Cochrane
Library

Cochrane Database of Systematic Reviews

Intra-abdominal drains for the prophylaxis of anastomotic leak in elective colorectal surgery (Review)

Rolph R, Duffy JMN, Alagaratnam S, Ng P, Novell R

Rolph R, Duffy JMN, Alagaratnam S, Ng P, Novell R.
Intra-abdominal drains for the prophylaxis of anastomotic leak in elective colorectal surgery.
Cochrane Database of Systematic Reviews 2004, Issue 4. Art. No.: CD002100.
DOI: [10.1002/14651858.CD002100.pub2](https://doi.org/10.1002/14651858.CD002100.pub2).

www.cochranelibrary.com

TABLE OF CONTENTS

HEADER	1
ABSTRACT	1
PLAIN LANGUAGE SUMMARY	2
SUMMARY OF FINDINGS	3
BACKGROUND	5
OBJECTIVES	5
METHODS	5
Figure 1.	7
RESULTS	9
DISCUSSION	11
AUTHORS' CONCLUSIONS	12
ACKNOWLEDGEMENTS	12
REFERENCES	14
CHARACTERISTICS OF STUDIES	16
DATA AND ANALYSES	20
Analysis 1.1. Comparison 1 All studies stratified by type of drain, Outcome 1 Clinical anastomotic dehiscence.	21
Analysis 1.2. Comparison 1 All studies stratified by type of drain, Outcome 2 Mortality.	22
Analysis 1.3. Comparison 1 All studies stratified by type of drain, Outcome 3 Re-intervention.	23
Analysis 1.4. Comparison 1 All studies stratified by type of drain, Outcome 4 Radiological anastomotic dehiscence.	23
Analysis 1.5. Comparison 1 All studies stratified by type of drain, Outcome 5 Wound infection.	24
Analysis 2.1. Comparison 2 All studies stratified by level of anastomosis, Outcome 1 Clinical anastomotic dehiscence.	25
Analysis 2.2. Comparison 2 All studies stratified by level of anastomosis, Outcome 2 Mortality.	26
Analysis 2.3. Comparison 2 All studies stratified by level of anastomosis, Outcome 3 Re-intervention.	26
Analysis 2.4. Comparison 2 All studies stratified by level of anastomosis, Outcome 4 Radiological anastomotic dehiscence.	26
Analysis 2.5. Comparison 2 All studies stratified by level of anastomosis, Outcome 5 Wound infection.	27
APPENDICES	27
WHAT'S NEW	28
HISTORY	29
CONTRIBUTIONS OF AUTHORS	29
DECLARATIONS OF INTEREST	29
SOURCES OF SUPPORT	29
INDEX TERMS	30

[Intervention Review]

Intra-abdominal drains for the prophylaxis of anastomotic leak in elective colorectal surgery

Rachel Rolph¹, James MN Duffy², Swethan Alagaratnam³, Paul Ng⁴, Richard Novell⁵

¹Department of Plastic and Reconstructive Surgery, Guys and St Thomas' NHS Foundation Trust, London, UK. ²iHOPE: International Collaboration to Harmonise Outcomes for Pre-eclampsia, Balliol College, University of Oxford, Oxford, UK. ³Department of Colorectal Surgery, Royal Free Hospital, London, UK. ⁴Department of Colorectal Surgery, St Thomas' Hospital, London, UK. ⁵University Department of Colorectal Surgery, Royal Free Hospital, London, UK

Contact address: Rachel Rolph, Department of Plastic and Reconstructive Surgery, Guys and St Thomas' NHS Foundation Trust, Westminster Bridge Road, London, SE1 7EH, UK. r.rolph1@nhs.net.

Editorial group: Cochrane Colorectal Cancer Group.

Publication status and date: Edited (no change to conclusions), published in Issue 3, 2016.

Citation: Rolph R, Duffy JMN, Alagaratnam S, Ng P, Novell R. Intra-abdominal drains for the prophylaxis of anastomotic leak in elective colorectal surgery. *Cochrane Database of Systematic Reviews* 2004, Issue 4. Art. No.: CD002100. DOI: [10.1002/14651858.CD002100.pub2](https://doi.org/10.1002/14651858.CD002100.pub2).

