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TABLE OF CONTENTS

HEADER	1
ABSTRACT	1
PLAIN LANGUAGE SUMMARY	2
SUMMARY OF FINDINGS	3
BACKGROUND	7
OBJECTIVES	7
METHODS	7
RESULTS	9
Figure 1.	10
Figure 2.	12
Figure 3.	14
Figure 4.	15
DISCUSSION	16
AUTHORS' CONCLUSIONS	17
ACKNOWLEDGEMENTS	18
REFERENCES	19
CHARACTERISTICS OF STUDIES	21
DATA AND ANALYSES	31
Analysis 1.1. Comparison 1 Tranexamic acid vs placebo, Outcome 1 Mortality.	33
Analysis 1.2. Comparison 1 Tranexamic acid vs placebo, Outcome 2 Mortality scenario analysis, treatment failure if participant missing.	33
Analysis 1.3. Comparison 1 Tranexamic acid vs placebo, Outcome 3 Mortality in trials with low risk of bias.	34
Analysis 1.4. Comparison 1 Tranexamic acid vs placebo, Outcome 4 Mortality in relation to endoscopic therapy.	35
Analysis 1.5. Comparison 1 Tranexamic acid vs placebo, Outcome 5 Mortality in relation to language of publication.	35
Analysis 1.6. Comparison 1 Tranexamic acid vs placebo, Outcome 6 Rebleeding.	36
Analysis 1.7. Comparison 1 Tranexamic acid vs placebo, Outcome 7 Rebleeding scenario analysis, treatment failure if participant missing.	37
Analysis 1.8. Comparison 1 Tranexamic acid vs placebo, Outcome 8 Rebleeding in trials with low risk of bias.	37
Analysis 1.9. Comparison 1 Tranexamic acid vs placebo, Outcome 9 Rebleeding in relation to endoscopic therapy.	38
Analysis 1.10. Comparison 1 Tranexamic acid vs placebo, Outcome 10 Rebleeding in relation to language of publication.	39
Analysis 1.11. Comparison 1 Tranexamic acid vs placebo, Outcome 11 Myocardial infarction, pulmonary embolism and cerebral infarction.	40
Analysis 1.12. Comparison 1 Tranexamic acid vs placebo, Outcome 12 Deep venous thrombosis.	40
Analysis 1.13. Comparison 1 Tranexamic acid vs placebo, Outcome 13 Any thromboembolic event.	40
Analysis 1.14. Comparison 1 Tranexamic acid vs placebo, Outcome 14 Surgery.	41
Analysis 1.15. Comparison 1 Tranexamic acid vs placebo, Outcome 15 Transfusion required.	41
Analysis 2.1. Comparison 2 Tranexamic acid vs cimetidine or lansoprazole, Outcome 1 Mortality.	42
Analysis 2.2. Comparison 2 Tranexamic acid vs cimetidine or lansoprazole, Outcome 2 Bleeding.	42
Analysis 2.3. Comparison 2 Tranexamic acid vs cimetidine or lansoprazole, Outcome 3 Surgery.	42
Analysis 2.4. Comparison 2 Tranexamic acid vs cimetidine or lansoprazole, Outcome 4 Transfusion.	43
ADDITIONAL TABLES	43
APPENDICES	44
WHAT'S NEW	46
CONTRIBUTIONS OF AUTHORS	47
DECLARATIONS OF INTEREST	47
SOURCES OF SUPPORT	47
DIFFERENCES BETWEEN PROTOCOL AND REVIEW	47
INDEX TERMS	48

[Intervention Review]

Tranexamic acid for upper gastrointestinal bleeding

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ABSTRACT

Background

Tranexamic acid reduces haemorrhage through its antifibrinolytic effects. In a previous version of the present review, we found that tranexamic acid may reduce mortality. This review includes updated searches and new trials.

Objectives

To assess the effects of tranexamic acid versus no intervention, placebo or other antiulcer drugs for upper gastrointestinal bleeding.

Search methods

We updated the review by performing electronic database searches (Cochrane Central Register of Controlled Trials (CENTRAL), MEDLINE, EMBASE, Science Citation Index) and manual searches in July 2014.

Selection criteria

Randomised controlled trials, irrespective of language or publication status.

Data collection and analysis

We used the standard methodological procedures of the The Cochrane Collaboration. All-cause mortality, bleeding and adverse events were the primary outcome measures. We performed fixed-effect and random-effects model meta-analyses and presented results as risk ratios (RRs) with 95% confidence intervals (CIs) and used I^2 as a measure of between-trial heterogeneity. We analysed tranexamic acid versus placebo or no intervention and tranexamic acid versus antiulcer drugs separately. To analyse sources of heterogeneity and robustness of the overall results, we performed subgroup, sensitivity and sequential analyses.

Main results

We included eight randomised controlled trials on tranexamic acid for upper gastrointestinal bleeding. Additionally, we identified one large ongoing pragmatic randomised controlled trial from which data are not yet available. Control groups were randomly assigned to placebo (seven trials) or no intervention (one trial). Two trials also included a control group randomly assigned to antiulcer drugs (lansoprazole or cimetidine). The included studies were published from 1973 to 2011. The number of participants randomly assigned ranged from 47 to 216 (median 204). All trials reported mortality. In total, 42 of 851 participants randomly assigned to tranexamic acid and 71 of 850 in the control group died (RR 0.60, 95% CI 0.42 to 0.87; P value 0.007; $I^2 = 0\%$). The analysis was not confirmed when all participants in the intervention group with missing outcome data were included as treatment failures, or when the analysis was limited to trials with low risk of attrition bias. Rebleeding was diagnosed for 117 of 826 participants in the tranexamic acid group and for 146 of 825 participants in the

control group (RR 0.80, 95% CI 0.64 to 1.00; P value 0.07; $I^2 = 49\%$). We were able to evaluate the risk of serious adverse events on the basis of only four trials. Our analyses showed 'no evidence of a difference between tranexamic acid and control interventions regarding the risk of thromboembolic events.' Tranexamic acid appeared to reduce the risk of surgery in a fixed-effect meta-analysis (RR 0.73, 95% CI 0.56 to 0.95), but this result was no longer statistically significant in a random-effects meta-analysis (RR 0.61, 95% CI 0.35 to 1.04; P value 0.07). No difference was apparent between tranexamic acid and placebo in the assessment of transfusion (RR 1.02, 95% CI 0.94 to 1.11; $I^2 = 0\%$), and meta-analyses that compared tranexamic acid versus antiulcer drugs did not identify beneficial or detrimental effects of tranexamic acid for any of the outcomes assessed.

Authors' conclusions

This review found that tranexamic acid appears to have a beneficial effect on mortality, but a high dropout rate in some trials means that we cannot be sure of this until the findings of additional research are published. At the time of this update in 2014, one large study (8000 participants) is in progress, so this review will be much more informative in a few years. Further examination of tranexamic acid would require inclusion of high-quality randomised controlled trials. Timing of randomisation is essential to avoid attrition bias and to limit the number of withdrawals. Future trials may use a pragmatic design and should include all participants with suspected bleeding or with endoscopically verified bleeding, as well as a tranexamic placebo arm and co-administration of pump inhibitors and endoscopic therapy. Assessment of outcome measures in such studies should be clearly defined. Endoscopic examination with appropriate control of severe bleeding should be performed, as should endoscopic verification of clinically significant rebleeding. In addition, clinical measures of rebleeding should be included. Other important outcome measures include mortality (30-day or in-hospital), need for emergency surgery or blood transfusion and adverse events (major or minor).

PLAIN LANGUAGE SUMMARY

Tranexamic acid, an agent that promotes blood clotting, for serious or uncontrolled upper gastrointestinal bleeding

Background

Upper gastrointestinal bleeding is a common reason for emergency hospital admission. The prognosis is serious. Some patients may die as the result of uncontrolled bleeding.

Review question

Tranexamic acid is an antifibrinolytic agent. This drug reduces the breakdown of fibrin; fibrin provides the framework for the formation of a blood clot, which is needed to stop the bleeding. Clinical trials suggest that tranexamic acid could reduce mortality in upper gastrointestinal bleeding.

Study characteristics

This review includes data from eight randomised trials on tranexamic acid. Two trials also assessed antiulcer drugs. Only one trial used additional endoscopic therapy, as the remaining trials were performed before this intervention was introduced into clinical practice.

Key results

These trials found that tranexamic acid appears to have a beneficial effect on mortality, but a high dropout rate in some trials means that we cannot be sure of these findings until additional research is published. Tranexamic acid did not reduce mortality in the trials that included antiulcer drugs or endoscopic therapy. Additional randomised controlled trials are needed before we can determine whether tranexamic acid has a beneficial effect on serious or uncontrolled upper gastrointestinal bleeding.

Quality of the evidence

Many patients who were randomly assigned were subsequently excluded from the assessment. The main source of bias was therefore attrition. The overall quality of the evidence was moderate to low.

SUMMARY OF FINDINGS

Summary of findings for the main comparison. Tranexamic acid vs placebo for upper gastrointestinal bleeding

Tranexamic acid vs placebo for upper gastrointestinal bleeding

Patient or population: patients with upper gastrointestinal bleeding

Settings:

Intervention: tranexamic acid vs placebo

Outcomes	Illustrative comparative risks* (95% CI)		Relative ef- fect (95% CI)	Number of participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	Control	Tranexamic acid vs placebo				
Mortality Clinical Follow-up: median 5 days	Study population		RR 0.6 (0.42-0.87)	1701 (8 studies)	⊕⊕⊕⊖ moderate a,b	
	84 per 1000	50 per 1000 (34-72)				
	Moderate					
	83 per 1000	50 per 1000 (34-71)				
Rebleeding Clinical and endoscopic assessment Follow-up: median 5 days	Study population		RR 0.72 (0.50-1.03)	1651 (7 studies)	⊕⊕⊖⊖ low a,b,c	
	177 per 1000	142 per 1000 (113-177)				
	Moderate					
	196 per 1000	157 per 1000 (125-196)				
Any thromboembolic event Clinical and radiological assessment Follow-up: median 5 days	Study population		RR 1.86 (0.66-5.24)	1095 (4 studies)	⊕⊕⊕⊖ moderate a,b	
	11 per 1000	20 per 1000 (7-57)				
	Moderate					
	10 per 1000	19 per 1000				

	(7-52)			
Surgery Number of participants who underwent surgery Follow-up: median 5 days	Study population	RR 0.61 (0.35-1.04)	1551 (7 studies)	⊕⊕⊕⊖ moderate ^a
	142 per 1000 103 per 1000 (79-135)			
	Moderate			
	154 per 1000 112 per 1000 (86-146)			
Transfusion Number needing blood transfusion Follow-up: median 3 weeks	Study population	RR 1.02 (0.94-1.1)	931 (5 studies)	⊕⊕⊕⊕ very low ^{a,d}
	564 per 1000 558 per 1000 (507-620)			
	Moderate			
	583 per 1000 577 per 1000 (525-641)			

*The basis for the **assumed risk** (e.g. 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.

^aMost trials had high risk of attrition bias.

^bNot possible to evaluate because number of trials was limited.

^cStatistical between-trial heterogeneity approached 50%.

^dThe number of participants who needed transfusion is an indirect measure of bleeding and varies among clinical sites.

