-H, Fixed, 95% CI)	1.83 [1.11, 3.02]
2 Repeated need for rapid tranquillisation	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
2.1 needing additional injection during 24 hours	1	67	Risk Ratio (M-H, Fixed, 95% CI)	1.05 [0.87, 1.27]
2.2 more than 3 injections during 24 hours	1	67	Risk Ratio (M-H, Fixed, 95% CI)	3.05 [0.92, 10.10]
3 Global outcome: 1. No overall improvement	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
3.1 by 30 minutes	1	45	Risk Ratio (M-H, Fixed, 95% CI)	2.67 [1.25, 5.68]
3.2 at 1 hour	1	45	Risk Ratio (M-H, Fixed, 95% CI)	14.77 [0.88, 247.54]
3.3 at > 1 hour	1	45	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
4 Adverse effects: 1. General	1	67	Risk Ratio (M-H, Fixed, 95% CI)	1.16 [0.62, 2.18]
4.1 one or more drug-related adverse effect during 24 hours	1	67	Risk Ratio (M-H, Fixed, 95% CI)	1.16 [0.62, 2.18]
5 Adverse effects: 2a. Specific - anticholinergic	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
5.1 dry mouth (only reported if occurred in $\geq 9\%$) during 24 hours	1	67	Risk Ratio (M-H, Fixed, 95% CI)	0.91 [0.20, 4.21]
6 Adverse effects: 2b. Specific - cardiovascular	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
6.1 dizziness (only reported if occurred in $\geq 9\%$) during 24 hours	1	67	Risk Ratio (M-H, Fixed, 95% CI)	1.37 [0.24, 7.69]
6.2 hypertension - use of additional clonidine during 24 hours	1	67	Risk Ratio (M-H, Fixed, 95% CI)	0.91 [0.06, 14.02]
7 Adverse effects: 2c. Specific - movement disorders	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
7.1 ataxia (only reported if occurred in $>9\%$) during 24 hours	1	67	Risk Ratio (M-H, Fixed, 95% CI)	0.30 [0.03, 2.78]
7.2 dystonia (only reported if occurred in $>9\%$) during 24 hours	1	67	Risk Ratio (M-H, Fixed, 95% CI)	8.25 [0.46, 147.45]
7.3 hypertonia/rigidity (only reported if occurred in $>9\%$) during 24 hours	1	67	Risk Ratio (M-H, Fixed, 95% CI)	2.74 [0.30, 25.05]
7.4 tremor (only reported if occurred in $>9\%$) during 24 hours	1	67	Risk Ratio (M-H, Fixed, 95% CI)	1.83 [0.17, 19.21]
7.5 speech disorder (only reported if occurred in $>9\%$) during 24 hours	1	67	Risk Ratio (M-H, Fixed, 95% CI)	1.22 [0.30, 5.03]
7.6 use of additional benztropine during 24 hours	1	67	Risk Ratio (M-H, Fixed, 95% CI)	2.74 [0.81, 9.25]

Analysis 17.1. Comparison 17 HALOPERIDOL vs COMBINATIONS: b. HALOPERIDOL + LORAZEPAM, Outcome 1 Tranquillisation or asleep - asleep.

Study or subgroup	Haloperidol n/N	Combination n/N	Risk Ratio M-H, Fixed, 95% CI	Weight	Risk Ratio M-H, Fixed, 95% CI
17.1.1 not asleep by 3 hours					
Battaglia 1997	24/35	12/32		100%	1.83[1.11,3.02]
Subtotal (95% CI)	35	32		100%	1.83[1.11,3.02]
Total events: 24 (Haloperidol), 12 (Combination)					
Heterogeneity: $\tau^2=0$; $\chi^2=0$, $df=0$ ($P<0.0001$); $I^2=100\%$					
Test for overall effect: $Z=2.36$ ($P=0.02$)					
Total (95% CI)	35	32		100%	1.83[1.11,3.02]
Total events: 24 (Haloperidol), 12 (Combination)					
Heterogeneity: $\tau^2=0$; $\chi^2=0$, $df=0$ ($P<0.0001$); $I^2=100\%$					
Test for overall effect: $Z=2.36$ ($P=0.02$)					
			haloperidol	0.5 0.7 1 1.5 2	combination

Analysis 17.2. Comparison 17 HALOPERIDOL vs COMBINATIONS: b. HALOPERIDOL + LORAZEPAM, Outcome 2 Repeated need for rapid tranquillisation.

Analysis 17.3. Comparison 17 HALOPERIDOL vs COMBINATIONS: b. HALOPERIDOL + LORAZEPAM, Outcome 3 Global outcome: 1. No overall improvement.

Analysis 17.4. Comparison 17 HALOPERIDOL vs COMBINATIONS: b. HALOPERIDOL + LORAZEPAM, Outcome 4 Adverse effects: 1. General.

Analysis 17.5. Comparison 17 HALOPERIDOL vs COMBINATIONS: b. HALOPERIDOL + LORAZEPAM, Outcome 5 Adverse effects: 2a. Specific - anticholinergic.

Analysis 17.6. Comparison 17 HALOPERIDOL vs COMBINATIONS: b. HALOPERIDOL + LORAZEPAM, Outcome 6 Adverse effects: 2b. Specific - cardiovascular.

Study or subgroup	Haloperidol n/N	Combination n/N	Risk Ratio M-H, Fixed, 95% CI	Weight	Risk Ratio M-H, Fixed, 95% CI
Test for subgroup differences: $\chi^2=0.06$, $df=1$ ($P=0.81$), $I^2=0\%$					
	haloperidol		0.01 0.1 1 10 100		combination

Analysis 17.7. Comparison 17 HALOPERIDOL vs COMBINATIONS: b. HALOPERIDOL + LORAZEPAM, Outcome 7 Adverse effects: 2c. Specific - movement disorders.

Study or subgroup	Haloperidol n/N	Combination n/N	Risk Ratio M-H, Fixed, 95% CI	Weight	Risk Ratio M-H, Fixed, 95% CI
17.7.1 ataxia (only reported if occurred in >9%) during 24 hours					
Battaglia 1997	1/35	3/32		100%	0.3[0.03,2.78]
Subtotal (95% CI)	35	32		100%	0.3[0.03,2.78]
Total events: 1 (Haloperidol), 3 (Combination)					
Heterogeneity: Not applicable					
Test for overall effect: $Z=1.05$ ($P=0.29$)					
17.7.2 dystonia (only reported if occurred in >9%) during 24 hours					
Battaglia 1997	4/35	0/32		100%	8.25[0.46,147.45]
Subtotal (95% CI)	35	32		100%	8.25[0.46,147.45]
Total events: 4 (Haloperidol), 0 (Combination)					
Heterogeneity: Not applicable					
Test for overall effect: $Z=1.43$ ($P=0.15$)					
17.7.3 hypertonia/rigidity (only reported if occurred in >9%) during 24 hours					
Battaglia 1997	3/35	1/32		100%	2.74[0.3,25.05]
Subtotal (95% CI)	35	32		100%	2.74[0.3,25.05]
Total events: 3 (Haloperidol), 1 (Combination)					
Heterogeneity: Not applicable					
Test for overall effect: $Z=0.89$ ($P=0.37$)					
17.7.4 tremor (only reported if occurred in >9%) during 24 hours					
Battaglia 1997	2/35	1/32		100%	1.83[0.17,19.21]
Subtotal (95% CI)	35	32		100%	1.83[0.17,19.21]
Total events: 2 (Haloperidol), 1 (Combination)					
Heterogeneity: Not applicable					
Test for overall effect: $Z=0.5$ ($P=0.62$)					
17.7.5 speech disorder (only reported if occurred in >9%) during 24 hours					
Battaglia 1997	4/35	3/32		100%	1.22[0.3,5.03]
Subtotal (95% CI)	35	32		100%	1.22[0.3,5.03]
Total events: 4 (Haloperidol), 3 (Combination)					
Heterogeneity: Not applicable					
Test for overall effect: $Z=0.27$ ($P=0.78$)					
17.7.6 use of additional benzotropine during 24 hours					
Battaglia 1997	9/35	3/32		100%	2.74[0.81,9.25]
Subtotal (95% CI)	35	32		100%	2.74[0.81,9.25]
Total events: 9 (Haloperidol), 3 (Combination)					
Heterogeneity: Not applicable					
	haloperidol		0.05 0.2 1 5 20		combination