Copyright © 2016 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.

ABSTRACT

Background

Elective colorectal surgery can involve formation of bowel anastomoses, which may be complicated by postoperative anastomotic leaks. Routine intra-operative drain placement aims to help clinicians diagnose and treat postoperative leaks. There is little agreement on the prophylactic use of drains for elective colorectal anastomoses. Once anastomotic leakage has occurred, it is generally agreed that drains should be used for therapeutic purposes. However, on prophylactic use no such agreement exists.

Objectives

To assess the effectiveness and safety of a prophylactic drain after elective colorectal anastomosis.

Search methods

We searched the Cochrane Colorectal Cancer Group's Specialized Register (February 2015), the Cochrane Central Register of Controlled Trials (CENTRAL) (2015, Issue 2), Ovid MEDLINE (1950 to February 2015) and Ovid EMBASE (1974 to February 2015). We also searched trial registers for ongoing and registered trials, Clinicaltrials.gov and the World Health Organization (WHO) search platform International Clinical Trials Registry Platform.

Selection criteria

We included randomised controlled trials (RCTs) comparing drainage with non-drainage regimens after anastomoses in elective colorectal surgery.

Data collection and analysis

Two review authors independently performed selection of studies, assessment of trial quality and extraction of relevant data; a third review author resolved disagreements. We used GRADE methods to evaluate the quality of evidence.

Main results

Of the 908 participants enrolled (three RCTs), 454 were allocated for drainage and 454 for no drainage. We found no new RCTs for this review update. Two trials reported the primary outcome measure of anastomotic dehiscence. There was no statistically significant difference in anastomotic dehiscence in participants treated with intra-abdominal drainage routinely compared to no treatment (risk ratio (RR) 1.40, 95% confidence intervals (CI) 0.45 to 4.40; $I^2 = 0\%$; 2 RCTs; 809 participants). There was no statistically significant difference in mortality

(RR 0.77, 95% CI 0.41 to 1.45; $I^2 = 0\%$; 3 RCTs; 908 participants); surgical re-intervention (RR 1.11, 95% CI 0.67 to 1.82; $I^2 = 29\%$; 3 RCTs; 908 participants); radiological dehiscence (RR 0.85, 95% CI 0.39 to 1.83; $I^2 = 0\%$; 2 RCTs; 809 participants) and wound infection (RR 0.82, 95% CI 0.45 to 1.51; $I^2 = 0\%$; 3 RCTs; 908 participants) in participants treated with routine prophylactic drainage compared to no treatment undergoing elective colorectal surgery. The quality of evidence was low according to GRADE method assessment.

Authors' conclusions

There was insufficient evidence for the use of prophylactic drains after elective colorectal anastomoses. The conclusions of this review were limited due to the nature of the available clinical data; The three included RCTs performed different interventions with relatively small sample sizes of eligible participants.

PLAIN LANGUAGE SUMMARY

Prophylactic anastomotic drainage for colorectal surgery

Background

We designed this review to compare drainage with non-drainage in people having colorectal surgery (a procedure that is used to repair damage to the colon, rectum, anus and muscle of the lower belly) with bowel anastomosis (where two ends of the bowel are joined to each other). Elective colorectal surgery (where surgery is planned in advance rather than carried out as an emergency) often involves removal of part of the large bowel for a variety of diagnoses with subsequent anastomosis. There is the option for the surgeon to place a drain at the time of surgery to prevent leakage from the anastomosis. This is called a prophylactic drain. Cochrane review authors assessed the evidence for the routine use of prophylactic drain placement after the formation of colorectal anastomoses.

Study characteristics

We included three clinical trials involving 908 participants. The trials were conducted in Germany and France. All trials compared routine anastomotic drainage versus no anastomotic drainage after elective colorectal surgery. The evidence was current to February 2015.