Summary of findings 2. Tranexamic acid vs cimetidine or lansoprazole for upper gastrointestinal bleeding

Tranexamic acid vs cimetidine or lansoprazole for upper gastrointestinal bleeding

Patient or population: patients with upper gastrointestinal bleeding

Settings:

Intervention: tranexamic acid vs cimetidine or lansoprazole

Outcomes	Illustrative comparative risks* (95% CI)		Relative ef- fect (95% CI)	Number of participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	Control	Tranexamic acid vs cimetidine or lansopra- zole				
Mortality Number of participants who died Follow-up: median 5 days	Study population		RR 0.91 (0.50-1.64)	720 (2 studies)	⊕⊕⊕⊕ moderate	
	61 per 1000	56 per 1000 (31-101)				
	Moderate					
	48 per 1000	44 per 1000 (24-79)				
Rebleeding Clinical and endoscopic assessment Follow-up: median 5 days	Study population		RR 0.87 (0.64-1.2)	720 (2 studies)	⊕⊕⊕⊕ moderate	a,b
	188 per 1000	166 per 1000 (121-226)				
	Moderate					
	161 per 1000	142 per 1000 (103-193)				
Surgery Number of participants who underwent surgery Follow-up: median 5 days	Study population		RR 0.83 (0.54-1.26)	720 (2 studies)	⊕⊕⊕⊕ moderate	a,b
	139 per 1000	115 per 1000 (79-168)				
	Moderate					
	105 per 1000	87 per 1000 (60-127)				
Transfusion Number of participants who required at least 1 blood transfusion Follow-up: median 5 days	Study population		RR 0.97 (0.78-1.22)	720 (2 studies)	⊕⊕⊕⊕ low	a,b,c
	573 per 1000	579 per 1000 (510-654)				

Moderate	
599 per 1000	605 per 1000 (533-683)

*The basis for the **assumed risk** (e.g. 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.

^aHigh risk of bias based on assessment of attrition.

^bNot possible to evaluate because number of trials was limited.

^cThis outcome measure is a surrogate estimate for bleeding.

BACKGROUND

Description of the condition

Upper gastrointestinal bleeding is a common reason for emergency hospital admission and a common complication in hospitalised patients (Rockall 1995; Blatchford 1997). A systematic review of general, population-based epidemiological studies found that the incidence of upper gastrointestinal bleeding among patients treated with non-steroidal anti-inflammatory drugs was 0.8 per 1000 (Hernandez 2002). The risk of upper gastrointestinal bleeding has been found to increase significantly with age, co-morbidity and use of non-steroidal anti-inflammatory drugs (Yavorski 1995; Paspatis 2000; Ng 2006). About 80% of patients with upper gastrointestinal bleeding will spontaneously stop bleeding, without recurrence (Laine 1994). The highest mortality and morbidity rates are seen in the remaining 20%, who experience recurrent or continued bleeding. Among patients referred to endoscopy for suspected upper gastrointestinal bleeding, 30-day mortality is 10% to 14% (van Leerdam 2003; Barkun 2004; Barkun 2010).

Description of the intervention

Several endoscopic therapies have been found to be effective in clinical trials (Kahi 2005). However, some hospital departments may not have access to acute endoscopy. In other cases, patients may refuse to undergo endoscopy. Identification of drugs that may achieve haemostasis, stabilising patients until endoscopy can be performed, is therefore essential.

How the intervention might work

Tranexamic acid reduces fibrinolysis by slowing down the conversion of plasminogen to plasmin. The resulting reduction in fibrinolysis prevents the breakdown of blood clots, which may result in haemostasis but increased risk of thromboembolic complications. This drug was introduced for menorrhagia in 1968 (Vermeylen 1968) and is used to reduce blood loss during surgery (Laupacis 1997; Cid 2005). A large multi-centre trial found that tranexamic acid reduces mortality in individuals with bleeding trauma by 9% and results in no apparent increase in thromboembolic events (CRASH-2). It is possible that a similar effect can be achieved in cases of upper gastrointestinal bleeding.

Why it is important to do this review

Endoscopic therapy and proton pump inhibitors serve as the cornerstone in the treatment of bleeding from peptic ulcers (Lau 2013). These treatments are highlighted in recent evidence-based guidelines (Dworzynski 2012), which do not recommend tranexamic acid for the management of upper gastrointestinal bleeding. Randomised trials have assessed the effects of tranexamic acid among patients with suspected or verified upper gastrointestinal bleeding (Cormack 1973; Biggs 1976; Engquist 1979; Bergqvist 1980; Barer 1983; Holstein 1987; Hawkey 2001). A meta-analysis of these trials revealed that tranexamic acid reduces the risks of rebleeding and mortality (Henry 1989). However, results of individual trials varied considerably. Furthermore, the overall result has been characterised as disproportionately skewed by inclusion of a trial in which mortality in the control group was surprisingly high (Barer 1983; Palmer 2002). We have previously published a systematic review on tranexamic acid versus placebo (Gluud 2008). A randomised trial on tranexamic

acid for upper gastrointestinal bleeding was published after our previous meta-analysis had been completed (Bagnenko 2011). This trial was included in a recent review, which determined that we still have insufficient evidence for definitive conclusions (Manno 2014). We performed this updated systematic review on tranexamic acid versus placebo, cimetidine or lansoprazole for upper gastrointestinal bleeding.

OBJECTIVES

To assess the effects of tranexamic acid versus no intervention, placebo or other antiulcer drugs for upper gastrointestinal bleeding.

METHODS

Criteria for considering studies for this review

Types of studies

Randomised trials, irrespective of language, blinding, length of follow-up or publication status.

Types of participants

Individuals with suspected or endoscopically verified upper gastrointestinal bleeding, irrespective of the bleeding source.

Types of interventions

Primary analyses included trials on tranexamic acid versus placebo or no intervention. Secondary analyses compared tranexamic acid versus any other antiulcer drug.

Types of outcome measures

Primary outcomes

- Mortality.
- Adverse events. We defined serious adverse events as all adverse events considered serious by study participants or investigators.

Secondary outcomes

- Rebleeding.
- Surgery.

Search methods for identification of studies

Electronic searches

Electronic searches of the following were performed July 2014.

- Cochrane Central Register of Controlled Trials (CENTRAL) (Appendix 1).
- MEDLINE via Ovid SP (Appendix 2).
- EMBASE via Ovid SP (Appendix 3).
- Science Citation Index Expanded (Appendix 4).

Searching other resources

We scanned conference proceedings and reference lists from relevant trials, wrote to authors of included trials and searched the International Clinical Trials Registry Platform (ICTRP) to identify trials on tranexamic acid (Appendix 5).

Data collection and analysis

Selection of studies

Three review authors (LLG, SLK and CB) screened search results for potentially eligible trials and identified trials that were eligible for inclusion. Excluded trials were listed together with the reasons for exclusion. At each stage of the selection process, at least two review authors independently reviewed search results and selected trials for inclusion. Three review authors (LLG, SLK and CB) agreed on the final list.

Data extraction and management

Two review authors (LLG and SLK) independently extracted data using data collection forms designed to capture information specific to this review. CB verified the extracted data at this update. Disagreements were resolved through discussion before analyses were performed. Data on baseline participant characteristics (inclusion criteria, mean age, proportion of men and source of bleeding), dose and duration of treatment, country of origin, publication status, funding, duration of follow-up and risk of bias were gathered from the included trials and from correspondence with study authors. For trials published in Russian, two review authors (LLG and CB) extracted data obtained by machine translation (Google translate). Dimitrinka Nikolova from the Cochrane Hepato-Biliary Group read the original Russian language publication and verified the extracted data.

Assessment of risk of bias in included studies

At least two review authors (LLG and SLK or CB) independently assessed risk of bias in the included studies by using the risk of bias assessment tool provided in Chapter 8 of the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011). We compared these evaluations and discussed and resolved inconsistencies.

We rated the following domains separately for each of the included studies as 'low risk of bias,' 'high risk of bias' or 'unclear risk of bias' if the risk of bias was uncertain or unknown. These assessments are reported in the 'Risk of bias' table for each individual study included in the [Characteristics of included studies](#) section of the review.

- Allocation sequence was adequately generated ('sequence generation').
- Allocation was adequately concealed ('allocation concealment').
- Knowledge of allocated interventions was adequately prevented during the study ('blinding') (whether the trial was described as double-blind or single-blind, and whether blinding involved healthcare providers, outcome assessors or those performing data extraction or data analysis).
- Incomplete outcome data were adequately addressed.
- Reports of the study were free of suggestions of selective outcome reporting (whether clinically relevant outcome measures were defined and reported).
- The study was apparently free of other sources of bias that could put it at high risk of bias (e.g. potential conflicts of interest, pharmaceutical funding/support, or both).
- Other biases (sample size calculations and registration in clinical trial databases).

We categorised and reported the overall risk of bias of each of the included studies according to the following.

- Low risk of bias (plausible bias unlikely to seriously alter the results) if all criteria were met.
- Unclear risk of bias (plausible bias that raises some doubt about the results) if one or more criteria were assessed as unclear.
- High risk of bias (plausible bias that seriously weakens confidence in the results) if one or more criteria were not met.

We reported these assessments in the [Risk of bias in included studies](#) section of this review.

Measures of treatment effect

The effect measure consisted of risk ratios (RRs) with 95% confidence intervals (CIs).

Unit of analysis issues

As the primary outcome measure was mortality, only trials using a parallel-group design were included. Data on all intervention groups were analysed separately for all trials, including those with more than two parallel arms. Risk of selection bias was noted in the allocation of participants to intervention or control groups and in the administration of collateral interventions; therefore we did not include cluster-randomised trials.

Multi-armed trials

For trials with multiple intervention groups, we partitioned participants into individual allocation arms to perform pair-wise comparisons. For example, a three-arm trial with 30 participants (10 in each arm) allocated to tranexamic acid versus placebo versus antiulcer drugs would allow two pair-wise comparisons of tranexamic acid versus placebo (10 vs 10 participants) and tranexamic acid versus antiulcer drugs (10 vs 10 participants).

Dealing with missing data

We contacted study investigators to request data on all randomly assigned participants to perform intention-to-treat analyses. We used simple imputation to evaluate the potential influence of missing data: imputing failures, imputing successes and worst- and best-case scenarios (Higgins 2008).

Assessment of heterogeneity

We assessed clinical heterogeneity by examining trial conditions on the basis of characteristics of included trials, participants and interventions. We assessed statistical heterogeneity by using I^2 statistical values and reported heterogeneity as important when the I^2 statistic was $> 60\%$ (Higgins 2011).

Assessment of reporting biases

We attempted to obtain trial protocols to compare reported outcome measures in the protocol versus those in the published trial. For analyses with at least 10 trials, we planned to assess reporting biases and other dissemination biases by using funnel plots (Higgins 2011) and to perform regression analyses by using Harbord's modified test (Harbord 2006). Our analyses included only eight trials; therefore we did not carry out a statistical analysis of reporting bias.

Data synthesis

We performed analyses in [RevMan 2014](#) (The Nordic Cochrane Centre, Copenhagen, Denmark), STATA version 13 (Stata Corp., Texas, USA) and Trial Sequential Analysis (The Cochrane Hepato-Biliary Group, Copenhagen, Denmark). We performed all meta-analyses using both random-effects and fixed-effect models. Fixed-effect model meta-analyses are reported only when the results of the two models differ (e.g. one model shows no difference between interventions and the other shows an intervention effect).

We report the results of analyses with the total number of participants. When it was possible to calculate an effect size, we report this with 95% confidence intervals. When the calculated effect size was statistically significant (defined as P value < 0.05), we state whether the result favoured the intervention or control condition.

Subgroup analysis and investigation of heterogeneity

We performed separate analyses of trials on tranexamic acid versus placebo or no intervention and trials on tranexamic acid versus antiulcer drugs, as well as subgroup analyses of trials with low risk of bias based on assessment of the separate domains. We also analysed subgroups of trials that used endoscopic therapy and trials published in English or Russian.

Sensitivity analysis

We performed two further analyses to determine the effects of missing outcome data and a per-protocol analysis to evaluate the influence of missing data. In the first scenario, all participants in the intervention arm with missing outcome data were included as treatment failures and participants in the control group with missing outcome data were considered as treatment successes. In the per-protocol analyses, we excluded participants with missing outcome data. We performed a post hoc analysis that excluded a trial with a very high event rate in the control group ([Barer 1983](#)).