Comparison 18. HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Repeated need for tranquillisation	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
1.1 need for additional drugs for tranquillisation up to 2 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	0.27 [0.10, 0.71]
1.2 need for additional drugs for tranquillisation by 12 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	0.88 [0.69, 1.13]
2 Specific behaviour: 1a. Agitation - average scores - i. up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
2.1 endpoint score at 1 hour (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-8.5 [-9.93, -7.07]
2.2 endpoint score at 2 hours (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-6.7 [-7.46, -5.94]
3 Specific behaviour: 1b. Agitation - average scores - ii. up to 4 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
3.1 endpoint score at 4 hours (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-5.5 [-6.48, -4.52]
4 Specific behaviour: 1c. Agitation - average scores - iii. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
4.1 endpoint score at 6 hours (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-3.70 [-4.56, -2.84]
4.2 endpoint score at 12 hours (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-11.2 [-12.24, -10.16]
5 Specific behaviour: 2a. Aggression - average scores - i. up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
5.1 endpoint score at 1 hour (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-1.20 [-2.44, 0.04]
5.2 endpoint score at 2 hours (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-2.40 [-4.21, -0.59]
6 Specific behaviour: 2b. Aggression - average scores - ii. up to 4 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
6.1 endpoint score at 4 hours (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-0.10 [-0.68, 0.48]
7 Specific behaviour: 2c. Aggression - average scores - iii. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
7.1 endpoint score at 6 hours (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.40 [0.30, 0.50]
7.2 endpoint score at 12 hours (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-1.9 [-2.58, -1.22]
8 Global outcomes: 1. General - need for additional measures	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
8.1 additional restraint, seclusion or special observation during 12 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	0.29 [0.13, 0.61]
9 Adverse effects: 1b. General - one or more adverse events	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
9.1 during 12 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	0.73 [0.41, 1.32]
10 Adverse effects: 1a. General - serious	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
10.1 death	1	60	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
11 Adverse effects: 2a. Specific - arousal - i. various	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
11.1 excessive sedation by 12 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	0.25 [0.08, 0.80]
12 Adverse effects: 2b. Specific - arousal - ii. average scores - i. up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
12.1 endpoint score at 1 hour (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-0.5 [-1.01, 0.01]
12.2 endpoint score at 2 hours (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-0.10 [-0.56, 0.36]
13 Adverse effects: 2c. Specific - arousal - ii. average scores - ii. up to 4 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
13.1 endpoint score at 4 hours (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-0.30 [-0.78, 0.18]
14 Adverse effects: 2d. Specific - arousal - iii. average scores - iii. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
14.1 endpoint score at 6 hours (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-0.10 [-0.40, 0.20]
14.2 endpoint score at 12 hours (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.10 [-0.26, 0.46]
15 Adverse effects: 2e. Specific - cardiovascular	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
15.1 hypotension by 12 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	0.09 [0.01, 1.57]
16 Adverse effects: 2f. Specific - movement disorders	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
16.1 EPS by 12 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	1.67 [0.44, 6.36]

Analysis 18.1. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 1 Repeated need for tranquillisation.

Analysis 18.2. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 2 Specific behaviour: 1a. Agitation - average scores - i. up to 2 hours.

Analysis 18.3. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 3 Specific behaviour: 1b. Agitation - average scores - ii. up to 4 hours.

Analysis 18.4. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 4 Specific behaviour: 1c. Agitation - average scores - iii. up to 24 hours.

Analysis 18.5. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 5 Specific behaviour: 2a. Aggression - average scores - i. up to 2 hours.

Analysis 18.6. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 6 Specific behaviour: 2b. Aggression - average scores - ii. up to 4 hours.

Analysis 18.7. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 7 Specific behaviour: 2c. Aggression - average scores - iii. up to 24 hours.

Analysis 18.8. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 8 Global outcomes: 1. General - need for additional measures.

Analysis 18.9. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 9 Adverse effects: 1b. General - one or more adverse events.

Analysis 18.10. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 10 Adverse effects: 1a. General - serious.

Analysis 18.11. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 11 Adverse effects: 2a. Specific - arousal - i. various.

Analysis 18.12. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 12 Adverse effects: 2b. Specific - arousal - ii. average scores - i. up to 2 hours.

Analysis 18.13. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 13 Adverse effects: 2c. Specific - arousal - ii. average scores - ii. up to 4 hours.

Analysis 18.14. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 14 Adverse effects: 2d. Specific - arousal - iii. average scores - iii. up to 24 hours.

Analysis 18.15. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 15 Adverse effects: 2e. Specific - cardiovascular.

Analysis 18.16. Comparison 18 HALOPERIDOL vs COMBINATIONS: c. HALOPERIDOL + MIDAZOLAM, Outcome 16 Adverse effects: 2f. Specific - movement disorders.