Key results

This review showed no apparent difference in anastomotic leak, death, radiological (x-ray) evidence of anastomotic leak, wound infection or need for re-operation. We found insufficient evidence to support the use of routine prophylactic drains after elective colorectal anastomosis. We based our conclusion on limited evidence with relatively small numbers of participants; this means that it is difficult to detect differences between treatment groups that may be present.

Quality of the evidence

The quality of the evidence was low, making it impossible to draw firm conclusions about the use of routine prophylactic drains after elective colorectal anastomosis. Additional studies are needed to strengthen the conclusion drawn by this systematic review and to provide further analysis using modern colorectal surgery. We found no new evidence since the previous version of our systematic review of 2004.

SUMMARY OF FINDINGS

Summary of findings for the main comparison. Intra-abdominal drainage compared with no drainage for elective colorectal surgery

Intra-abdominal drainage compared with no drainage for elective colorectal surgery

Patient or population: unselected population undergoing elective colorectal anastomosis

Settings: high resource settings

Intervention: intra-abdominal drainage

Comparison: no drainage

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of participants (studies)	Quality of the evidence (GRADE)
	Assumed risk	Corresponding risk			
	No drainage	Intra-abdominal drainage			
Anastomotic dehiscence	Unselected population undergoing elective colorectal anastomosis		RR 1.40 (0.45 to 4.40)	809 (2)	⊕⊕⊕⊕ low
	12 per 1000*	17 per 1000 (5 to 52.8)			
Mortality	Unselected population undergoing elective colorectal anastomosis		RR 0.77 (0.41 to 1.45)	908 (3)	⊕⊕⊕⊕ low
	46 per 1000*	47 per 1000 (19 to 67)			

*The basis for the **assumed risk** (e.g. the median control group risk) 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 (www.gradepro.org)

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.

Assumed risk: anastomotic dehiscence occurred 5 times in 406 participants in the no drainage (control) group. Mortality occurred 21 times in 454 participants in the no drainage (control) group.

The evidence was downgraded to low quality according to GRADE methods; there was substantial clinical heterogeneity and further research in current laparoscopic colorectal practice is very likely to have an important impact on our confidence in the estimate of effect of drainage. In addition, there was a low event rate for anastomotic leak and mortality within the studies assessed, which may not accurately reflect current clinical estimates.

BACKGROUND

Description of the condition

Formation of colorectal anastomoses is common in people undergoing colorectal surgery. A large number of intra-abdominal conditions requiring bowel resection result in the requirement for a bowel anastomosis, for example, colorectal carcinoma and inflammatory bowel disease. There are varying techniques for anastomosis formation, including hand-sewn, stapled and, less commonly, tissue adhesive techniques (Sajid 2012; Sliker 2013; Vakalopoulos 2013). Complications of bowel anastomoses include anastomotic leak (AL), bleeding, fistulae and strictures (Davis 2013). Although a consensus definition of AL is lacking, most surgeons agree that extravasation of contrast on imaging and faecal material seen in drains constitutes a clinical leak (Adams 2013). In simple terms, AL can be defined as the leak of bowel contents from a surgical join between two hollow viscera. Computed tomography (CT) imaging is often used to confirm clinical suspicion of the diagnosis; however, overall sensitivity of CT scanning to diagnose leakage is estimated at 0.68 (95% confidence interval (CI) 0.59 to 0.75) for colonic resection (Kornmann 2013).

AL can be divided into generalised (gross abdominal faecal contamination) or local (localised faecal contamination in the peri-anastomotic space). Practically, there is a large difference between the management of a generalised versus local leak. A generalised leak will result in diffuse peritonitis, acute sepsis and necessitates immediate surgical re-intervention. This will be performed as an emergency procedure (laparoscopic or open) to wash out the contaminated abdomen, take down or reinforce the anastomosis, defunction the person using a stoma to divert faeces from the leak, or a combination of these.

A small contained localised leak in the peri-anastomotic space may produce a localised collection requiring radiological drainage. If the person has already been defunctioned with a stoma at the time of the initial resection, there may be little clinical sequelae. Small contained leaks can lead to enterocutaneous fistula formation or anastomotic stricturing with a significant negative impact on outcome.