Trial sequential analysis

We performed a post hoc trial sequential analysis to determine the risk of bias associated with cumulative testing and to evaluate futility in the assessment of mortality and bleeding ([Higgins 2008](#); [Wetterslev 2008](#)). We performed the analysis with power set to 80%, alpha to 5% and model-based diversity and with relative risk reduction (RRR) to 30%. We set the control incidence to 10% for the analysis of mortality and to 30% for the analysis of bleeding.

'Summary of findings' tables

We prepared a 'Summary of findings' table ([Guyatt 2008](#)) using GRADEpro software ([Gradepro 3.6](#)) and we included information on the results of our primary outcomes in relation to risk of heterogeneity, duration of follow-up and quality of the evidence.

RESULTS

Description of studies

Results of the search

The original electronic searches identified 84 hits, and the updated search 37 hits ([Appendix 6](#)). The manual search identified one additional record. After reading the titles and abstracts, we retrieved 15 records for further assessment. One record referred to an ongoing trial ([ISRCTN11225767](#)) on tranexamic acid for gastrointestinal bleeding ([Characteristics of ongoing studies](#)). We will include this trial in future updates if the results become available ([Figure 1](#)). We excluded four records that did not assess tranexamic acid for upper gastrointestinal bleeding. In total, we included eight randomised controlled trials in our analyses ([Cormack 1973](#); [Biggs 1976](#); [Engquist 1979](#); [Bergqvist 1980](#); [Barer 1983](#); [Holstein 1987](#); [Hawkey 2001](#); [Bagnenko 2011](#)). We received additional information about study design and outcomes from the primary investigators of one of the included trials ([Hawkey 2001](#)). For the remaining trials, we had access only to published data. The included trials were published as full paper articles, from 1973 to 2011. One trial was published in Russian ([Bagnenko 2011](#)), and the remaining in English.

Figure 1. Study flow diagram.

Included studies

Design

All trials were randomised, parallel arm. One trial was open (Bagnenko 2011), and the remaining trials were double blind with a placebo control.

Sample sizes

The numbers of participants randomly assigned ranged from 47 to 216 (median 204).

Setting

All trials were performed at hospitals.

Participants

The trials included participants admitted with suspected upper gastrointestinal bleeding confirmed by endoscopy or clinically through gastric lavage, hematemesis or melena. Participants with previous or ongoing thromboembolic disease or renal disease and pregnant women were excluded from the trials. Three trials included only participants with severe bleeding (Engquist 1979; Bergqvist 1980; Bagnenko 2011). In the remaining trials, proportions of participants with circulatory involvement ranged from 1% to 21%. Mean participant age ranged from 56 to 62 years in the tranexamic acid groups and from 56 to 65 years in the control groups. Five trials reported that a proportion of participants had oesophageal varices (mean proportion 8%, range 5% to 16%). In four trials (Barer 1983; Holstein 1987; Hawkey 2001; Bagnenko 2011), participants underwent endoscopy within 24 hours after admission. One trial reported that 12% of included participants did not undergo the planned endoscopy. Two trials evaluated participants with endoscopy but did not specify the time frame (Engquist 1979; Bergqvist 1980). The remaining two trials did not include an endoscopic evaluation (Cormack 1973) or had access to endoscopy only during a portion of the trial (Biggs 1976).

Interventions

Tranexamic acid was administered intravenously in one trial (Barer 1983) and orally in three trials (Cormack 1973; Bergqvist 1980; Hawkey 2001). The remaining trials used intravenous followed by oral administration. The total daily dose of tranexamic acid ranged from 4 to 8 g. Duration of therapy ranged from two to seven days.

Comparisons

One trial included a no intervention control group (Bagnenko 2011), and the remaining trials included a placebo control. Two trials were multi-armed and included control groups randomly assigned to the histamine receptor (H₂) agonist cimetidine (Barer

1983) or the proton pump inhibitor lansoprazole alone or with tranexamic acid (Hawkey 2001). One trial allowed co-intervention with the histamine receptor agonist famotidine (Bagnenko 2011). In five trials, a variety of co-interventions, including novaluzide and cimetidine or ranitidine, were administered to participants in both treatment arms (Cormack 1973; Biggs 1976; Engquist 1979; Bergqvist 1980; Holstein 1987). Two trials used endoscopic therapy to control bleeding (Hawkey 2001; Bagnenko 2011).

Outcomes

One trial reported endoscopically verified rebleeding (Bagnenko 2011). Another trial reported rebleeding as assessed by hematemesis, melena or hypotension plus a drop in haemoglobin or rebleeding seen at endoscopy (Hawkey 2001). Two trials defined rebleeding on the basis of a drop in haemoglobin (Holstein 1987) or a drop in haemoglobin, hematemesis or melena (Barer 1983). The remaining trials did not define rebleeding.

Excluded studies

We excluded four trials that were published as full paper articles (Hollanders 1982; Tam 1989; Adachi 2001; Sabovic 2003). These trials were excluded because they were observational, assessed certexate or did not include participants with upper gastrointestinal bleeding (Excluded studies).

Risk of bias in included studies

Allocation

One trial (Bagnenko 2011) reported an adequate method for allocation sequence generation (Figure 2). The remaining trials did not describe how the allocation sequence was generated. One trial did not describe how allocation was concealed (Bagnenko 2011). In the remaining trials, allocation was adequately concealed through double-blind administration of the intervention or placebo.

Figure 2. 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 (performance bias and detection 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
Bagnenko 2011	?	?	-	-	-	+	+	+
Barer 1983	+	+	+	+	+	?	+	+
Bergqvist 1980	?	+	+	+	+	-	-	+
Biggs 1976	?	+	+	+	+	?	+	+
Cormack 1973	?	+	+	+	+	?	+	+
Engquist 1979	?	+	+	+	+	-	+	+
Hawkey 2001	?	+	?	+	+	+	+	+
Holstein 1987	?	+	+	+	+	-	+	+

Blinding

One trial was open (Bagnenko 2011). The remaining trials were double blind with a placebo control.

Incomplete outcome data

Attrition was one of the main sources of bias. Five trials reported losses to follow-up (Engquist 1979; Bergqvist 1980; Barer 1983; Holstein 1987; Hawkey 2001). Three trials gave the impression

that no dropouts or withdrawals had occurred, although this was not specifically reported (Cormack 1973; Biggs 1976; Bagnenko 2011). One in five participants (20%) were withdrawn or excluded after randomisation. Reasons for exclusion included lack of verified bleeding, malignant disease and terminal illness, or the treatment was administered too late.

Selective reporting

One trial did not report bleeding and was classed as having high risk of reporting bias (Bergqvist 1980).

Other potential sources of bias

None of the trials had other potential sources of bias.

Effects of interventions

See: [Summary of findings for the main comparison Tranexamic acid vs placebo for upper gastrointestinal bleeding](#); [Summary of findings 2 Tranexamic acid vs cimetidine or lansoprazole for upper gastrointestinal bleeding](#)

Comparison 1. Tranexamic acid versus placebo

Primary outcomes

Mortality

All trials reported mortality (Analysis 1.1). Forty-two of 851 participants randomly assigned to tranexamic acid and 71 of 850 in

the control group died. The fixed-effect meta-analysis showed that tranexamic acid reduced mortality (RR 0.60, 95% CI 0.42 to 0.87; P value 0.007). No statistical heterogeneity was noted between trials ($I^2 = 0\%$).

Sensitivity analyses and trial sequential analysis

The analysis was not confirmed in a scenario analysis in which all participants with missing outcome data were included as treatment failures (Analysis 1.2), but it was confirmed in a per-protocol analysis (data not shown).

Analyses of studies with low risk of bias confirmed that attrition was the main source of bias (Analysis 1.3). The single remaining trial with low risk of attrition bias (Hawkey 2001) found no effects of the intervention. A similar result was seen when trials that used endoscopic therapy to control bleeding were analysed (Analysis 1.4). No difference (test for subgroup differences P value 0.67) was noted between trials published in English and those translated from Russian (Analysis 1.5).

One of the trials has been criticised for reporting a high control group event rate (Barer 1983; Palmer 2002). This trial included 516 participants and reported a large weight in the analysis (49%). In a post hoc analysis that excluded this trial, tranexamic acid was seen to have no effect on mortality (RR 0.73, 95% CI 0.45 to 1.19; analysis not shown). In our post hoc trial sequential analysis (Figure 3), the required information size was not met, suggesting that the meta-analysis is inconclusive.

Figure 3. Trial sequential analysis of eight trials on tranexamic acid versus placebo or no intervention. Outcome measure is mortality. Analysis was performed with alpha 5% and power 80%. Model-based heterogeneity correction was 0%, relative risk reduction 30% and control group incidence 10%. Graph shows the Z-curve, which is the cumulative result of analysis with trials added according to year of publication. Horizontal line represents 'traditional' 5% level of significance, and inward sloping red line shows trial sequential monitoring boundary. Vertical line represents required information size. Analysis shows that the Z-curve crosses the trial sequential monitoring boundary, suggesting that the result of the meta-analysis is confirmed when analysis is adjusted for cumulative testing. Total number of included participants (N = 1701) is only 62% of required information size (N=2714). The meta-analysis therefore remains inconclusive.

Bleeding

Seven trials reported rebleeding ([Analysis 1.6](#)). In total, rebleeding was diagnosed for 117 of 826 participants in the tranexamic acid group and for 146 of 825 participants in the control group.

Sensitivity analyses and trial sequential analysis

The difference was not statistically significant (RR 0.72, 95% CI 0.50 to 1.03; P value 0.07). A similar conclusion was reached in worst-

case scenario analysis ([Analysis 1.7](#)) and per-protocol analysis (data not shown). Analyses of trials with low risk of bias found no effects of tranexamic acid on bleeding ([Analysis 1.8](#)), and no differences were noted between trials stratified according to use of endoscopic therapy ([Analysis 1.9](#)) or language ([Analysis 1.10](#)). In the trial sequential analysis ([Figure 4](#)), 95% of the required information size was reached. This analysis suggested that the meta-analysis was inconclusive.

Figure 4. Trial sequential analysis of seven trials on tranexamic acid versus placebo or no intervention. Outcome measure is bleeding. Analysis was performed with alpha 5% and power 80%. Model-based heterogeneity correction was 53%, relative risk reduction 30% and control group incidence 10%. Graph shows the Z-curve, which is the cumulative result of analysis with trials added according to year of publication. Horizontal line represents 'traditional' 5% level of significance, and inward sloping red line shows trial sequential monitoring boundary. Vertical line represents required information size. Analysis shows that the Z-curve does not cross the trial sequential monitoring boundary, and the total number of participants (N = 1651) is 95% of required information size (N = 1734). The meta-analysis therefore remains inconclusive.

Adverse events

Several participants randomly assigned to tranexamic acid experienced abdominal pain, nausea, vomiting and thrombophlebitis at the injection site (Table 1). We were unable to perform meta-analyses on these adverse events because data were not provided for both treatment and control groups.

Three trials (Analysis 1.13) on 1048 participants reported thromboembolic events (Engquist 1979; Barer 1983; Holstein 1987). Among participants randomly assigned to tranexamic acid, two cases of myocardial infarction, two cases of pulmonary embolism and one case of cerebral infarction occurred. In the placebo group, two cases of myocardial infarction and two cases of cerebral infarction were recorded. When data on these serious thromboembolic events were combined, the difference was not statistically significant (RR 1.37, 95% CI 0.36 to 5.28; Analysis 1.11). Six cases of deep venous thrombosis occurred among participants randomly assigned to tranexamic acid compared with two cases

in the placebo group (RR 2.32, 95% CI 0.60 to 8.89; Analysis 1.12). The numbers of participants with any thrombotic event were not significantly different between treatment and control groups (RR 1.86, 95% CI 0.66 to 5.24; Analysis 1.13).