Comparison 19. HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Tranquillisation or asleep - not tranquil or asleep - i. by 30 minutes	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
1.1 at 20 minutes	1	316	Risk Ratio (M-H, Fixed, 95% CI)	1.60 [1.18, 2.16]
2 Tranquillisation or asleep - not tranquil or asleep - ii. up to 2 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
2.1 at 40 minutes	1	316	Risk Ratio (M-H, Fixed, 95% CI)	1.26 [0.83, 1.91]
2.2 at 1 hour	1	316	Risk Ratio (M-H, Fixed, 95% CI)	1.42 [0.85, 2.37]
2.3 at 2 hours	1	316	Risk Ratio (M-H, Fixed, 95% CI)	1.32 [0.68, 2.56]
3 Repeated need for tranquillisation - need for additional drugs for tranquillisation	2	436	Risk Ratio (M-H, Fixed, 95% CI)	0.94 [0.68, 1.29]
3.1 by 2 hours	2	376	Risk Ratio (M-H, Fixed, 95% CI)	0.78 [0.43, 1.41]
3.2 by 12 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	1.10 [0.81, 1.49]
4 Specific behaviour: 1a. Agitation - average scores - i. up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
4.1 endpoint score at 1 hour (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-24.5 [-27.32, -21.68]
4.2 endpoint score at 2 hours (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-9.40 [-10.39, -8.41]
5 Specific behaviour: 1b. Agitation - average scores - ii. up to 4 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
5.1 endpoint score at 4 hours (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	3.80 [3.27, 4.33]
6 Specific behaviour: 1c. Agitation - average scores - iii. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
6.1 endpoint score at 6 hours (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	2.6 [2.13, 3.07]
6.2 endpoint score at 12 hours (OASS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.80 [0.55, 1.05]
7 Specific behaviour: 2a. Aggression - average scores - i. up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
7.1 endpoint score at 1 hour (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-4.50 [-6.28, -2.72]
7.2 endpoint score at 2 hours (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-0.70 [-1.89, 0.49]
8 Specific behaviour: 2b. Aggression - average scores - ii. up to 4 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
8.1 endpoint score at 4 hours (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.60 [0.49, 0.71]
9 Specific behaviour: 2c. Aggression - average scores - iii. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
9.1 endpoint score at 6 hours (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	1.1 [0.91, 1.29]
9.2 endpoint score at 12 hours (OAS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	1.8 [1.67, 1.93]
10 Specific behaviour: 2d. Aggression - further aggressive episode - i. up to 24 hours	1	316	Risk Ratio (M-H, Fixed, 95% CI)	1.06 [0.68, 1.65]
11 Global outcomes: 1. General - need for additional measures	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
11.1 restraints needed by 2 hours	1	316	Risk Ratio (M-H, Fixed, 95% CI)	1.21 [0.84, 1.76]
11.2 additional restraint, seclusion or special observation during 12 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	1.2 [0.41, 3.51]
12 Global outcomes: 2. Specific	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
12.1 doctor called to see patient	1	316	Risk Ratio (M-H, Fixed, 95% CI)	1.50 [1.05, 2.14]
12.2 refuse oral drugs	1	316	Risk Ratio (M-H, Fixed, 95% CI)	0.99 [0.61, 1.60]
13 Adverse effects: 1a. General	2	376	Risk Ratio (M-H, Fixed, 95% CI)	2.01 [1.07, 3.80]
13.1 one or more adverse effects up to 24 hours	2	376	Risk Ratio (M-H, Fixed, 95% CI)	2.01 [1.07, 3.80]
14 Adverse effects: 1b. General - serious	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
14.1 seizure	1	316	Risk Ratio (M-H, Fixed, 95% CI)	2.05 [0.19, 22.39]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
14.2 death	1	60	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
15 Adverse effects: 2a. Specific - arousal - i. various	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
15.1 excessive sedation by 12 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	3.0 [0.33, 27.23]
16 Adverse effects: 2b. Specific - arousal - ii. average scores - i. up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
16.1 endpoint score at 1 hour (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.10 [-0.38, 0.58]
16.2 endpoint score at 2 hours (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-0.10 [-0.50, 0.30]
17 Adverse effects: 2c. Specific - arousal - ii. average scores - ii. up to 4 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
17.1 endpoint score at 4 hours (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-0.30 [-0.78, 0.18]
18 Adverse effects: 2d. Specific - arousal - ii. average scores - iii. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
18.1 endpoint score at 6 hours (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	-0.20 [-0.48, 0.08]
18.2 endpoint score at 12 hours (RSS, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.0 [-0.28, 0.28]
19 Adverse effects: 2e. Specific - cardiovascular	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
19.1 hypotension by 12 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	0.14 [0.01, 2.65]
20 Adverse effects: 2f. Specific - movement disorders	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
20.1 acute dystonia	1	316	Risk Ratio (M-H, Fixed, 95% CI)	19.48 [1.14, 331.92]
20.2 EPS by 12 hours	1	60	Risk Ratio (M-H, Fixed, 95% CI)	1.0 [0.32, 3.10]
21 Service outcomes: 1. Not discharged by 14 days	1	316	Risk Ratio (M-H, Fixed, 95% CI)	0.87 [0.72, 1.05]
22 Leaving the study early	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
22.1 any reason	1	316	Risk Ratio (M-H, Fixed, 95% CI)	1.03 [0.54, 1.94]
22.2 absconded before receiving drug	1	316	Risk Ratio (M-H, Fixed, 95% CI)	0.21 [0.01, 4.24]
22.3 incomplete information in notes	1	316	Risk Ratio (M-H, Fixed, 95% CI)	1.12 [0.51, 2.46]
22.4 seizure before drug was given	1	316	Risk Ratio (M-H, Fixed, 95% CI)	3.08 [0.13, 74.95]
22.5 transfer to another hospital/notes lost	1	316	Risk Ratio (M-H, Fixed, 95% CI)	1.03 [0.21, 5.00]
22.6 withdrew consent	1	316	Risk Ratio (M-H, Fixed, 95% CI)	1.03 [0.06, 16.25]

Analysis 19.1. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 1 Tranquillisation or asleep - not tranquil or asleep - i. by 30 minutes.

Analysis 19.2. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 2 Tranquillisation or asleep - not tranquil or asleep - ii. up to 2 hours.

Analysis 19.3. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 3 Repeated need for tranquillisation - need for additional drugs for tranquillisation.

Analysis 19.4. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 4 Specific behaviour: 1a. Agitation - average scores - i. up to 2 hours.

Analysis 19.5. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 5 Specific behaviour: 1b. Agitation - average scores - ii. up to 4 hours.

Analysis 19.6. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 6 Specific behaviour: 1c. Agitation - average scores - iii. up to 24 hours.

Analysis 19.7. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 7 Specific behaviour: 2a. Aggression - average scores - i. up to 2 hours.

Analysis 19.8. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 8 Specific behaviour: 2b. Aggression - average scores - ii. up to 4 hours.

Analysis 19.9. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 9 Specific behaviour: 2c. Aggression - average scores - iii. up to 24 hours.

Analysis 19.10. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 10 Specific behaviour: 2d. Aggression - further aggressive episode - i. up to 24 hours.

Analysis 19.11. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 11 Global outcomes: 1. General - need for additional measures.

Analysis 19.12. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 12 Global outcomes: 2. Specific.

Analysis 19.13. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 13 Adverse effects: 1a. General.

Analysis 19.14. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 14 Adverse effects: 1b. General - serious.

Analysis 19.15. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 15 Adverse effects: 2a. Specific - arousal - i. various.

Analysis 19.16. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 16 Adverse effects: 2b. Specific - arousal - ii. average scores - i. up to 2 hours.

Analysis 19.17. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 17 Adverse effects: 2c. Specific - arousal - ii. average scores - ii. up to 4 hours.

Analysis 19.18. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 18 Adverse effects: 2d. Specific - arousal - ii. average scores - iii. up to 24 hours.

Analysis 19.19. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 19 Adverse effects: 2e. Specific - cardiovascular.

Analysis 19.20. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 20 Adverse effects: 2f. Specific - movement disorders.

Study or subgroup	haloperidol n/N	combination n/N	Risk Ratio M-H, Fixed, 95% CI	Weight	Risk Ratio M-H, Fixed, 95% CI
Test for overall effect: Not applicable					
Test for subgroup differences: $\chi^2=3.63$, $df=1$ ($P=0.06$), $I^2=72.49\%$					
	haloperidol		0.002 0.1 1 10 500	combination	

Analysis 19.21. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 21 Service outcomes: 1. Not discharged by 14 days.

Study or subgroup	haloperidol n/N	combination n/N	Risk Ratio M-H, Fixed, 95% CI	Weight	Risk Ratio M-H, Fixed, 95% CI
Huf 2007	83/156	98/160		100%	0.87[0.72,1.05]
Total (95% CI)	156	160		100%	0.87[0.72,1.05]
Total events: 83 (haloperidol), 98 (combination)					
Heterogeneity: Not applicable					
Test for overall effect: $Z=1.44$ ($P=0.15$)					
	haloperidol		0.5 0.7 1 1.5 2	combination	

Analysis 19.22. Comparison 19 HALOPERIDOL vs COMBINATIONS: d. HALOPERIDOL + PROMETHAZINE, Outcome 22 Leaving the study early.