AL is associated with prolonged hospital stay; increased morbidity (Montedori 2010; Tsujinaka 2011); and, more pertinently, increased local recurrence, reduced survival following cancer resections and mortality (Mirnezami 2011). A number of studies have used multivariate analysis of participant cohorts to identify risk factors for AL: these are male sex, history of radiotherapy, increased intra-operative times, peri-operative blood transfusion, a high American Society of Anesthesiologists (ASA) score and body mass index greater than 30 kg/m² (Boccola 2011; Klein 2012; Kube 2010; McDermott 2015). The incidence of AL varies from 2% to 39% and is inversely proportional to the distance of the anastomosis from the anal verge (Montedori 2010; Vignali 1997). A variety of clinical markers are currently being developed to predict and diagnose AL accurately and as early as possible. These include elevated cytokines, metallo-proteinases, serum C-reactive protein and reduced intra-operative rectal stump blood flow as measured by laser Doppler (Almeida 2012; Cini 2013; Kao 2012; Vignali 2000).

AL remains a major and frequent complication following colorectal surgery, and is associated with significant postoperative morbidity and mortality. The incidence of AL after colon surgery is reported

to be 1% to 3%, and after colorectal surgery around 10% (Alberts 2003; Borowski 2010). Mortality increases significantly after AL, with AL associated with a 16% mortality compared to 3% without AL (Bakker 2014). It still remains a feared complication in colorectal surgery. Surgical strategies to minimise the risk of AL include creating an anastomosis under no tension, with an adequate blood supply and in a surgical field free of infection.

Description of the intervention

The use of routine prophylactic drains when performing an elective colorectal anastomosis has been debated since the late 1980s. The usefulness of the drain to act as an early indicator of AL and a means of localised drainage to prevent AL remain key points in this debate. This is weighed against drain-related complications including drain-site wound infection, pain, herniation and bleeding. There is the suggestion that drain placement may itself be a risk factor for AL as placement of the drain in close proximity to the anastomosis may mechanically disrupt the tissue or blood supply to the join (Tsujinaka 2011).

How the intervention might work

The prophylactic use of a drain adjacent to the anastomosis could evacuate peri-anastomotic fluid collections, reducing the risk of sepsis and pressure in and around the anastomosis and thus decrease the risk of AL.

The purpose of this review was to compare outcomes after elective colorectal surgery with and without the use of prophylactic anastomotic drains.

Why it is important to do this review

This review assessed and summarised the current evidence regarding the efficacy of routine abdominal drainage for colorectal anastomosis. This review is an update of a previous Cochrane review (Jesus 2004). The comparisons within the review should assist people and their clinicians in choosing an appropriate surgical procedure with a better knowledge of the current evidence. The review serves to highlight the current limitations in the literature and to highlight the need for further research.

OBJECTIVES

To assess the effectiveness and safety of a prophylactic drain after elective colorectal anastomosis.

METHODS

Criteria for considering studies for this review

Types of studies

Published and unpublished randomised controlled trials (RCTs) and cluster-RCTs. We excluded non-randomised and quasi-randomised trials as they are associated with a high risk of bias.

Types of participants

All people undergoing elective primary colorectal anastomosis.

Types of interventions

We included trials if they compared an intra-abdominal drain with placebo (e.g. blind ended drain) or no treatment.

Types of outcome measures

Primary outcomes

- Anastomotic dehiscence (as defined by the trialists).

Secondary outcomes

- Mortality (30-day all-cause mortality).
- Surgical re-intervention (return to theatre for washout or intervention for AL (re-operation)).
- Radiological anastomotic dehiscence (as defined by the trialists).
- Wound infection (as defined by the trialists).

Search methods for identification of studies

We searched for all published and unpublished RCTs, without language restriction and in consultation with the Colorectal Cancer Review Group Trials Search Co-ordinator (Marija Barbateskovic).

Electronic searches

We searched the following electronic databases, trial registers and websites:

- Cochrane Colorectal Cancer Group's Specialized Register (February 2015);
- Cochrane Central Register of Controlled Trials (CENTRAL, 2015, Issue 2) ([Appendix 1](#));
- Ovid MEDLINE (1950 to February 2015) ([Appendix 2](#));
- Ovid EMBASE (1974 to February 2015) ([Appendix 3](#)).