Secondary outcomes

Surgery

Seven studies with 1551 participants reported the numbers of participants who required surgery (Analysis 1.14). Between-trial heterogeneity was important ($I^2 = 63\%$). Tranexamic acid appeared to reduce the risk of surgery in a fixed-effect meta-analysis (RR 0.73, 95% CI 0.56 to 0.95), but this result was no longer statistically significant at the 5% level, when a random-effects meta-analysis was used (RR 0.61, 95% CI 0.35 to 1.04; Analysis 1.14).

Transfusions

We planned to analyse the transfusion requirements of included participants, but we did not identify the necessary data. We therefore performed a post hoc analysis of the total numbers of participants who needed at least one blood transfusion ([Analysis 1.15](#)) and found no apparent differences between tranexamic acid and placebo (RR 1.02, 95% CI 0.94 to 1.11 in 931 participants; five studies; $I^2 = 0\%$).

Comparison 2. Tranexamic acid versus antiulcer drugs (cimetidine or lansoprazole)

Primary outcomes

Mortality and rebleeding

Two trials compared tranexamic acid with cimetidine ([Barer 1983](#)) or lansoprazole ([Hawkey 2001](#)). These trials found no significant effects of tranexamic acid on mortality (RR 0.91, 95% CI 0.50 to 1.64; 720 participants; $I^2 = 0\%$; [Analysis 2.1](#)) or bleeding (RR 0.87, 95% CI 0.64 to 1.20; $I^2 = 0\%$; [Analysis 2.2](#)).

Adverse events

We were unable to perform meta-analyses on adverse events ([Table 1](#)). [Barer 1983](#) reported one case of fatal stroke in the tranexamic acid group and two cases of confusion in the control group. [Hawkey 2001](#) reported no adverse events in the tranexamic acid or control groups.

Surgery and transfusions

Trials comparing tranexamic acid versus cimetidine or lansoprazole ([Barer 1983](#); [Hawkey 2001](#)) found no differences between allocation groups regarding the need for surgery (RR 0.83, 95% CI 0.54 to 1.26; 720 participants; two studies; $I^2 = 3\%$; [Analysis 2.3](#)) or blood transfusion (RR 0.97, 95% CI 0.78 to 1.22; 720 participants; two studies; $I^2 = 64\%$; [Analysis 2.4](#)).

'Summary of findings' tables

As shown in the 'Summary of findings' tables on tranexamic acid versus placebo ([Summary of findings for the main comparison](#)) or cimetidine or lansoprazole ([Summary of findings 2](#)), the quality of the evidence was downgraded to moderate or low because of risk of bias.

DISCUSSION

Summary of main results

This review includes eight randomised controlled trials and a total of 1701 participants with acute upper gastrointestinal bleeding. The meta-analyses found that tranexamic acid appears to have a beneficial effect on mortality, but a high dropout rate in some trials means that we cannot be sure of this until additional research is published. Reduced mortality did not clearly reflect reduced bleeding or surgery. The result of the primary meta-analysis was not stable in analyses that adjusted for risk of attrition bias or in trial sequential analyses. When trials with high risk of attrition bias were removed, only one trial was left ([Hawkey 2001](#)) in the analysis, so although we may need to be cautious about the potential effects of dropouts, the evidence is still fairly strong in favour of tranexamic acid. Likewise, only two of the included trials used endoscopic therapy ([Hawkey 2001](#); [Bagnenko 2011](#)). These trials found no clear effects of tranexamic acid on bleeding was noted and that

attrition bias may affect the assessment of mortality, therefore the combined evidence does not allow any recommendations to be made. Additional randomised controlled trials are needed to determine the effects of tranexamic acid. A large randomised double-blind trial on tranexamic acid for gastrointestinal bleeding is ongoing ([ISRCTN11225767](#)). This trial has started to randomly assign the first participants. Results of this trial are expected to finally confirm or refute the effects of tranexamic acid for individuals with upper gastrointestinal bleeding.

Overall completeness and applicability of evidence

During recent years, standard care and assessment of outcomes have changed considerably ([Dworzynski 2012](#)). Previously, patients were not offered full diagnostic endoscopy or endoscopic interventions that are used as standard care today. It is noteworthy that the trial that is the most favourable towards tranexamic acid was published in 1983 ([Barer 1983](#)). The clinical question today is not whether tranexamic acid is better than placebo, but whether tranexamic acid is better than or may be used in combination with current treatments. One of the included trials found no significant difference between tranexamic acid and lansoprazole when used alone, or when the two treatments were combined ([Hawkey 2001](#)). This trial was not designed to assess clinical outcomes, and its statistical power may well have been too small to detect clinically relevant effects. On the other hand, this trial does suggest that additional research is necessary.

Several reasons may explain why we found that tranexamic acid reduces mortality but not bleeding. One possible explanation is that tranexamic acid may be effective in subgroups of patients with a serious prognosis. One of the included trials assessed the influence of the severity of bleeding on the intervention effect ([Hawkey 2001](#)). This trial included 414 participants with suspected upper gastrointestinal bleeding and asked admitting investigators to classify participant risk as high or low on the basis of their presentation. No specific criteria were used. Overall, high-risk participants were more likely to die or need surgery. The effect of tranexamic acid was not related to risk stratification.

Quality of the evidence

In clinical guidelines on the management of upper gastrointestinal non-variceal bleeding, tranexamic acid may be considered, but it is not recommended as routine therapy ([Palmer 2002](#); [Barkun 2003](#)). This treatment generally is not recommended for variceal bleeding, and in several trials, participants with varices were excluded after randomisation. However, meta-analyses in the present review include participants with upper gastrointestinal bleeding due to oesophageal varices. No significant association was found between treatment effect and the proportion of participants with varices. These analyses are only hypothesis generating but suggest that the effects of tranexamic acid on mortality may also be seen in this patient group. Likewise, we found no significant differences between trials in which participants received tranexamic acid in a daily dose of 12 g for two days and those in which the dose was 4 g for seven days. It may be interesting to determine whether dose or treatment duration is related to the risk of thromboembolic events, but we did not have sufficient data to analyse this question.

Vested interests were reported in some of the included trials. We were unable to identify specific design features that introduced bias due to vested interests and the fact that we included only eight

trials limited the possibility of further analyses. Nevertheless, we cannot exclude that this factor can lead to bias. On the other hand, the trial that received full funding (Hawkey 2001) found no effect of tranexamic acid on any of the outcomes assessed. The largest trials received tranexamic acid and placebo from pharmaceutical companies (Barer 1983; Hawkey 2001). The trial with the most positive assessment of tranexamic acid received only medications (Barer 1983).

Mortality is an outcome measure that is relatively stable in relation to performance and ascertainment bias, but not to attrition bias. We found that attrition was the main source of bias. Included trials randomly assigned participants early, and many excluded participants after the source of bleeding had been identified by endoscopy. Exclusion of participants appeared to be done before blinding was broken, although this is not specifically stated. As the result of risk of bias and clinical heterogeneity between trials, which is inevitable given that trials were conducted over several decades, the overall quality of the evidence was deemed moderate or low (Summary of findings for the main comparison; Summary of findings 2).

Potential biases in the review process

Potential biases in the review process are limited. In particular, we attempted to avoid bias in identification and selection of trials, but one possible limitation is that despite exhaustive searching, it is possible that other trials have been conducted, and although we searched extensively, we did not identify unpublished trials or trials published in abstract form. We included only one paper that was published in Russian (Bagненко 2011). Remaining trials were published in English. Because the number of identified trials was limited, we were not able to analyse the risk of publication bias. In theory, meta-analyses are observational, which may lead to biases. A written protocol is necessary to determine biases in the review process and in general, the original protocol must be followed. However, we made methodological changes to our protocol that were based on recent evidence and guidelines (Guyatt 2008; Higgins 2008; Higgins 2011). These changes were mainly related to assessment of risk of bias. We also performed post hoc sequential analyses to improve assessment of biases.

Agreements and disagreements with other studies or reviews

Although our review methods did include assessment of all reported adverse events, we did not carry out a separate search for adverse effects; therefore one of the most important limitations of the present review is the fact that information about adverse events was limited. The recent debate on the use of antifibrinolytic drugs highlights the need for valid safety data. In 2006, an observational study of patients undergoing revascularisation was reported (Mangano 2006). This study included 1295 participants who received aprotinin and 822 who received tranexamic acid. Multi-variable analyses found that aprotinin, but not tranexamic acid, significantly increased the risk of renal failure and cardiovascular or cerebral adverse events. Although results support the safety of tranexamic acid, the participant cohort was considerably different from that included in the present review. In particular, the trials included in the present review excluded participants with previous thromboembolic or renal disease. Therefore, it may be argued that one of the most important limitations of the present review is that limited information about adverse events was

available. Although we found no significant association between tranexamic acid and risk of thromboembolic events, our analyses did not have sufficient statistical power to allow clear inferences. Theoretically, tranexamic acid increases the risk of thrombosis due to unopposed fibrin generation. Case reports have associated tranexamic acid with thromboembolic events, which may be fatal (Rydin 1976; Agnelli 1982; Endo 1988; Woo 1989; Taparia 2002). The trials in the present review excluded participants with previous thromboembolic events, although this is not generally accepted as a contraindication in clinical practice. Likewise these trials excluded participants with renal disease. One observational study has assessed the effects of tranexamic acid on patients undergoing dialysis. Tranexamic acid seems relatively safe to use in this patient group. However, the study included only 20 participants, so additional evidence is needed (Sabovic 2003).

AUTHORS' CONCLUSIONS

Implications for practice

This review found no evidence to support or refute the use of tranexamic acid for upper gastrointestinal bleeding, in terms of mortality, bleeding, surgery or transfusion requirements. We found limited information about adverse events. At present, tranexamic acid cannot be recommended for routine clinical practice.

Implications for research

This review found no evidence that tranexamic acid offers benefit to patients with upper gastrointestinal bleeding in terms of mortality, bleeding, surgery or requirement for blood transfusion. At the time of this update in 2014, additional large pragmatic randomised trials seem warranted. We are aware of one large study (8000 participants) that is in progress, so this review will be much more informative in a few years. However, proton pump inhibitors and endoscopic interventions for severe gastrointestinal bleeding, including injection and thermal and mechanical methods such as haemoclips, may serve as alternative effective interventions (Dworzynski 2012).

Further examination of the efficacy and safety of tranexamic acid would involve high-quality randomised controlled trials. The timing of randomisation is essential in avoiding attrition bias and limiting the number of withdrawals. The trial may use a pragmatic design and include all participants with suspected bleeding based on haemoglobin levels, gastric lavage, hematemesis or melena. Alternatively, only participants with endoscopically verified bleeding may be randomly assigned. The pragmatic approach could mean inclusion of participants with concomitant disease who may have higher risk of bleeding. When such participants are excluded from trials, we may overestimate benefits and underestimate adverse events. Therefore the advantage of a pragmatic approach would be that investigators could assess whether the intervention works in these patients, that is, such trials would have high external validity.

Future trials should assess tranexamic acid delivered orally or intravenously over an appropriate length of time. The trials included in this review assessed a daily dose of up to 16 to 42 g. A placebo arm (for comparison with tranexamic acid) should be included, and interventions should be administered in accordance with current treatment recommendations. Accordingly, co-interventions such as proton pump inhibitors and endoscopic

therapy must be offered. Assessment of outcome measures should be clearly defined, and reports of adverse events should be included. Endoscopic examination with appropriate control of severe bleeding should be included, and endoscopic verification of clinically significant rebleeding should be performed. In addition, clinical measures of rebleeding should be included. Other important outcome measures include mortality (30-day

or in-hospital), along with need for emergency surgery, blood transfusion and adverse events (major or minor).