Study or subgroup	haloperidol n/N	combination n/N	Risk Ratio M-H, Fixed, 95% CI	Weight	Risk Ratio M-H, Fixed, 95% CI
19.22.1 any reason					
Huf 2007	17/156	17/160		100%	1.03[0.54,1.94]
Subtotal (95% CI)	156	160		100%	1.03[0.54,1.94]
Total events: 17 (haloperidol), 17 (combination)					
Heterogeneity: Not applicable					
Test for overall effect: $Z=0.08$ ($P=0.94$)					
19.22.2 absconded before receiving drug					
Huf 2007	0/156	2/160		100%	0.21[0.01,4.24]
Subtotal (95% CI)	156	160		100%	0.21[0.01,4.24]
Total events: 0 (haloperidol), 2 (combination)					
Heterogeneity: Not applicable					
Test for overall effect: $Z=1.03$ ($P=0.31$)					
19.22.3 incomplete information in notes					
Huf 2007	12/156	11/160		100%	1.12[0.51,2.46]
Subtotal (95% CI)	156	160		100%	1.12[0.51,2.46]
Total events: 12 (haloperidol), 11 (combination)					
Heterogeneity: Not applicable					
Test for overall effect: $Z=0.28$ ($P=0.78$)					
19.22.4 seizure before drug was given					
Huf 2007	1/156	0/160		100%	3.08[0.13,74.95]
Subtotal (95% CI)	156	160		100%	3.08[0.13,74.95]
Total events: 1 (haloperidol), 0 (combination)					
	haloperidol		0.005 0.1 1 10 200	combination

Comparison 20. HALOPERIDOL vs COMBINATIONS: e. HALOPERIDOL + RISPERIDONE

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Specific behaviour: 1a. Agitation - average scores - i. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
1.1 endpoint score at 24 hours (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2 Specific behaviour: 1b. Agitation - average scores - ii. over 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
2.1 endpoint score at 3 days (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2.2 endpoint score at 5 days (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2.3 endpoint score at 1 week (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2.4 endpoint score at 2 weeks (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
3 Global outcome: No improvement	1		Risk Ratio (M-H, Fixed, 95% CI)	Totals not selected
3.1 at 72 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
3.2 at 5 days	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
3.3 at 1 week	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
3.4 at 2 weeks	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
4 Adverse effects: 1a. Specific - arousal level - i. various	1		Risk Ratio (M-H, Fixed, 95% CI)	Totals not selected
4.1 insomnia during 14 days	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
4.2 somnolence during 14 days	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
5 Adverse effects: 1b. Specific - movement disorders - i. various	1		Risk Ratio (M-H, Fixed, 95% CI)	Totals not selected
5.1 tremor during 14 days	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
5.2 akathisia during 14 days	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
6 Adverse effects: 1c. Specific - miscellaneous	1		Risk Ratio (M-H, Fixed, 95% CI)	Totals not selected
6.1 Dry mouth during 14 days	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
6.2 Constipation during 14 days	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
6.3 Increased salivation during 14 days	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]

Analysis 20.1. Comparison 20 HALOPERIDOL vs COMBINATIONS: e. HALOPERIDOL + RISPERIDONE, Outcome 1 Specific behaviour: 1a. Agitation - average scores - i. up to 24 hours.

Study or subgroup	Haloperidol		Comparison		Mean Difference Fixed, 95% CI	Mean Difference Fixed, 95% CI
	N	Mean(SD)	N	Mean(SD)		
20.1.1 endpoint score at 24 hours (PANSS-EC, high=worse)						
Liu 2012	50	30 (5.5)	50	26.2 (7)		3.82[1.35,6.29]
				haloperidol	-20 -10 0 10 20	comparison

Analysis 20.2. Comparison 20 HALOPERIDOL vs COMBINATIONS: e. HALOPERIDOL + RISPERIDONE, Outcome 2 Specific behaviour: 1b. Agitation - average scores - ii. over 24 hours.

Study or subgroup	Haloperidol		Comparison		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
20.2.1 endpoint score at 3 days (PANSS-EC, high=worse)						
Liu 2012	50	17.6 (6.6)	50	17.2 (7.1)		0.34[-2.34,3.02]
20.2.2 endpoint score at 5 days (PANSS-EC, high=worse)						
Liu 2012	50	15.8 (7)	50	15.1 (6.4)		0.74[-1.9,3.38]
					haloperidol	comparison

Study or subgroup	Haloperidol		Comparison		Mean Difference Fixed, 95% CI	Mean Difference Fixed, 95% CI
	N	Mean(SD)	N	Mean(SD)		
20.2.3 endpoint score at 1 week (PANSS-EC, high=worse)						
Liu 2012	50	15 (5.6)	50	14.6 (5.4)		0.42[-1.73,2.57]
20.2.4 endpoint score at 2 weeks (PANSS-EC, high=worse)						
Liu 2012	50	12.5 (3.5)	50	13.1 (4.1)		-0.64[-2.13,0.85]
					haloperidol	comparison

Analysis 20.3. Comparison 20 HALOPERIDOL vs COMBINATIONS: e. HALOPERIDOL + RISPERIDONE, Outcome 3 Global outcome: No improvement.

Study or subgroup	Haloperidol n/N	Comparison n/N	Risk Ratio M-H, Fixed, 95% CI	Risk Ratio M-H, Fixed, 95% CI
20.3.1 at 72 hours				
Liu 2012	3/50	8/50		0.38[0.11,1.33]
20.3.2 at 5 days				
Liu 2012	1/50	4/50		0.25[0.03,2.16]
20.3.3 at 1 week				
Liu 2012	2/50	3/50		0.67[0.12,3.82]
20.3.4 at 2 weeks				
Liu 2012	0/50	1/50		0.33[0.01,7.99]
			haloperidol	comparison

Analysis 20.4. Comparison 20 HALOPERIDOL vs COMBINATIONS: e. HALOPERIDOL + RISPERIDONE, Outcome 4 Adverse effects: 1a. Specific - arousal level - i. various.

Study or subgroup	Haloperidol n/N	Comparison n/N	Risk Ratio M-H, Fixed, 95% CI	Risk Ratio M-H, Fixed, 95% CI
20.4.1 insomnia during 14 days				
Liu 2012	6/50	5/50		1.2[0.39,3.68]
20.4.2 somnolence during 14 days				
Liu 2012	11/50	12/50		0.92[0.45,1.88]
haloperidol 0.01 0.1 1 10 100 comparison				

Analysis 20.5. Comparison 20 HALOPERIDOL vs COMBINATIONS: e. HALOPERIDOL + RISPERIDONE, Outcome 5 Adverse effects: 1b. Specific - movement disorders - i. various.

Study or subgroup	Haloperidol n/N	Comparison n/N	Risk Ratio M-H, Fixed, 95% CI					Risk Ratio M-H, Fixed, 95% CI	
20.5.1 tremor during 14 days									
			haloperidol	0.01	0.1	1	10	100	comparison

Analysis 20.6. Comparison 20 HALOPERIDOL vs COMBINATIONS: e. HALOPERIDOL + RISPERIDONE, Outcome 6 Adverse effects: 1c. Specific - miscellaneous.

Comparison 21. HALOPERIDOL vs COMBINATIONS: f. OLANZAPINE + MAGNESIUM VALPROATE

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Specific behaviour: 1. Agitation - average scores - over 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
1.1 endpoint score at 72 hours (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
1.2 endpoint score at 7 days (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
1.3 endpoint score at 14 days (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2 Global outcome: No improvement	1	64	Risk Ratio (M-H, Fixed, 95% CI)	1.06 [0.38, 2.95]
3 Mental state: Average scores - i. over 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
3.1 endpoint score at 72 hours (PANSS total, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
3.2 endpoint score at 7 days (PANSS total, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
3.3 endpoint score at 14 days (PANSS total, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
4 Adverse effects: General - over 24 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	Totals not selected
4.1 one or more drug related adverse effects - by 14 days	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]

Analysis 21.1. Comparison 21 HALOPERIDOL vs COMBINATIONS: f. OLANZAPINE + MAGNESIUM VALPROATE, Outcome 1 Specific behaviour: 1. Agitation - average scores - over 24 hours.