Other electronic searches that we performed included:

- trial registers for ongoing and registered trials: ClinicalTrials.gov ([clinicaltrials.gov](#)) and World Health Organization (WHO)

search platform International Clinical Trials Registry Platform (www.who.int/trialsearch/Default.aspx);

- Citation indexes (scientific.thomson.com/products/sci);
- PubMed (www.ncbi.nlm.nih.gov/pubmed/);
- OpenGrey database (opengrey.eu/) and Google for grey literature;
- Conference abstracts in the Web of Knowledge (wokinfo.com).

Searching other resources

We handsearched reference lists of articles retrieved by the search and contacted experts in the field to obtain additional data. In addition, we handsearched relevant journals and conference abstracts that were not covered in the Colorectal Cancer Review Group Trials register, in liaison with the Trials Search Co-ordinator (UK colorectal and general surgery conference proceedings (Association of Surgeons in Training (ASiT), Association of Surgeons of Great Britain and Ireland (ASGBI), Association of Coloproctology of Great Britain and Ireland (ACPGBI)) and newsletters were reviewed since 2011).

Data collection and analysis

Selection of studies

Three review authors (RR, SA and JD) performed an initial screen of the titles and abstracts retrieved by the search. We retrieved the full texts of all potentially eligible studies. Two review authors (RR and JD) independently examined these full-text articles for compliance with the inclusion criteria and selected studies eligible for inclusion in the review. We included all relevant trials irrespective of whether they reported on measured outcome data. We corresponded with study investigators, as required, to clarify study eligibility or to seek further data where necessary. We resolved disagreements about study eligibility by discussion or by a third review author (RN). See [Figure 1](#) for the selection process in the PRISMA flow chart.

Figure 1. PRISMA study flow diagram.

Data extraction and management

Two review authors (RR and JD) independently extracted the data from eligible studies using a data extraction form designed and pilot-tested by the review authors. We resolved any disagreements by discussion or by a third review author (PN). Data extracted included study characteristics and outcome data. Where studies had multiple publications, we used the main trial report as the reference and derived additional details from the secondary papers. We corresponded with study investigators for further data on methods and results, as required.

Assessment of risk of bias in included studies

Review authors (RR and JD) independently assessed the included studies for risk of bias using the Cochrane 'Risk of bias' assessment tool in the *Cochrane Handbook for Systematic Reviews of Interventions* (Cochrane Bias Methods Group; Higgins 2011) to assess:

- selection bias: allocation (random sequence generation and allocation concealment);
- performance/detection bias: blinding of participants and personnel, blinding of outcome assessors;
- attrition bias: incomplete outcome data;
- reporting bias: selective reporting;
- other bias (differences in baseline characteristics, sample size calculations).

We judged each domain as high, low or unclear risk of bias according to criteria used in the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011).

Selection bias

Was the allocation sequence adequately generated?

- Score 'low risk' if a random component in the sequence generation process was described (e.g. referring to a random number table).
- Score 'high risk' when a non-random method is used (e.g. performed by date of admission).
- Score 'unclear risk' if not specified in the paper.

Was the allocation adequately concealed?

- Score 'low risk' if participants and investigators enrolling participants could not foresee assignment (e.g. because used a centralised randomisation scheme, an on-site computer system or sealed opaque envelopes).
- Score 'high risk' if non-randomised studies.
- Score 'unclear risk' if not specified in the paper.

Performance/detection bias

Was knowledge of the allocated interventions adequately prevented during the study?

- Score 'low risk' if the authors stated explicitly that the primary outcome variables were assessed blindly, or the outcomes were objective (e.g. length of hospital stay).
- Score 'high risk' if the outcomes were not assessed blindly.
- Score 'unclear risk' if not specified in the paper.

Attrition bias

Were incomplete outcome data adequately addressed?