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REFERENCES

References to studies included in this review

Bagnenko 2011 {published data only}

Bagnenko SF, Verbitskii VG. [Antifibrinolytic therapy for the treatment of massive ulcerative gastro-intestinal bleedings]. *Khirurgiia* 2011;**4**:42-6. [PUBMED: 21721282]

Barer 1983 {published data only}

* Barer D, Ogilvie A, Henry D, Dronfield M, Coggon D, French S, et al. Cimetidine and tranexamic acid in the treatment of acute upper-gastrointestinal-tract bleeding. *New England Journal of Medicine* 1983;**308**:1571-5.

Ogilvie AL, Barer D, Dronfield MW. Trial of cimetidine, tranexamic acid and placebo in the management of acute upper gastrointestinal haemorrhage [abstract]. *Gut* 1981;**22**:F20.

Bergqvist 1980 {published data only}

Bergqvist D, Dahlgren S, Hessman Y. Local inhibition of the fibrinolytic system in patients with massive upper gastrointestinal hemorrhage. *Uppsala Journal of Medical Sciences* 1980;**85**:173-8.

Biggs 1976 {published data only}

Biggs JC, Hugh TB, Dodds AJ. Tranexamic acid and upper gastrointestinal haemorrhage - a double-blind trial. *Gut* 1976;**17**:729-34.

Cormack 1973 {published data only}

Cormack F, Chakrabarti RR, Jouhar AJ, Fearnley GR. Tranexamic acid in upper gastrointestinal haemorrhage. *Lancet* 1973;**1**:1207-8.

Engquist 1979 {published data only}

Engqvist A, Brostrom O, von Feilitzen F, Halldin M, Nystrom B, Ost A, et al. Tranexamic acid in massive haemorrhage from the upper gastrointestinal tract: a double-blind study. *Scandinavian Journal of Gastroenterology* 1979;**14**:839-44.

Hawkey 2001 {published data only}

Hawkey GM, Cole AT, McIntyre AS, Long RG, Hawkey CJ. Drug treatments in upper gastrointestinal bleeding: value of endoscopic findings as surrogate end points. *Gut* 2001;**49**:372-9.

Holstein 1987 {published data only}

* von Holstein CC, Eriksson SB, Kallen R. Tranexamic acid as an aid to reducing blood transfusion requirements in gastric and duodenal bleeding. *BMJ (Clinical Research Edition)* 1987;**294**:7-10.

von Holstein CC, Eriksson SB, Kallen R. Tranexamic acid in gastric and duodenal bleeding. *Scandinavian Journal of Gastroenterology. Supplement* 1987;**137**:71-4.

References to studies excluded from this review

Adachi 2001 {published data only}

Adachi K, Suetsugu H, Moriyama N, Kazumori H, Kawamura A, Fujishiro H, et al. Influence of *Helicobacter pylori* infection

and cetraxate on gastric mucosal blood flow during healing of endoscopic mucosal resection-induced ulcers. *Journal of Gastroenterology and Hepatology* 2001;**16**:1211-6.

Hollanders 1982 {published data only}

Hollanders D, Thomson JM, Schofield PF. Tranexamic acid therapy in ulcerative colitis. *Postgraduate Medical Journal* 1982;**58**(676):87-91.

Sabovic 2003 {published data only}

Sabovic M, Lavre J, Vujkovic B. Tranexamic acid is beneficial as adjunctive therapy in treating major upper gastrointestinal bleeding in dialysis patients. *Nephrology, Dialysis, Transplantation* 2003;**18**:1388-91.

Tam 1989 {published data only}

Tam HC, Lee YC, Wong HY, Ng NS. Healing of gastric ulcer in the elderly: a double-blind study of cetraxate versus ranitidine. *Journal of Clinical Gastroenterology* 1989;**11**:183-6.

References to ongoing studies

ISRCTN11225767 {unpublished data only}

Tranexamic acid for the treatment of gastrointestinal haemorrhage: an international randomised, double blind placebo controlled trial. ISRCTN 2012.

Additional references

Agnelli 1982

Agnelli G, Gresele P, De Cunto M, Gallai V, Nenci GG. Tranexamic acid, intrauterine contraceptive devices and fatal cerebral arterial thrombosis. Case report. *British Journal of Obstetrics and Gynaecology* 1982;**89**:681-2.

Barkun 2003

Barkun A, Bardou M, Marshall JK, Nonvariceal Upper GI Bleeding Consensus Conference Group. Consensus recommendations for managing patients with nonvariceal upper gastrointestinal bleeding. *Annals of Internal Medicine* 2003;**139**:843-57.

Barkun 2004

Barkun A, Sabbah S, Enns R, Armstrong D, Gregor J, Fedorak RN, et al. The Canadian Registry on Nonvariceal Upper Gastrointestinal Bleeding and Endoscopy (RUGBE): Endoscopic hemostasis and proton pump inhibition are associated with improved outcomes in a real-life setting. *The American Journal of Gastroenterology* 2004;**99**(7):1238-46. [PUBMED: 15233660]

Barkun 2010

Barkun AN, Bardou M, Kuipers EJ, Sung J, Hunt RH, Martel M, Sinclair P, International Consensus Upper Gastrointestinal Bleeding Conference Group. International consensus recommendations on the management of patients with nonvariceal upper gastrointestinal bleeding. *Annals of Internal Medicine* 2010;**152**:101-13.

Blatchford 1997

Blatchford O, Davidson LA, Murray WR, Blatchford M, Pell J. Acute upper gastrointestinal haemorrhage in west of Scotland: case ascertainment study. *BMJ (Clinical Research Edition)* 1997;**315**:510-4.

Cid 2005

Cid J, Lozano M. Tranexamic acid reduces allogeneic red cell transfusions in patients undergoing total knee arthroplasty: results of a meta-analysis of randomized controlled trials. *Transfusion* 2005;**45**:1302-7.

CRASH-2

CRASH-2 Collaborators. Effect of tranexamic acid in traumatic brain injury: a nested randomised, placebo controlled trial (CRASH-2 Intracranial Bleeding Study). *BMJ (Clinical Research Edition)* 2011;**343**:d3795. [PUBMED: 21724564]

Dworzynski 2012

Dworzynski K, Pollit V, Kelsey A, Higgins B, Palmer K. Management of acute upper gastrointestinal bleeding: summary of NICE guidance. *BMJ (Clinical Research Edition)* 2012;**344**:e3412. [PUBMED: 22695897]

Endo 1988

Endo Y, Nishimura S, Miura A. Deep-vein thrombosis induced by tranexamic acid in idiopathic thrombocytopenic purpura. *JAMA* 1988;**259**:3561-2.

Gradepro 3.6 [Computer program]

The Grading of Recommendations Assessment, Development and Evaluation (GRADE) Working Group. GRADEpro. [Computer program]. Version 3.2 for Windows. Jan Brozek, Andrew Oxman, Holger Schünemann, 2008. The Grading of Recommendations Assessment, Development and Evaluation (GRADE) Working Group, 2008.

Guyatt 2008

Guyatt GH, Oxman AD, Vist GE, Kunz R, Falck-Ytter Y, Alonso-Coello P, et al. GRADE: an emerging consensus on rating quality of evidence and strength of recommendations. *BMJ (Clinical Research Edition)* 2008;**336**(7650):924-6. [PUBMED: 18436948]

Harbord 2006

Harbord RM, Egger M, Sterne JA. A modified test for small-study effects in meta-analyses of controlled trials with binary endpoints. *Statistics in Medicine* 2006;**25**(20):3443-57. [PUBMED: 16345038]

Henry 1989

Henry DA, O'Connell DL. Effects of fibrinolytic inhibitors on mortality from upper gastrointestinal haemorrhage. *BMJ (Clinical Research Edition)* 1989;**298**:1142-6.

Hernandez 2002

Hernandez-Diaz S, Rodriguez LA. Incidence of serious upper gastrointestinal bleeding/perforation in the general population: review of epidemiologic studies. *Journal of Clinical Epidemiology* 2002;**55**:157-63.

Higgins 2008

Higgins JPT, Green S (editors). *Cochrane Handbook for Systematic Reviews of Interventions* Version 5.0.0 [updated February 2008]. The Cochrane Collaboration, 2008. www.cochrane-handbook.org.

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. www.cochrane-handbook.org.

Kahi 2005

Kahi CJ, Jensen DM, Sung JJ, Bleau BL, Jung HK, Eckert G, et al. Endoscopic therapy versus medical therapy for bleeding peptic ulcer with adherent clot: a meta-analysis. *Gastroenterology* 2005;**129**:855-62.

Laine 1994

Laine L, Peterson WL. Bleeding peptic ulcer. *New England Journal of Medicine* 1994;**331**(11):717-27. [PUBMED: 8058080]

Lau 2013

Lau JY, Barkun A, Fan DM, Kuipers EJ, Yang YS, Chan FK. Challenges in the management of acute peptic ulcer bleeding. *Lancet* 2013;**381**(9882):2033-43. [PUBMED: 23746903]

Laupacis 1997

Laupacis A, Fergusson D. Drugs to minimize perioperative blood loss in cardiac surgery: meta-analyses using perioperative blood transfusion as the outcome. The International Study of Peri-operative Transfusion (ISPOT) Investigators. *Anesthesia and Analgesia* 1997;**85**:1258-67.

Mangano 2006

Mangano DT, Tudor IC, Dietzel C. The risk associated with aprotinin in cardiac surgery. *The New England Journal of Medicine* 2006;**354**:353-65.

Manno 2014

Manno D, Ker K, Roberts I. How effective is tranexamic acid for acute gastrointestinal bleeding?. *BMJ (Clinical Research Edition)* 2014;**348**:g1421. [PUBMED: 24535627]

Ng 2006

Ng W, Wong WM, Chen WH, Tse HF, Lee PY, Lai KC, et al. Incidence and predictors of upper gastrointestinal bleeding in patients receiving low-dose aspirin for secondary prevention of cardiovascular events in patients with coronary artery disease. *World Journal of Gastroenterology* 2006;**12**:2923-7.

Palmer 2002

Palmer KR. Non-variceal upper gastrointestinal haemorrhage: guidelines. *Gut* 2002;**Suppl 4**:1-6.

Paspatis 2000

Paspatis GA, Matrella E, Kapsoritakis A, Leontithis C, Papanikolaou N, Chlouverakis GJ, et al. An epidemiological study of acute upper gastrointestinal bleeding in Crete, Greece. *European Journal of Gastroenterology & Hepatology* 2000;**12**:1215-20.

RevMan 2014 [Computer program]

The Nordic Cochrane Centre, The Cochrane Collaboration.
Review Manager (RevMan). Version Version 5.3. Copenhagen:
The Nordic Cochrane Centre, The Cochrane Collaboration, 2014.

Rockall 1995

Rockall TA, Logan RF, Devlin HB, Northfield TC. Incidence of and mortality from acute upper gastrointestinal haemorrhage in the United Kingdom. Steering Committee and members of the National Audit of Acute Upper Gastrointestinal Haemorrhage. *BMJ (Clinical Research Edition)* 1995;**311**:222-6.

Rydin 1976

Rydin E, Lundberg PO. Tranexamic acid and intracranial thrombosis. *Lancet* 1976;**2**:49.

Taparia 2002

Taparia M, Cordingley FT, Leahy MF. Pulmonary embolism associated with tranexamic acid in severe acquired haemophilia. *European Journal of Haematology* 2002;**68**:307-9.

van Leerdam 2003

van Leerdam ME, Vreeburg EM, Rauws EA, Geraedts AA, Tijssen JG, Reitsma JB, et al. Acute upper GI bleeding: did anything change? Time trend analysis of incidence and outcome of acute upper GI bleeding between 1993/1994 and 2000. *The American Journal of Gastroenterology* 2003;**98**(7):1494-9. [PUBMED: 12873568]

Vermeylen 1968

Vermeylen J, Verhaegen-Declercq ML, Fierens F, Verstraete M. A double blind study of the effect of tranexamic acid in essential

menorrhagia. *Bulletin de la Société Royale Belge de Gynécologie et D'Obstétrique* 1968;**38**:385-90.