Study or subgroup	Haloperidol		Combination		Mean Difference Fixed, 95% CI	Mean Difference Fixed, 95% CI
	N	Mean(SD)	N	Mean(SD)		
21.1.1 endpoint score at 72 hours (PANSS-EC, high=worse)						
Shi 2010	31	10.4 (2.4)	33	10.4 (2.9)		-0.04[-1.33,1.25]
21.1.2 endpoint score at 7 days (PANSS-EC, high=worse)						
Shi 2010	31	8.6 (3.1)	33	6.9 (2.8)		1.61[0.15,3.07]
21.1.3 endpoint score at 14 days (PANSS-EC, high=worse)						
Shi 2010	31	7.6 (2.8)	33	5.8 (2)		1.82[0.61,3.03]
					haloperidol	combination

haloperidol -4 -2 0 2 4 combination

Analysis 21.2. Comparison 21 HALOPERIDOL vs COMBINATIONS: f. OLANZAPINE + MAGNESIUM VALPROATE, Outcome 2 Global outcome: No improvement.

Study or subgroup	Haloperidol		Combination		Risk Ratio M-H, Fixed, 95% CI	Weight	Risk Ratio M-H, Fixed, 95% CI
	n/N	n/N	n/N	n/N			
Shi 2010	6/31	6/33				100%	1.06[0.38,2.95]
Total (95% CI)	31	33				100%	1.06[0.38,2.95]
Total events: 6 (Haloperidol), 6 (Combination)							
Heterogeneity: Not applicable							
Test for overall effect: Z=0.12(P=0.9)							

haloperidol 0.01 0.1 1 10 100 combination

Analysis 21.3. Comparison 21 HALOPERIDOL vs COMBINATIONS: f. OLANZAPINE + MAGNESIUM VALPROATE, Outcome 3 Mental state: Average scores - i. over 24 hours.

Study or subgroup	Haloperidol		Combination		Mean Difference Fixed, 95% CI	Mean Difference Fixed, 95% CI
	N	Mean(SD)	N	Mean(SD)		
21.3.1 endpoint score at 72 hours (PANSS total, high=worse)						
Shi 2010	31	77 (12.7)	33	78.6 (12.7)		-1.58[-7.79,4.63]
				haloperidol		combination

haloperidol -10 -5 0 5 10 combination

Analysis 21.4. Comparison 21 HALOPERIDOL vs COMBINATIONS: f. OLANZAPINE + MAGNESIUM VALPROATE, Outcome 4 Adverse effects: General - over 24 hours.

Comparison 22. HALOPERIDOL vs COMBINATIONS: g. QUETIAPINE + MAGNESIUM VALPROATE

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Specific behaviour: 1a. Agitation - binary measures	1	60	Risk Ratio (M-H, Fixed, 95% CI)	1.17 [0.44, 3.06]
1.1 <25% reduction, no effect (PANSS-EC)	1	60	Risk Ratio (M-H, Fixed, 95% CI)	1.17 [0.44, 3.06]
2 Specific behaviour: 1b. Agitation - average scores - i. over 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
2.1 endpoint score at 3 days (PANSS-EC, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	0.02 [-2.31, 2.35]
2.2 endpoint score at 7 days (PANSS-EC, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	1.16 [-0.98, 3.30]
2.3 endpoint score at 14 days (PANSS-EC, high=worse)	1	60	Mean Difference (IV, Fixed, 95% CI)	1.24 [-0.49, 2.97]

Analysis 22.1. Comparison 22 HALOPERIDOL vs COMBINATIONS: g. QUETIAPINE + MAGNESIUM VALPROATE, Outcome 1 Specific behaviour: 1a. Agitation - binary measures.

Analysis 22.2. Comparison 22 HALOPERIDOL vs COMBINATIONS: g. QUETIAPINE + MAGNESIUM VALPROATE, Outcome 2 Specific behaviour: 1b. Agitation - average scores - i. over 24 hours.

Comparison 23. HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Tranquillisation or asleep	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1.1 not asleep at 2 hours	1	205	Risk Ratio (M-H, Fixed, 95% CI)	1.07 [0.82, 1.39]
1.2 not asleep at 4 hours	1	205	Risk Ratio (M-H, Fixed, 95% CI)	0.82 [0.66, 1.03]
2 Specific behaviour: 1a. Agitation - average scores - i. up to 2 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
2.1 change score at 2 hours (PANSS-EC, high=worse)	1	108	Mean Difference (IV, Fixed, 95% CI)	-0.5 [-3.80, 2.80]
3 Specific behaviour: 1b. Agitation - average scores - i. up to 4 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
3.1 change score at 4 hours (PANSS-EC, high=worse)	1	126	Mean Difference (IV, Fixed, 95% CI)	-2.40 [-5.20, 0.40]
4 Specific behaviour: 1c. Agitation - average scores - iii. up to 24 hours	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
4.1 score at 24 hours (PANSS-EC, high=worse)	2	305	Mean Difference (IV, Fixed, 95% CI)	0.71 [-0.56, 1.98]
5 Specific behaviour: 1d. Agitation - average scores - iv. over 24 hours	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
5.1 score at 3 days (PANSS-EC, high=worse)	2	305	Mean Difference (IV, Fixed, 95% CI)	-0.41 [-1.66, 0.83]
5.2 score at 5 days (PANSS-EC, high=worse)	2	305	Mean Difference (IV, Fixed, 95% CI)	0.14 [-1.24, 1.52]
5.3 endpoint score at 1 week (PANSS-EC, high=worse)	1	100	Mean Difference (IV, Fixed, 95% CI)	-2.56 [-4.57, -0.55]
5.4 endpoint score at 2 weeks (PANSS-EC, high=worse)	1	100	Mean Difference (IV, Fixed, 95% CI)	-0.88 [-2.34, 0.58]
6 Global outcome: No improvement	1		Risk Ratio (M-H, Fixed, 95% CI)	Totals not selected
6.1 at 72 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
6.2 at 5 days	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
6.3 at 1 week	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
6.4 at 2 weeks	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
7 Mental state: 1. Average scores - i. over 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
7.1 change score at 120 hours (PANSS total, high=worse)	1	205	Mean Difference (IV, Fixed, 95% CI)	-3.5 [-7.07, 0.07]
8 Adverse effects: 1. General	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
8.1 overall adverse events during 120 hours	1	205	Risk Ratio (M-H, Fixed, 95% CI)	1.72 [1.29, 2.29]
9 Adverse effects: 2a. Specific - arousal - i. various	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
9.1 insomnia during 5 days (only re-reported if occurred in ≥5%)	1	205	Risk Ratio (M-H, Fixed, 95% CI)	1.54 [0.45, 5.31]
9.2 insomnia during 14 days	1	100	Risk Ratio (M-H, Fixed, 95% CI)	0.5 [0.20, 1.23]
9.3 somnolence during 14 days	1	100	Risk Ratio (M-H, Fixed, 95% CI)	2.75 [0.94, 8.06]
10 Adverse effects: 2b. Specific - cardiovascular - i. various	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
10.1 tachycardia during 120 hours (only reported if occurred in ≥5%)	1	205	Risk Ratio (M-H, Fixed, 95% CI)	1.29 [0.36, 4.66]
11 Adverse effects: 2c. Specific - movement disorders - i. various	2		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
11.1 akathisia during 5 days (only re-reported if occurred in ≥5%)	1	205	Risk Ratio (M-H, Fixed, 95% CI)	1.63 [1.12, 2.38]
11.2 akathisia during 14 days	1	100	Risk Ratio (M-H, Fixed, 95% CI)	2.33 [0.98, 5.58]
11.3 EPS during 5 days (only reported if occurred in ≥5%)	1	205	Risk Ratio (M-H, Fixed, 95% CI)	2.22 [1.52, 3.23]
11.4 tremor during 14 days	1	100	Risk Ratio (M-H, Fixed, 95% CI)	3.2 [1.27, 8.07]
12 Adverse effects: 2d. Specific - miscellaneous	1		Risk Ratio (M-H, Fixed, 95% CI)	Totals not selected
12.1 Dry mouth during 14 days	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
12.2 Constipation during 14 days	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
12.3 Increased salivation during 14 days	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]

Analysis 23.1. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 1 Tranquillisation or asleep.