- Score 'low risk' if missing outcome measures were unlikely to bias the results (e.g. reasons for missingness unlikely to be related to true outcome, missing outcome data balanced in numbers across intervention groups with similar reasons for missing data or missing data were imputed using appropriate methods).
- Score 'high risk' if missing outcome data were likely to bias the results.
- Score 'unclear risk' if not specified in the paper.

Reporting bias

Were reports of the study free from selective outcome reporting?

- Score 'low risk' if there is no evidence that outcomes were selectively reported (e.g. the study had a protocol pre-specifying the outcomes, or all relevant outcomes in the methods section were reported in the results section).
- Score 'high risk' if some pre-specified outcomes were subsequently omitted from the results.
- Score 'unclear risk' if not specified in the paper.

Were reports of the study free from selective analysis reporting?

- Score 'low risk' for each outcome if there was no evidence that analyses were selectively reported (e.g. analyses were defined in the methods section of the protocol or paper).
- Score 'high risk' if there was evidence of selective analysis reporting (e.g. multiple adjusted analyses were carried out and only one reported, or unusual cut-points were used for categorising an outcome).
- Score 'unclear' risk if unclear from the paper.

We resolved disagreements by discussion or by a third review author (RN). We described all the judgements fully and presented the conclusions in the 'Risk of bias' table. We used sensitivity analyses to incorporate the risk of bias into the interpretation of review findings. We took care to search for within trial selective reporting, such as trials failing to report obvious outcomes or reporting them in insufficient detail to allow inclusion. We sought published protocols and compared the outcomes between the protocol and the final published study.

Measures of treatment effect

All outcomes were dichotomous. We used the numbers of events in the control and intervention groups of each study and calculated risk ratios (RRs). We planned to reverse the direction of effect of individual studies, if required, to ensure consistency across trials. We presented 95% CI for all outcomes. We compared the magnitude and direction of effect reported by studies with how they are presented in the review, taking into account the clinical significance of these differences.

Unit of analysis issues

All included RCTs were parallel in design, therefore the unit of analysis was the individual participant. Should we identify any cluster-randomised trials in future updates, we will include cluster RCTs in the analyses along with individually randomised trials. We will adjust their sample sizes or standard errors using an

estimate of the intracluster correlation co-efficient (ICC) derived from the trial (if possible), from a similar trial or from a study of a similar population. If we use ICCs from other sources, we will report this and conduct sensitivity analyses to investigate the effect of variation in the ICC. If we identify both cluster-randomised trials and individually randomised trials, we plan to synthesise the relevant information. We will consider it reasonable to combine the results from both if there is little heterogeneity between the study designs and the interaction between the effect of intervention and the choice of randomisation unit is considered to be unlikely.

Dealing with missing data

We analysed studies on an intention-to-treat basis, as far as possible. If data were unclear or missing, we contacted the authors of the studies for further details. If there were no further data or details made available from the contacted authors, we assessed the studies on the data provided and excluded them from overall analysis if they did not meet the inclusion criteria.

Assessment of heterogeneity

We considered whether the clinical and methodological characteristics of the included studies were sufficiently similar for meta-analysis to provide a clinically meaningful summary. We assessed statistical heterogeneity using the I^2 statistic. An I^2 value greater than 50% indicated substantial heterogeneity (Higgins 2011).

Assessment of reporting biases

Reporting biases arise when the dissemination of research findings is influenced by the nature and direction of results. Some types of reporting bias (e.g. publication bias, multiple publication bias, language bias) reduce the likelihood that all studies eligible for a review will be retrieved. If all eligible studies are not retrieved, the review may also be biased. In view of the difficulty of detecting and correcting for publication bias and other reporting biases, we aimed to minimise the potential impact by ensuring a comprehensive search for eligible studies and by being alert for duplication of data. We sought published protocols and compared the outcomes between the protocol and the final published study. As the review included only three RCTs, the use of funnel plots to evaluate publication bias was not appropriate.