Wetterslev 2008

Wetterslev J, Thorlund K, Brok J, Gluud C. Trial sequential analysis may establish when firm evidence is reached in cumulative meta-analysis. *Journal of Clinical Epidemiology* 2008;**61**:64-75.

Woo 1989

Woo KS, Tse LK, Woo JL, Vallance-Owen J. Massive pulmonary thromboembolism after tranexamic acid antifibrinolytic therapy. *British Journal of Clinical Practice* 1989;**43**:465-6.

Yavorski 1995

Yavorski RT, Wong RK, Maydonovitch C, Battin LS, Furnia A, Amundson DE. Analysis of 3,294 cases of upper gastrointestinal bleeding in military medical facilities. *American Journal of Gastroenterology* 1995;**90**:568-73.

References to other published versions of this review

Gluud 2008

Gluud LL, Klingenberg SL, Langholz E. Systematic review: tranexamic acid for upper gastrointestinal bleeding. *Alimentary Pharmacology and Therapeutics* 2008;**27**:752-8.

* Indicates the major publication for the study

CHARACTERISTICS OF STUDIES

Characteristics of included studies [ordered by study ID]

Bagnenko 2011

Methods	Randomised trial on tranexamic acid vs no intervention
Participants	<p>Inclusion criteria: patients admitted with endoscopically verified severe upper gastrointestinal bleeding</p> <p>Number of participants randomly assigned: 47 Mean age tranexamic acid/control: 62/64 years Proportion:</p> <ul style="list-style-type: none"> • of men: 62% • with peptic ulcer: 100% • with oesophageal varices: 0% • with massive bleeding: 100%
Interventions	Tranexamic acid 10 mg/kg IV or oral 3 times/d vs no intervention
Outcomes	All-cause mortality, rebleeding and surgery
Endoscopy	Performed within 24 hours of admission
Definition of re-bleeding	Endoscopically verified bleeding

Tranexamic acid for upper gastrointestinal bleeding (Review)

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Bagnenko 2011 (Continued)

Duration of therapy	3 days	
Notes	No funding reported	
<i>Risk of bias</i>		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not described
Allocation concealment (selection bias)	Unclear risk	Not described
Blinding (performance bias and detection bias) All outcomes	High risk	Open
Blinding of participants and personnel (performance bias) All outcomes	High risk	Open
Blinding of outcome assessment (detection bias) All outcomes	High risk	Open
Incomplete outcome data (attrition bias) All outcomes	Low risk	No losses to follow-up
Selective reporting (reporting bias)	Low risk	Mortality and bleeding are reported
Other bias	Low risk	No other apparent biases are reported

Barer 1983

Methods	Randomised trial on tranexamic acid vs cimetidine vs placebo
Participants	<p>Inclusion criteria: patients admitted with upper gastrointestinal bleeding confirmed by observation of haematemesis or melena</p> <p>Number of participants randomly assigned: 516</p> <p>Mean age: 60-63 years</p> <p>Proportion:</p> <ul style="list-style-type: none"> • of men: 65% • with peptic ulcer: 58% • with oesophageal varices: 5% • with massive bleeding: 13%
Interventions	Tranexamic acid 1 g IV 4 times/d then 1 g orally 4 times/d vs cimetidine 400 mg IV 4 times/d then 400 mg orally 4 times/d vs placebo

Barer 1983 (Continued)

Outcomes	All-cause mortality, rebleeding, surgery and adverse events
Endoscopy	Performed within 24 hours of admission
Definition of re-bleeding	Severe hematemesis or fresh melena or a fall in haemoglobin of at least 2 g/dL within 24 hours after the first day of admission
Duration of therapy	7 days
Notes	Kabi Vitrum Ltd and Smith Kline and French supplied medications; details of funding not supplied

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Allocation sequence based on random numbers
Allocation concealment (selection bias)	Low risk	Central packaging of coded drug containers
Blinding (performance bias and detection bias) All outcomes	Low risk	Double blinding (using placebo)
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Participants and personnel blinded (using placebo)
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Outcomes assessed in a blinded manner (using placebo)
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	In total, 99 participants were excluded after randomisation. Reporting of follow-up and handling of missing outcome data are clear
Selective reporting (reporting bias)	Low risk	Mortality and bleeding are reported
Other bias	Low risk	No other apparent biases are reported

Bergqvist 1980

Methods	Randomised trial on tranexamic acid vs placebo
Participants	<p>Inclusion criteria: patients admitted with massive upper gastrointestinal bleeding confirmed by haematemesis or melena and circulatory involvement (criteria not specified) Number of participants randomly assigned: 50</p> <p>Mean age tranexamic acid/placebo: 61/58 years Proportion:</p> <ul style="list-style-type: none"> • of men: 79% • with peptic ulcer: 65%

Tranexamic acid for upper gastrointestinal bleeding (Review)

Bergqvist 1980 (Continued)

- with oesophageal varices: 7%
- with massive bleeding: not reported

Interventions	Tranexamic acid 2 g orally 6 times daily vs placebo
Outcomes	All-cause mortality and surgery
Endoscopy	Performed after admission (time frame not specified)
Definition of re-bleeding	Not described
Duration of therapy	2 days
Notes	No funding reported

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not described
Allocation concealment (selection bias)	Low risk	Blinded administration of tranexamic acid or placebo
Blinding (performance bias and detection bias) All outcomes	Low risk	Double-blind placebo-controlled trial
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Participants and personnel blinded (using placebo)
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Outcomes assessed in a blinded manner (using placebo)
Incomplete outcome data (attrition bias) All outcomes	High risk	Outcomes for participants who were excluded or withdrawn from treatment (14%) are not reported
Selective reporting (reporting bias)	High risk	Continued bleeding or rebleeding is not reported
Other bias	Low risk	No other apparent biases are reported

Biggs 1976

Methods	Randomised trial comparing tranexamic acid vs placebo
Participants	Inclusion criteria: patients admitted with upper gastrointestinal bleeding observed by medical officer or confirmed by gastric lavage or observation of melena Number of participants randomly assigned: 200 Mean age: not reported

Tranexamic acid for upper gastrointestinal bleeding (Review)

Biggs 1976 (Continued)

Proportion:

- of men: 78%
- with peptic ulcer: 54%
- with oesophageal varices: 6%
- with massive bleeding: 21%

Interventions	Tranexamic acid 1 g IV and 1 g orally 4 times/d then 1 g orally 4 times/d vs placebo
Outcomes	All-cause mortality, rebleeding, surgery and adverse events
Endoscopy	Some participants were evaluated with full endoscopy (time frame and number not specified). Early in the trial, endoscopy was unavailable
Definition of re-bleeding	Not described
Duration of therapy	5 days
Notes	Funding from Fauldings Australia Ltd

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not described
Allocation concealment (selection bias)	Low risk	Blinded administration of tranexamic acid or placebo
Blinding (performance bias and detection bias) All outcomes	Low risk	Double blinding (using placebo)
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Participants and personnel blinded (using placebo)
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Outcomes assessed in a blinded manner (using placebo)
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No withdrawals or exclusions are reported
Selective reporting (reporting bias)	Low risk	Mortality and bleeding are reported
Other bias	Low risk	No other apparent biases are reported

Cormack 1973

Methods	Randomised trial comparing tranexamic acid vs placebo
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Tranexamic acid for upper gastrointestinal bleeding (Review)

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Cormack 1973 (Continued)

Participants	<p>Inclusion criteria: patients admitted with upper gastrointestinal bleeding confirmed by haematemesis or melena</p> <p>Number of participants randomly assigned: 150</p> <p>Mean age: not reported</p> <p>Proportion of:</p> <ul style="list-style-type: none"> men: 67% with peptic ulcer: not reported with oesophageal varices: not reported with massive bleeding: 11%
Interventions	Tranexamic acid 1.5 g orally 4 times/d vs placebo
Outcomes	All-cause mortality. Continued bleeding, rebleeding and surgery reported only as a composite outcome
Endoscopy	Not performed
Definition of re-bleeding	Not defined
Duration of therapy	7 days
Notes	Funding from AB Kabi, Stockholm, Sweden

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not described
Allocation concealment (selection bias)	Low risk	Blinded administration of tranexamic acid or placebo
Blinding (performance bias and detection bias) All outcomes	Low risk	Double blinding (using placebo)
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Participants and personnel blinded (using placebo)
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Outcomes assessed in a blinded manner (using placebo)
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No withdrawals or dropouts are reported
Selective reporting (reporting bias)	Low risk	Mortality and bleeding are reported
Other bias	Low risk	No other apparent biases are reported

Engquist 1979

Methods	Randomised trial comparing tranexamic acid vs placebo
Participants	<p>Inclusion criteria: patients admitted with massive upper gastrointestinal bleeding defined as circulatory embarrassment, loss of 1800 mL blood (4 blood units), or haemoglobin concentration lowered by at least 30 g/L</p> <p>Number of participants randomly assigned: 204</p> <p>Mean age tranexamic acid/placebo: 59/56 years</p> <p>Proportion:</p> <ul style="list-style-type: none"> • of men: 78% • with peptic ulcer: 47% • with oesophageal varices: 16% • with massive bleeding: 100%
Interventions	Tranexamic acid 1 g IV 6 times/d then 1.5 g orally 4 times/d vs placebo
Outcomes	Mortality, bleeding, surgery and adverse events
Endoscopy	Performed (time frame not specified)
Definition of re-bleeding	Not defined
Duration of therapy	7 days
Notes	Funding not reported

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not described
Allocation concealment (selection bias)	Low risk	Blinded administration of tranexamic acid or placebo
Blinding (performance bias and detection bias) All outcomes	Low risk	Double blinding (using placebo)
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Participants and personnel blinded (using placebo)
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Outcomes assessed in a blinded manner (using placebo)
Incomplete outcome data (attrition bias) All outcomes	High risk	Outcome measures are not reported for participants who were withdrawn or excluded (27%)
Selective reporting (reporting bias)	Low risk	Mortality and bleeding are reported

Engquist 1979 (Continued)

Other bias	Low risk	No other apparent biases are reported
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Hawkey 2001

Methods	Randomised trial comparing tranexamic acid alone vs lansoprazole vs tranexamic acid and lansoprazole vs placebo
Participants	Inclusion criteria: patients admitted with suspected upper gastrointestinal bleeding Number of participants randomly assigned: 206 Mean age tranexamic acid/control groups: 58/58 years Proportion: <ul style="list-style-type: none"> • of men: 61% • with peptic ulcer: 37% • with oesophageal varices: 14% • with massive bleeding: 16%
Interventions	Tranexamic acid 2 g orally (bolus) then 1 g orally 4 times/d vs lansoprazole 60 mg orally (bolus) then 30 mg orally vs tranexamic acid and lansoprazole vs placebo
Outcomes	All-cause mortality, bleeding-related mortality and surgery. Adverse events are reported without information about intervention arms
Endoscopy	Endoscopy was performed on the day after admission or earlier for 359 of 414 included participants. The number of participants who were not evaluated with endoscopy was 12 for placebo and 21 for tranexamic acid
Definition of re-bleeding	New haematemesis, melena or hypotension plus a drop in haemoglobin or rebleeding seen at endoscopy
Duration of therapy	4 days
Notes	Funding from Lederle Laboratories Additional data received

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not described
Allocation concealment (selection bias)	Low risk	Blinded administration of tranexamic acid or placebo
Blinding (performance bias and detection bias) All outcomes	Unclear risk	Double blinding (using placebo)
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Participants and personnel blinded (using placebo)

Tranexamic acid for upper gastrointestinal bleeding (Review)

Hawkey 2001 (Continued)