Analysis 23.2. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 2 Specific behaviour: 1a. Agitation - average scores - i. up to 2 hours.

Analysis 23.3. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 3 Specific behaviour: 1b. Agitation - average scores - i. up to 4 hours.

Analysis 23.4. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 4 Specific behaviour: 1c. Agitation - average scores - iii. up to 24 hours.

Analysis 23.5. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 5 Specific behaviour: 1d. Agitation - average scores - iv. over 24 hours.

Analysis 23.6. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 6 Global outcome: No improvement.

Analysis 23.7. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 7 Mental state: 1. Average scores - i. over 24 hours.

Analysis 23.8. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 8 Adverse effects: 1. General.

Study or subgroup	haloperidol n/N	combination n/N	Risk Ratio M-H, Fixed, 95% CI	Weight	Risk Ratio M-H, Fixed, 95% CI		
Test for overall effect: Z=3.68(P=0)							
	haloperidol	0.01	0.1	1	10	100	combination

Analysis 23.9. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 9 Adverse effects: 2a. Specific - arousal - i. various.

Study or subgroup	haloperidol n/N	combination n/N	Risk Ratio M-H, Fixed, 95% CI	Weight	Risk Ratio M-H, Fixed, 95% CI		
23.9.1 insomnia during 5 days (only reported if occurred in ≥5%)							
Fang 2012	6/101	4/104		100%	1.54[0.45,5.31]		
Subtotal (95% CI)	101	104		100%	1.54[0.45,5.31]		
Total events: 6 (haloperidol), 4 (combination)							
Heterogeneity: Not applicable							
Test for overall effect: Z=0.69(P=0.49)							
23.9.2 insomnia during 14 days							
Liu 2012	6/50	12/50		100%	0.5[0.2,1.23]		
Subtotal (95% CI)	50	50		100%	0.5[0.2,1.23]		
Total events: 6 (haloperidol), 12 (combination)							
Heterogeneity: Not applicable							
Test for overall effect: Z=1.51(P=0.13)							
23.9.3 somnolence during 14 days							
Liu 2012	11/50	4/50		100%	2.75[0.94,8.06]		
Subtotal (95% CI)	50	50		100%	2.75[0.94,8.06]		
Total events: 11 (haloperidol), 4 (combination)							
Heterogeneity: Not applicable							
Test for overall effect: Z=1.84(P=0.07)							
Test for subgroup differences: Chi²=6.04, df=1 (P=0.05), I²=66.88%							
	haloperidol	0.01	0.1	1	10	100	combination

Analysis 23.10. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 10 Adverse effects: 2b. Specific - cardiovascular - i. various.

Study or subgroup	haloperidol n/N	combination n/N	Risk Ratio M-H, Fixed, 95% CI	Weight	Risk Ratio M-H, Fixed, 95% CI		
23.10.1 tachycardia during 120 hours (only reported if occurred in ≥5%)							
Fang 2012	5/101	4/104		100%	1.29[0.36,4.66]		
Subtotal (95% CI)	101	104		100%	1.29[0.36,4.66]		
Total events: 5 (haloperidol), 4 (combination)							
Heterogeneity: Not applicable							
Test for overall effect: Z=0.38(P=0.7)							
	haloperidol	0.01	0.1	1	10	100	combination

Analysis 23.11. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 11 Adverse effects: 2c. Specific - movement disorders - i. various.

Analysis 23.12. Comparison 23 HALOPERIDOL vs COMBINATIONS: h. RISPERIDONE + CLONAZEPAM, Outcome 12 Adverse effects: 2d. Specific - miscellaneous.

Comparison 24. HALOPERIDOL vs COMBINATIONS: i. ZIPRASIDONE + CLONAZEPAM

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Specific behaviour: 1a. Agitation - average scores - i. up to 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
1.1 endpoint score at 12 hours (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2 Specific behaviour: 1b. Agitation - average scores - ii. over 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
2.1 endpoint score at 48 hours (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2.2 endpoint score at 3 days (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2.3 endpoint score at 5 days (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2.4 endpoint score at 7 days (PANSS-EC, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
3 Global outcome: 1. Average scores - i. over 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
3.1 endpoint score at 7 days (CGI-S, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
4 Mental state: 1. Average scores - i. over 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
4.1 endpoint score at 7 days (PANSS total, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
4.2 endpoint score at 7 days (PANSS positive sub-scale, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
4.3 endpoint score at 7 days (PANSS negative sub-scale, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
4.4 endpoint score at 7 days (PANSS general psychopathology sub-scale, high=worse)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
5 Leaving the study early	1		Risk Ratio (M-H, Fixed, 95% CI)	Totals not selected
5.1 any reason	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
5.2 adverse effects	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
5.3 lost at follow-up	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
5.4 non compliance	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]

Analysis 24.1. Comparison 24 HALOPERIDOL vs COMBINATIONS: i. ZIPRASIDONE + CLONAZEPAM, Outcome 1 Specific behaviour: 1a. Agitation - average scores - i. up to 24 hours.

Study or subgroup	Haloperidol		Combination		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
24.1.1 endpoint score at 12 hours (PANSS-EC, high=worse)						
Zhou 2014	35	16.6 (3)	36	16.9 (3.5)		-0.28[-1.8,1.24]
					haloperidol -5 -2.5 0 2.5 5 combination	

Analysis 24.2. Comparison 24 HALOPERIDOL vs COMBINATIONS: i. ZIPRASIDONE + CLONAZEPAM, Outcome 2 Specific behaviour: 1b. Agitation - average scores - ii. over 24 hours.

Study or subgroup	Haloperidol		Combination		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
24.2.1 endpoint score at 48 hours (PANSS-EC, high=worse)						
Zhou 2014	35	12.3 (3.6)	36	13.8 (3.8)		-1.54[-3.26,0.18]
24.2.2 endpoint score at 3 days (PANSS-EC, high=worse)						
Zhou 2014	35	9.8 (2.4)	36	10.2 (2.8)		-0.42[-1.63,0.79]
24.2.3 endpoint score at 5 days (PANSS-EC, high=worse)						
Zhou 2014	35	9.1 (3.5)	36	9.4 (3.3)		-0.22[-1.78,1.34]
24.2.4 endpoint score at 7 days (PANSS-EC, high=worse)						
Zhou 2014	35	7.9 (2.9)	36	8.1 (3.1)		-0.19[-1.57,1.19]
					haloperidol -5 -2.5 0 2.5 5 combination	

Analysis 24.3. Comparison 24 HALOPERIDOL vs COMBINATIONS: i. ZIPRASIDONE + CLONAZEPAM, Outcome 3 Global outcome: 1. Average scores - i. over 24 hours.

Study or subgroup	Haloperidol		Combination		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
24.3.1 endpoint score at 7 days (CGI-S, high=worse)						
Zhou 2014	35	5.4 (0.6)	36	5.4 (0.5)		0[-0.26,0.26]
					haloperidol -100 -50 0 50 100 combination	

Analysis 24.4. Comparison 24 HALOPERIDOL vs COMBINATIONS: i. ZIPRASIDONE + CLONAZEPAM, Outcome 4 Mental state: 1. Average scores - i. over 24 hours.

Analysis 24.5. Comparison 24 HALOPERIDOL vs COMBINATIONS: i. ZIPRASIDONE + CLONAZEPAM, Outcome 5 Leaving the study early.