Blinding of outcome assessment (detection bias) All outcomes	Low risk	Outcomes assessed in a blinded manner (using placebo)
Incomplete outcome data (attrition bias) All outcomes	Low risk	Although several participants were withdrawn or lost to follow-up, all randomly assigned participants are accounted for
Selective reporting (reporting bias)	Low risk	Mortality and bleeding are reported
Other bias	Low risk	No other apparent biases are reported

Holstein 1987

Methods	Randomised trial comparing tranexamic acid vs placebo	
Participants	Inclusion criteria: patients admitted with suspected (criteria not specified) upper gastrointestinal bleeding. After randomisation, endoscopy was performed, and all patients without a benign gastric or duodenal bleeding source were excluded Number of participants randomly assigned: 128 Mean age tranexamic acid/placebo: 62/65 years Proportion: <ul style="list-style-type: none"> • of men: 70% • with peptic ulcer: 90% • with oesophageal varices: 0% • with massive bleeding: 1% 	
Interventions	Tranexamic acid 1 g IV 6 times/d then 1.5 g orally 4 times/d vs placebo	
Outcomes	Mortality, bleeding, surgery and adverse events	
Endoscopy	Performed within 24 hours of admission	
Definition of re-bleeding	Drop in haemoglobin of at least 20 g/L	
Duration of therapy	6 days	
Notes	KabiVitrum AB supplied tranexamic acid and placebo and gave advice about the trial design	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Not described
Allocation concealment (selection bias)	Low risk	Blinded administration of tranexamic acid or placebo
Blinding (performance bias and detection bias) All outcomes	Low risk	Double blinding (using placebo)

Tranexamic acid for upper gastrointestinal bleeding (Review)

Holstein 1987 *(Continued)*

Blinding of participants and personnel (performance bias) All outcomes	Low risk	Participants and personnel blinded (using placebo)
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Outcomes assessed in a blinded manner (using placebo)
Incomplete outcome data (attrition bias) All outcomes	High risk	Outcome measures are not reported for participants who were excluded or lost to follow-up (47%)
Selective reporting (reporting bias)	Low risk	Mortality and bleeding are reported
Other bias	Low risk	No other apparent biases are reported

IV: intravenous.

Characteristics of excluded studies *[ordered by study ID]*

Study	Reason for exclusion
Adachi 2001	Randomised trial on lansoprazole alone or with cetraxate for healing of peptic ulcers.
Hollanders 1982	Randomised trial on tranexamic acid for ulcerative colitis
Sabovic 2003	Non-randomised trial on tranexamic acid as adjunctive therapy for upper gastrointestinal bleeding in patients with renal failure
Tam 1989	Randomised trial on cetraxate vs ranitidine for treatment of patients with gastric ulcer

Characteristics of ongoing studies *[ordered by study ID]*
ISRCTN11225767

Trial name or title	Tranexamic acid for the treatment of gastrointestinal haemorrhage: an international randomised, double-blind placebo-controlled trial (Haemorrhage ALleviation with Tranexamic acid? IntesTinal system HALT-IT)
Methods	Randomised double-blind trial
Participants	<ul style="list-style-type: none"> Inclusion criteria: all adult patients with acute significant upper or lower gastrointestinal bleeding when the responsible clinician is substantially uncertain as to the appropriateness of antifibrinolytic agents in the patient Exclusion criteria: fundamental eligibility criterion: the responsible clinician's 'uncertainty' as to whether an antifibrinolytic agent should be used in a particular patient with upper or lower gastrointestinal bleeding
Interventions	Tranexamic acid (loading dose 1 g over 10 minutes then infusion of 3 g over 24 hours) or matching placebo
Outcomes	Death in hospital within 28 days of randomisation (cause-specific mortality will also be recorded)

Tranexamic acid for upper gastrointestinal bleeding (Review)

ISRCTN11225767 (Continued)

Starting date	Date of first enrolment: January 2 2013
Contact information	Professor Ian Roberts, haltit@lshtm.ac.uk Ms Haleema Shakur, haltit@lshtm.ac.uk
Target sample size	8000 participants
Sponsor	London School of Hygiene and Tropical Medicine (UK)
Conditions assessed	Upper and lower gastrointestinal bleeding
Notes	Funded by Health Technology Assessment programme, which is part of the National Institute for Health Research (NIHR), UK

DATA AND ANALYSES

Comparison 1. Tranexamic acid vs placebo

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Mortality	8	1701	Risk Ratio (M-H, Random, 95% CI)	0.60 [0.42, 0.87]
2 Mortality scenario analysis, treatment failure if participant missing	8	1701	Risk Ratio (M-H, Random, 95% CI)	0.88 [0.43, 1.81]
3 Mortality in trials with low risk of bias	8		Risk Ratio (M-H, Random, 95% CI)	Subtotals only
3.1 Allocation sequence generation	1	516	Risk Ratio (M-H, Random, 95% CI)	0.46 [0.26, 0.82]
3.2 Allocation concealment	7	1654	Risk Ratio (M-H, Random, 95% CI)	0.61 [0.42, 0.89]
3.3 Blinding	7	1654	Risk Ratio (M-H, Random, 95% CI)	0.61 [0.42, 0.89]
3.4 Incomplete outcome data	1	206	Risk Ratio (M-H, Random, 95% CI)	0.8 [0.22, 2.89]
3.5 Selective reporting	7	1651	Risk Ratio (M-H, Random, 95% CI)	0.60 [0.41, 0.89]
4 Mortality in relation to endoscopic therapy	8	1701	Risk Ratio (M-H, Random, 95% CI)	0.60 [0.42, 0.87]
4.1 Endoscopic therapy not used	6	1448	Risk Ratio (M-H, Random, 95% CI)	0.60 [0.40, 0.88]
4.2 Endoscopic therapy used to control bleeding	2	253	Risk Ratio (M-H, Random, 95% CI)	0.66 [0.22, 2.00]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
5 Mortality in relation to language of publication	8	1701	Risk Ratio (M-H, Random, 95% CI)	0.60 [0.42, 0.87]
5.1 Trials published in English	7	1654	Risk Ratio (M-H, Random, 95% CI)	0.61 [0.42, 0.89]
5.2 Trials translated from Russian	1	47	Risk Ratio (M-H, Random, 95% CI)	0.38 [0.04, 3.38]
6 Rebleeding	7	1651	Risk Ratio (M-H, Random, 95% CI)	0.72 [0.50, 1.03]
7 Rebleeding scenario analysis, treatment failure if participant missing	7	1651	Risk Ratio (M-H, Random, 95% CI)	0.80 [0.51, 1.27]
8 Rebleeding in trials with low risk of bias	7	5581	Risk Ratio (M-H, Random, 95% CI)	0.78 [0.65, 0.93]
8.1 Allocation sequence generation	1	516	Risk Ratio (M-H, Random, 95% CI)	1.16 [0.83, 1.61]
8.2 Allocation concealment	6	1604	Risk Ratio (M-H, Random, 95% CI)	0.73 [0.50, 1.07]
8.3 Blinding	6	1604	Risk Ratio (M-H, Random, 95% CI)	0.73 [0.50, 1.07]
8.4 Incomplete outcome data	1	206	Risk Ratio (M-H, Random, 95% CI)	0.9 [0.38, 2.12]
8.5 Selective reporting	7	1651	Risk Ratio (M-H, Random, 95% CI)	0.72 [0.50, 1.03]
9 Rebleeding in relation to endoscopic therapy	7	1651	Risk Ratio (M-H, Random, 95% CI)	0.72 [0.50, 1.03]
9.1 Endoscopic therapy not used	5	1398	Risk Ratio (M-H, Random, 95% CI)	0.70 [0.45, 1.08]
9.2 Endoscopic therapy used to control bleeding	2	253	Risk Ratio (M-H, Random, 95% CI)	0.77 [0.36, 1.62]
10 Rebleeding in relation to language of publication	7	1651	Risk Ratio (M-H, Random, 95% CI)	0.72 [0.50, 1.03]
10.1 Trials published in English	6	1604	Risk Ratio (M-H, Random, 95% CI)	0.73 [0.50, 1.07]
10.2 Trials translated from Russian	1	47	Risk Ratio (M-H, Random, 95% CI)	0.45 [0.10, 2.11]
11 Myocardial infarction, pulmonary embolism and cerebral infarction	3	1048	Risk Ratio (M-H, Random, 95% CI)	1.37 [0.36, 5.28]
12 Deep venous thrombosis	3	1048	Risk Ratio (M-H, Random, 95% CI)	2.32 [0.60, 8.89]
13 Any thromboembolic event	4	1095	Risk Ratio (M-H, Random, 95% CI)	1.86 [0.66, 5.24]
14 Surgery	7	1551	Risk Ratio (M-H, Random, 95% CI)	0.61 [0.35, 1.04]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
15 Transfusion required	5	931	Risk Ratio (M-H, Random, 95% CI)	1.02 [0.94, 1.11]

Analysis 1.1. Comparison 1 Tranexamic acid vs placebo, Outcome 1 Mortality.

Analysis 1.2. Comparison 1 Tranexamic acid vs placebo, Outcome 2 Mortality scenario analysis, treatment failure if participant missing.

Analysis 1.3. Comparison 1 Tranexamic acid vs placebo, Outcome 3 Mortality in trials with low risk of bias.

Analysis 1.4. Comparison 1 Tranexamic acid vs placebo, Outcome 4 Mortality in relation to endoscopic therapy.

Analysis 1.5. Comparison 1 Tranexamic acid vs placebo, Outcome 5 Mortality in relation to language of publication.

Analysis 1.6. Comparison 1 Tranexamic acid vs placebo, Outcome 6 Rebleeding.

Analysis 1.7. Comparison 1 Tranexamic acid vs placebo, Outcome 7 Rebleeding scenario analysis, treatment failure if participant missing.

Analysis 1.8. Comparison 1 Tranexamic acid vs placebo, Outcome 8 Rebleeding in trials with low risk of bias.

Analysis 1.9. Comparison 1 Tranexamic acid vs placebo, Outcome 9 Rebleeding in relation to endoscopic therapy.

Analysis 1.10. Comparison 1 Tranexamic acid vs placebo, Outcome 10 Rebleeding in relation to language of publication.

Analysis 1.11. Comparison 1 Tranexamic acid vs placebo, Outcome 11 Myocardial infarction, pulmonary embolism and cerebral infarction.

Analysis 1.12. Comparison 1 Tranexamic acid vs placebo, Outcome 12 Deep venous thrombosis.

Analysis 1.13. Comparison 1 Tranexamic acid vs placebo, Outcome 13 Any thromboembolic event.

Analysis 1.14. Comparison 1 Tranexamic acid vs placebo, Outcome 14 Surgery.

Analysis 1.15. Comparison 1 Tranexamic acid vs placebo, Outcome 15 Transfusion required.

Comparison 2. Tranexamic acid vs cimetidine or lansoprazole

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Mortality	2	720	Risk Ratio (M-H, Random, 95% CI)	0.91 [0.50, 1.64]
2 Bleeding	2	720	Risk Ratio (M-H, Random, 95% CI)	0.87 [0.64, 1.20]
3 Surgery	2	720	Risk Ratio (M-H, Random, 95% CI)	0.83 [0.54, 1.26]
4 Transfusion	2	720	Risk Ratio (M-H, Random, 95% CI)	0.97 [0.78, 1.22]

Analysis 2.1. Comparison 2 Tranexamic acid vs cimetidine or lansoprazole, Outcome 1 Mortality.

Analysis 2.2. Comparison 2 Tranexamic acid vs cimetidine or lansoprazole, Outcome 2 Bleeding.

Analysis 2.3. Comparison 2 Tranexamic acid vs cimetidine or lansoprazole, Outcome 3 Surgery.

Analysis 2.4. Comparison 2 Tranexamic acid vs cimetidine or lansoprazole, Outcome 4 Transfusion.