ADDITIONAL TABLES

Table 1. Other relevant Cochrane reviews

Focus of review	Reference
Completed and maintained reviews	
'As required' medication regimens for seriously mentally ill people in hospital	Douglas-Hall 2015
Benzodiazepines for psychosis-induced aggression or agitation	Gillies 2013
Chlorpromazine for psychosis induced aggression or agitation	Ahmed 2010

Table 1. Other relevant Cochrane reviews *(Continued)*

Clotiapine for acute psychotic illnesses	Berk 2004
Containment strategies for people with serious mental illness	Muralidharan 2006
Droperidol for acute psychosis	Khokhar 2016
Haloperidol plus promethazine for psychosis-induced aggression	Huf 2016
Olanzapine IM or velotab for acutely disturbed/agitated people with suspected serious mental illnesses	Belgamwar 2005
Seclusion and restraint for serious mental illnesses	Sailas 2000
Zuclopenthixol acetate for acute schizophrenia and similar serious mental illnesses	Jayakody 2012
Reviews in the process of being completed	
Risperidone for psychosis induced aggression or agitation	Ahmed 2011
Haloperidol for long term aggression in psychosis	Khushu 2016
Loxapine inhaler for psychosis-induced aggression	Vangala 2012
Quetiapine for psychosis-induced aggression	Wilkie 2012
De-escalation techniques for psychosis-induced aggression	Du 2017

Table 2. Variations on blinding

Description	Study
Explicitly referred to how participants, study personnel and raters were blinded + tested blinding.	Stotsky 1977
Explicitly referred to how participants, study personnel and raters were blinded.	Breier 2002 , Calver 2015 , Nobay 2004
Report that drugs had identical appearance - no further details regarding rater blinding.	Fitzgerald 1969 , Fruensgaard 1977 , Paprocki 1977 , Resnick 1984 , Ritter 1972
Explicitly state that rater was blinded - no reference to participant blinding. Not all study personnel could have been blinded because person preparing drug (not blinded), often administered drug.	Bristol-Myers 2005b
Study personnel and raters blinded, not clear if attempts made to blind participants.	Battaglia 1997 , Battaglia 2002
Raters blinded - not clear if study personnel and participants were blinded.	Dorevitch 1999 , Foster 1997 , Salzman 1991
"Double blind" - no further information regarding blinding of participants, study personnel and raters.	Eli Lilly 2004 , Kewala 1984 , Kinnon 2001d , Man 1973

Table 2. Variations on blinding (Continued)

Described as double-blind, but during oral phase haloperidol described as being in capsule form, and aripiprazole in tablet form, therefore unclear whether drugs had identical appearance. Not explicit if raters were blinded.	Bristol Myers 2004a
Described as double-blind during IM phase - open during oral phase - no further information given regarding blinding.	Reschke 1974
Described as "modified double", whereby staff administering drugs were not blinded, but raters were - not clear if participants were blinded.	Tuason 1986
Single-blind studies - "open label, rater blind".	Baldacara 2011 , Currier 2004 , Lim 2010 , Shu 2010 , Yin 2012
Reported to be single-blind, but unclear who was blind.	Li 2006
Open-label and rater blinding for some outcomes (BPRS).	Taymeeyapradit 2002
Rater blinding for assessment of BPRS, but unclear regarding rater blinding for other outcomes and whether participants and study personnel were blind.	Bailine 1987
Open studies	Brook 2000 , Fang 2012 , Garza-Trevino 1989 , Guo 2007 , Higashima 2004 , Huf 2007
Not stated	Liang 2013 , Liu 2012 , Shi 2010 , Walther 2014 , Zhou 2014

BPRS - Brief Psychiatric Rating Scale

Table 3. Suggested design for a trial

Methods	Allocation: randomised, clearly described, concealed. Blindness: double, described and tested. Duration: 2 weeks.
Participants	Diagnosis: thought to have psychoses. N = 300.* Age: any. Sex: both. History: acutely ill, aggressive.
Interventions	1. Haloperidol IM: dose flexible within recommended limits. N = 150. 2. Benzodiazepine IM: dose flexible within recommended limits. N = 150.
Outcomes	All outcomes are grouped by time: by 30 minutes, up to two hours, up to four hours, up to 24 hours and finally over 24 hours. Tranquillisation or asleep (measured at 30 minutes, 2 hours, 4 hours and 24 hours). Mortality. Specific behaviours - self-harm, including suicide, injury to others, aggression. Global outcomes - overall improvement, use of additional medication, use of restraints/seclusion. Service outcomes - duration of hospital stay, re-admission. Mental state - no clinically important change in general mental state. Adverse effects - clinically important adverse effects. Leaving the study early - why. Economic outcomes.

Table 3. Suggested design for a trial (Continued)

Notes	* Powered to be able to identify a difference of ~20% between groups for primary outcome with adequate degree of certainty.
--------------	---

IM: intramuscular

Table 4. Adverse effects for 1 mg, 5 mg, 15 mg aripiprazole (Bristol-Myers 2005)

AE	Dose	1 mg (N = 57)	5 mg (N = 63)	15 mg (N = 58)
At least 1 AE		28	30	27
AE rated as serious		1	2	0
Tachycardia		3	2	0
Sinus tachycardia		1	0	0
Vomiting		1	0	3
Nausea		0	6	2
Dizziness		4	7	7
Headache		4	11	8
Somnolence		3	5	6
Akathisia		0	2	0
Dystonia		0	0	1
Agitation		0	0	3
Pain at injection site		Not reported	1	1
QTC abnormality		4	4	3

AE: adverse event

Table 5. Global outcome - data for olanzapine 2.5 mg, 5 mg and 7.5 mg (Breier 2001)

Outcome	Dose	Mean	SD	N
Mean dose of study drug	2.5 mg	4	1.5	48
	5 mg	6.9	2.7	45
	7.5 mg	9.8	3.8	46
Mean dose of benzodiazepines	2.5 mg	3.2	1.1	48
	5 mg	2.0	0	45
	7.5 mg	3.0	1.4	46

Table 6. Continuous skewed data rated at 2 hours for olanzapine 2.5 mg, 5 mg and 7.5 mg (Breier 2001)

Rating scale (Mean change at 2 hr)	Dose	Mean	SD	N
Specific behaviour: ACES	2.5 mg	1.3	1.5	48
	5 mg	2.3	1.9	45
	7.5 mg	2.4	1.7	46
Specific behaviour: ABS	2.5 mg	-5.8	5.5	48
	5 mg	-9.0	5.5	45
	7.5 mg	-10.5	5.6	46
Specific behaviour: PANSS-EC	2.5 mg	-5.5	4.6	48
	5 mg	-8.1	5.3	45
	7.5 mg	-8.7	5	46
Mental state: BPRS Total	2.5 mg	-8.2	9.1	48
	5 mg	-10.4	7.5	45
	7.5 mg	-12.0	7	46
Mental state: BPRS Positive	2.5 mg	-1.5	3.1	48
	5 mg	-1.7	2.8	45
	7.5 mg	-2.1	2.9	46

ABS: Agitated Behavior Scale

ACES: Agitation-Calmness Evaluation Scale

BPRS: Brief Psychiatric Rating Scale

PANSS-EC: Positive and Negative Syndrome Scale Excited Component

Table 7. Continuous skewed data for olanzapine 2.5 mg, 5 mg and 7.5 mg rated at 24 hours (Breier 2001)

Rating scale (mean change at 24 hr)	Dose	Mean	SD	N
Specific behaviour: ACES	2.5 mg	0.9	0.8	48
	5 mg	1.1	1.1	45
	7.5 mg	1	1	46
Specific behaviour: ABS	2.5 mg	-5.7	4.2	48
	5 mg	-6.7	5.9	45
	7.5 mg	-7.7	5.8	46

Table 7. Continuous skewed data for olanzapine 2.5 mg, 5 mg and 7.5 mg rated at 24 hours (Breier 2001) *(Continued)*