ADDITIONAL TABLES

Table 1. Adverse events

Trial	Serious adverse events			Non-serious adverse events	Non-serious adverse events
	Tranexam- ic acid	Control group	Intervention group not specified	Tranexamic acid	Control group
Barer 1983	Fatal stroke (n = 1)	Confu- sion (n = 1)	Pulmonary embolism (n = 5). Myocardial infarction (n = 8)	None described	Confusion (n = 1)
Biggs 1976	None de- scribed	None de- scribed	None described	Thrombophlebitis at injec- tion site (n = 3). Nausea or headache (n = 4)	Throm- bophlebitis at injection site (n = 2). Nausea or headache (n = 5). Fever (n = 2)
Engquist 1979	Pulmonary embolism (n = 2). My- ocardial in- farction (n = 2)	Cerebral infarction (n = 2)	None described	None described	None described
Hawkey 2001	None de- scribed	None de- scribed	Several participants experienced throm- boembolic complications. The numbers were described as not significantly differ- ent in treatment and control groups, but no specific data are provided	None described	None described
Holstein 1987	None de- scribed	Cerebral infarction (n = 2)	Five additional participants were exclud- ed as the result of thromboembolic dis- ease, but whether these participants were randomly assigned to tranexamic acid or placebo is not reported	Nausea and vomiting, tachycardia (n = 3). Hy- potension (n = 3). Throm- bophlebitis at injection site (n = 2). Deep venous thrombosis (n = 1)	None described

APPENDICES

Appendix 1. CENTRAL search strategy

CENTRAL July 2014

#1 stomach or antrum or antral or pyloric or pylorus or gastri* or epigastri* or duodenal or duodenum or gastro-duodenal or gastroduodenal or oeso*ag* or esp*ag* or upper GI or UGI or upper gastrointestinal

#2 h*emorrhag* or bleed* or re-bleed* or rebleed*

#3 (#1 AND #2)

#4 MeSH descriptor Gastrointestinal Hemorrhage explode all trees

#5 Gastrointestinal Hemorrhage

#6 h*ematemesis

#7 (#3 OR #4 OR #5)

#8 tranexamic acid or amchafibrin or anvitoff or cyklokapron or Espercil or exacyl or lysteda or spotof or t-amcha or tranhexamic acid or transamin or Transcam or ugurol

#9 (#7 AND #6)

Appendix 2. MEDLINE search strategy

Ovid MEDLINE 1950 to July 2014

1. (stomach or antrum or antral or pyloric or pylorus or gastri\$ or epigastri\$ or duodenal or duodenum or gastro-duodenal or gastroduodenal or oeso*ag* or esp*ag* or "upper GI" or UGI or "upper gastrointestinal ").mp. [mp=title, original title, abstract, name of substance word, subject heading word, unique identifier]

2. (h*emorrhag\$ or bleed\$ or re-bleed\$ or rebleed\$).mp. [mp=title, original title, abstract, name of substance word, subject heading word, unique identifier]

3. 1 and 2

4. exp Gastrointestinal Hemorrhage/

5. h*ematemesis.mp. [mp=title, original title, abstract, name of substance word, subject heading word, unique identifier]

6. or/3-5

7. ("tranexamic acid" or amchafibrin or anvitoff or cyklokapron or Espercil or exacyl or lysteda or spotof or t-amcha or "tranhexamic acid" or transamin or Transcam or ugurol).mp. [mp=title, original title, abstract, name of substance word, subject heading word, unique identifier]

8. 6 and 7

9. randomized controlled trial.pt.

10. controlled clinical trial.pt.

11. randomized.ab.

12. placebo.ab.

13. drug therapy.fs.

14. randomly.ab.

15. trial.ab.

16. groups.ab.

17. or/9-16

18. exp animals/ not humans.sh.

19. 17 not 18

20. 8 and 19

Appendix 3. EMBASE search strategy

EMBASE 1980 to July 2014

1. (stomach or antrum or antral or pyloric or pylorus or gastri\$ or epigastri\$ or duodenal or duodenum or gastro-duodenal or gastroduodenal or oeso*ag* or esp*ag* or "upper GI" or UGI).mp. [mp=title, abstract, subject headings, heading word, drug trade name, original title, device manufacturer, drug manufacturer name]

2. (h*emorrhag\$ or bleed\$ or re-bleed\$ or rebleed\$).mp. [mp=title, abstract, subject headings, heading word, drug trade name, original title, device manufacturer, drug manufacturer name]

3. 1 and 2

4. gastrointestinal hemorrhage/

5. duodenum bleeding/ or stomach hemorrhage/ or upper gastrointestinal bleeding/

6. h*ematemesis.mp. [mp=title, abstract, subject headings, heading word, drug trade name, original title, device manufacturer, drug manufacturer name]

7. or/3-6

8. ("tranexamic acid" or amchafibrin or anvitoff or cyklokapron or Espercil or exacyl or lysteda or spotof or t-amcha or "tranhexamic acid" or transamin or Transcam or ugurol).mp. [mp=title, abstract, subject headings, heading word, drug trade name, original title, device manufacturer, drug manufacturer name]

9. 7 and 8

10. Clinical trial/

11. Randomized controlled trial/

12. Randomization/

13. Single-Blind Method/

14. Double-Blind Method/

15. Cross-Over Studies/

16. Random Allocation/

17. Placebo/

18. Randomi?ed controlled trial\$.tw.

19. Rct.tw.

20. Random allocation.tw.

21. Randomly allocated.tw.

22. Allocated randomly.tw.

23. (allocated adj2 random).tw.

24. Single blind\$.tw.

25. Double blind\$.tw.

26. ((treble or triple) adj blind\$.tw.

27. Placebo\$.tw.

28. Prospective study/

29. or/10-28

30. Case study/

31. Case report.tw.

32. Abstract report/ or letter/

33. or/30-32

34. 29 not 33

35. 9 and 34

Appendix 4. Science Citation Index search strategy

7 #6 AND #5

6 TS=(random* or blind* or placebo* or meta-analysis)

5 #4 AND #3

4 TS=(tranexamic acid)

3 #2 AND #1

2 TS=(oeso*ag* or eso*ag* or stomach or gastric or ventricular or duodenum)

1 TS=(bleeding or hemorrhage or haemorrhage or re-bleeding or (recurren* and (bleed* or haemorrhage or hemorrhage)))

Appendix 5. International Clinical Trials Registry Platform (ICTRP) search strategy

ICTRP July 2014

(bleeding OR "BLEEDING" OR "Blood Loss" OR "EXTRAVASATION BLOOD" OR "HAEMORRHAGE NOS" OR "HEM" OR "hemorrhage" OR "HEMORRHAGE (NOS)" OR "HEMORRHAGE NOS" OR "Hemorrhage, unspecified" OR "Hemorrhages" OR "Loss of blood" tranexamic acid OR "Acid, Tranexamic" OR "AMCA" OR "AMCHA" OR "Cyclokapron" OR "Cyklokapron" OR "t-AMCHA" OR "TRANEXAMIC ACID" OR "trans-4-(Aminomethyl)cyclohexanecarboxylic Acid" AND (tranexamic acid OR "Acid, Tranexamic" OR "AMCA" OR "AMCHA" OR "Cyclokapron" OR "Cyklokapron" OR "t-AMCHA" OR "TRANEXAMIC ACID" OR). "trans-4-(Aminomethyl)cyclohexanecarboxylic Acid"))

Appendix 6. Search results

Eighty records were identified by initial electronic searches of the following databases.

- CENTRAL: 10 citations.
- MEDLINE: 22 citations.
- EMBASE: 62 citations.
- Science Citation Index: 6 citations.

Thirty-seven additional records were identified in the search update performed July 2014 ([Figure 1](#)).

WHAT'S NEW

Date	Event	Description
29 July 2014	New search has been performed	We originally planned to perform separate analyses on continued bleeding and rebleeding, but we were unable to extract the necessary data from the included trials

Date	Event	Description
		We have updated the assessment of bias according to the recommendations provided in the <i>Cochrane Handbook for Systematic Reviews of Interventions</i> of The Cochrane Collaboration
29 July 2014	New citation required but conclusions have not changed	One additional randomised controlled trial has been included in the analyses. The addition of this trial has not changed our conclusions.

CONTRIBUTIONS OF AUTHORS

LLG drafted the review and performed the statistical analyses. SLK and ELA participated in interpretation of the results and revision of the review. All review authors have approved the final version.

DECLARATIONS OF INTEREST

None of the review authors has reported any conflicts of interest with regard to the present work.

CB is the proprietor of Systematic Research Ltd and was paid by the CUGPD Editorial Group for her contributions to this review.

SOURCES OF SUPPORT

Internal sources

- The present review did not receive funding, Other.

External sources

- No sources of support supplied

DIFFERENCES BETWEEN PROTOCOL AND REVIEW

We updated the review in 2014 to comply with MECIR standards for the conduct and reporting of systematic reviews. We updated the format of the review for clarity and provided the following information about our methods.

- We included adverse effects as a primary outcome (formerly a secondary outcome) to comply with recent guidance on conduct and presentation of the systematic review.
- We clarified that because of the risk of selection bias in the allocation of participants to intervention or control groups and bias in the administration of collateral interventions, we did not include cluster-randomised trials.
- For trials with multiple intervention groups, we partitioned the numbers of participants in individual allocation arms into pair-wise comparisons.
- We used simple imputation to evaluate the potential influence of missing data: imputing failures, imputing successes, worst- and best-case.
- We provided additional explanations of our assessment and reporting of heterogeneity for clarity; our methods remain unchanged.
- We performed all meta-analyses using both random-effects and fixed-effect models. Fixed-effect meta-analyses are reported only when results of the two models differ (e.g. one model shows no difference between interventions and the other shows an intervention effect).
- We performed separate analyses of trials on tranexamic acid versus placebo or no intervention and trials on tranexamic acid versus antiulcer drugs. We performed subgroup analyses of trials with low risk of bias based on assessment of the separate domains. We also analysed subgroups of trials that used endoscopic therapy and trials published in English or Russian.
- We performed an analysis when all participants with missing outcome data were included as treatment failures, and we used a per-protocol analysis to evaluate the influence of missing data when we excluded participants with missing outcome data.
- For analyses with at least 10 trials, we planned to assess reporting biases and other dissemination biases on the basis of funnel plots (Higgins 2011) and regression analyses by using Harbord's modified test (Harbord 2006). Our analyses included only eight trials; therefore we did not carry out statistical analysis of reporting bias.
- We performed a post hoc trial sequential analysis to evaluate the risk of bias associated with cumulative testing and to evaluate futility in assessment of mortality and bleeding (Higgins 2008; Wetterslev 2008). We performed the analysis with power set to 80%, alpha to 5%, model-based diversity and relative risk reduction (RRR) to 25%. We set the control group incidence to 8% in our analysis of mortality and to 18% in our analysis of bleeding.

- We prepared a 'Summary of findings' table ([Guyatt 2008](#)) using GRADEpro software ([Gradepro 3.6](#)) and included information on results of our primary outcomes in relation to risk of heterogeneity, duration of follow-up and quality of the evidence.

INDEX TERMS

Medical Subject Headings (MeSH)

Administration, Oral; Aluminum Hydroxide [therapeutic use]; Anti-Ulcer Agents [therapeutic use]; Antifibrinolytic Agents [adverse effects] [*therapeutic use]; Cimetidine [therapeutic use]; Drug Combinations; Endoscopy, Gastrointestinal; Gastrointestinal Hemorrhage [*drug therapy] [mortality]; Injections, Intravenous; Lansoprazole [therapeutic use]; Magnesium [therapeutic use]; Magnesium Hydroxide [therapeutic use]; Randomized Controlled Trials as Topic; Tranexamic Acid [adverse effects] [*therapeutic use]

MeSH check words

Humans