Specific behaviour:	2.5 mg	-4.9	4.3	48
PANSS-EC	5 mg	-5.5	4.9	45
	7.5 mg	-5.5	4.1	46
Global outcomes:	2.5 mg	-0.3	0.5	48
CGI-S (24 hr)	5 mg	-0.5	0.8	45
	7.5 mg	-0.6	0.7	46
Mental state:	2.5 mg	-8.4	7.4	48
BPRS Total	5 mg	-9.2	7.8	45
	7.5 mg	-9.6	7.5	46
Mental state:	2.5 mg	-1.5	2.3	48
BPRS Positive	5 mg	-2.0	2.6	45
	7.5 mg	-1.9	2.7	46

ABS: Agitated Behavior Scale

ACES: Agitation-Calmness Evaluation Scale

BPRS: Brief Psychiatric Rating Scale

CGI-S: Clinical Global Impression – Severity

PANSS-EC: Positive and Negative Syndrome Scale Excited Component

Table 8. Binary data for olanzapine 2.5 mg, 5 mg, 7.5 mg (Breier 2001)

Dose	2.5 mg	5 mg	7.5 mg
Outcome	N = 48	N = 45	N = 46
Global outcome: > 1 injection	25	17	14
Leaving the study early: lack of efficacy	0	2	0
Adverse effects: EPS	0	0	0
Adverse effects: akathisia	0	2	0
Adverse effects: QT abnormality	0	4	2

EPS: Extrapyramidal Side Effects

Table 9. Continuous skewed data for aripiprazole 1 mg, 5 mg, 15 mg (Bristol-Myers 2005)

Rating scale	Dose	Mean	SD	N
(Mean change 2 hr after 1st injection)				

Table 9. Continuous skewed data for aripiprazole 1 mg, 5 mg, 15 mg (Bristol-Myers 2005) (Continued)

Specific behaviour:	1 mg	0.65	1.64	56
ACES	5 mg	1.01	1.65	62
	15 mg	0.99	1.73	58
Specific behaviour:	1 mg	-5.15	6.8	56
CABS	5 mg	-5.97	6.81	62
	15 mg	-7.04	7.01	58
Global outcomes:	1 mg	3.07	0.98	56
CGI-I	5 mg	2.82	1.10	62
	15 mg	2.66	1.07	58
Global outcomes:	1 mg	-0.63	1.15	56
CGI-S	5 mg	-0.82	1.12	62
	15 mg	-0.99	1.17	58
Mental state:	1 mg	-6.53	9.61	55
BPRS Total	5 mg	-8.16	9.66	61
	15 mg	-8.88	9.89	56
Mental state:	1 mg	-1.20	2.72	55
BPRS Positive	5 mg	-1.47	2.71	61
	15 mg	-1.86	2.82	57

ACES: Agitation-Calmness Evaluation Scale

BPRS: Brief Psychiatric Rating Scale

CABS: CABS - Corrigan Agitated Behavior Scale

CGI-I: Clinical Global Impression - Improvement

CGI-S: Clinical Global Impression - Severity

Table 10. Binary data for 1 mg, 5 mg and 15 mg aripiprazole (Bristol-Myers 2005)

	Dose	1 mg	5 mg	15 mg
Outcome	N	N = 57	N = 63	N = 58
PEC response $\geq$ 40%		21	31	32
Need for rescue medication		11	5	12
Discontinued for any reason		2	3	1
Discontinued due to adverse event		0	0	1

Table 10. Binary data for 1 mg, 5 mg and 15 mg aripiprazole (Bristol-Myers 2005) *(Continued)*

Withdrew consent	2	2	0
Discontinued due to other known cause	0	1	0

PEC: Psychiatric Emergency Centre

APPENDICES

Appendix 1. Previous searches

1.1 Search in 2011

1.1.1 Electronic searches

1.1.1.1 Cochrane Schizophrenia Group Trials Register

We searched the register (1st June 2011) using the phrase:

(*haloperi* or *R-1625* or *haldol* or *alased* or *aloperidi* or *bioperido* or *buterid* or *ceree* or *dozic* or *duraperido* or *fortuna* or *serena* or *serenel* or *seviu* or *sigaperid* or *sylad* or *zafri* in intervention of STUDY) AND (*aggress* or *violen* or *agitat* or *tranq* in title, abstract, index terms of REFERENCE or intervention of STUDY)

This register is compiled by systematic searches of major databases, handsearches and conference proceedings (see [group module](#)).

1.1.2 Searching other resources

1.1.2.1 Reference searching

We inspected references of all included studies for further relevant studies.

1.1.2.2 Personal contact

We attempted to contact the first author of each included study for information regarding unpublished trials.

WHAT'S NEW

Date	Event	Description
7 March 2017	New citation required but conclusions have not changed	Results of update have not changed the overall conclusions of the review; better evidence on potential droperidol use in the acute setting. 'Summary of findings' tables updated.
17 January 2017	New search has been performed	Nine new studies added, contributing with data on 1056 new individuals (total number of individuals in the updated review: 3546), six new comparisons, and six updates on previously existing comparisons.

HISTORY

Protocol first published: Issue 10, 2011

Review first published: Issue 11, 2012

Date	Event	Description
26 May 2016	Amended	Search was updated and 37 new references added to 'Classification Pending References' section of the review.

Date	Event	Description
1 June 2011	Amended	Cochrane Schizophrenia Group Trial Register searched: 48 studies included in the review, 32 studies included in the analyses.

CONTRIBUTIONS OF AUTHORS

Edoardo G Ostinelli - screened and retrieved papers (2016 search) against eligibility criteria, appraised quality of papers, extracted data from papers, entered data into RevMan and analysed data, interpreted data and took the lead in writing the 2016 update of the review.

Melanie J Powney - screened and retrieved papers (original search) against eligibility criteria, appraised quality of papers, extracted data from papers, entered data into RevMan and analysed data, interpreted data and took the lead in writing the protocol and review.

Xue Li - screened and retrieved papers (2016 search) against eligibility criteria, extracted data from papers.

Clive E Adams - screened and retrieved papers against eligibility criteria, extracted data from papers, helped write protocol and review (original search and 2016).

DECLARATIONS OF INTEREST

Melanie J Powney - none known.

Clive E Adams - was a principle investigator on one trial included in this review ([Huf 2007](#)) .

Edoardo G Ostinelli - none known.

Xue Li - at the time of writing the review, Xue was employed by Systematic Review Solutions Ltd, a company that produces sytematic reviews.

SOURCES OF SUPPORT

Internal sources

- Università degli Studi di Milano, Milan, Italy.
Employs review author Edoardo G Ostinelli
- Systematic Review Solutions Ltd, Nottingham, UK.
At the time of writing, employed review author Xue Li
- University of Nottingham, UK.

Training and support for review author Melanie J Brooke-Powney

External sources

- NIHR Cochrane Programme Grant 2011, UK.

Reference number: 10/4001/15

DIFFERENCES BETWEEN PROTOCOL AND REVIEW

We have re-ordered the main outcomes of interest to reflect order of outcomes in [Types of outcome measures](#)

We have clarified the time point of secondary outcome 'not tranquil or not asleep' to differentiate these outcomes from the primary outcome 'not tranquil or asleep'.

We have added the sub-heading 1. Tranquilisation or asleep to categorise the primary outcomes.

INDEX TERMS

Medical Subject Headings (MeSH)

Aggression [*drug effects] [psychology]; Antipsychotic Agents [administration & dosage] [adverse effects]; Dystonia [chemically induced]; Haloperidol [*administration & dosage] [adverse effects]; Hypnotics and Sedatives [administration & dosage] [adverse

effects]; Placebos [therapeutic use]; Psychomotor Agitation [*drug therapy]; Psychotic Disorders [*drug therapy] [psychology]; Randomized Controlled Trials as Topic; Sleep; Tranquilizing Agents [adverse effects] [*therapeutic use]

MeSH check words

Humans