Data synthesis

When studies were sufficiently similar, we combined the data using a fixed-effect model. The trials were clinically heterogeneous; however, there were not enough to inform the distribution of effects in a random-effects model, therefore the choice of a fixed-effect model was due to the small number of trials. An increase in the RRs of a particular outcome, which is detrimental to the participant (e.g. AL) is displayed graphically in the meta-analysis to the right of the centre line and a decrease in the RR of an outcome to the left of the centre line.

Subgroup analysis and investigation of heterogeneity

We performed subgroup analysis to determine if the incidence of AL in the two drainage groups was affected by the level of colorectal anastomosis (intra- or extra-peritoneal) and type of drainage (active or passive drains).

Sensitivity analysis

We conducted sensitivity analyses for the primary outcome to determine whether the conclusions were robust to arbitrary decisions made regarding the eligibility and analysis. These analyses included consideration of whether the review conclusions would have differed if we restricted eligibility to studies without high risk of bias.

Summary of findings

We assessed the quality of evidence for the outcomes anastomotic dehiscence and mortality for intra-abdominal drainage versus no drainage for elective colorectal surgery using the Grading of Recommendations, Assessment, Development and Evaluation (GRADE) approach and created a 'Summary of findings' table using the GRADEpro Guideline Development tool (GRADEpro).

The GRADE approach classifies the quality of evidence in one of four grades:

- high: further research is very unlikely to change our confidence in the estimate of effect;
- moderate: further research is likely to have an impact on our confidence in the estimate of effect and may change the estimate;
- low: 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: any estimate of effect is very uncertain.

The quality of evidence can be downgraded by one (serious concern) or two levels (very serious concern) for the following reasons: risk of bias, inconsistency (unexplained heterogeneity, inconsistency of results), indirectness (indirect population, intervention, control or outcomes) and imprecision (wide CIs, single trial). The quality can also be upgraded by one level due to a large summary effect estimate.

RESULTS

Description of studies

Results of the search

The search resulted in a total of 837 records. After deleting duplicates, we screened 577 records. We excluded 568 clearly irrelevant records leaving nine potentially eligible studies of which we retrieved full texts (Mennigen 1989; Merad 1998; Merad 1999; Brown 2001; Hoffmann 1987; Johnson 1989; Sagar 1993; Sagar 1995; Hagmuller 1990). We excluded five studies for reasons listed in the Characteristics of excluded studies table (Brown 2001; Hoffmann 1987; Johnson 1989; Sagar 1993; Sagar 1995), and listed one study as awaiting classification (Hagmuller 1990) as we were unable to retrieve the original paper from authors (see Characteristics of studies awaiting classification table). Finally, three studies met our inclusion criteria (Mennigen 1989; Merad 1998; Merad 1999). Figure 1 shows the study flow diagram.

See: Characteristics of included studies table, Characteristics of excluded studies table and Characteristics of ongoing studies table.

Included studies

Study design and setting

The review included three RCTs (Mennigen 1989; Merad 1998; Merad 1999). One trial was conducted in a single centre (Mennigen 1989), and two trials were conducted in multiple centres (Merad 1998; Merad 1999). The trials were conducted in France (Merad 1998; Merad 1999), and Germany (Mennigen 1989).

Participants

The studies randomised 908 adults (Mennigen 1989; Merad 1998; Merad 1999). Participant's ages ranged from 15 to 98 years. Male to female ratios were approximately balanced. The majority of participants underwent colorectal resection for carcinoma. The other reasons for resection included benign tumours, diverticular disease and inflammatory bowel disease.

Interventions

One study used silicone passive drain (Mennigen 1989), one study used closed suction and silicone passive drains (Merad 1998), and one study used closed suction drain (Merad 1999). Drains were positioned at the site of anastomosis. They used silk sutures to secure the drain.

Primary outcome

- Anastomotic leak

All trials reported anastomotic dehiscence. Anastomotic dehiscence was determined clinically by clinical signs and radiological imaging. Definitions of outcome measures were consistent between the three studies. However, radiological anastomotic dehiscence was subject to radiologist interpretation on each of the sites and, therefore, may have differed between the studies in reporting rates of AL. AL usually presents within seven to 10 days postoperatively; however, reporting for AL is at 30 days postoperatively.

Secondary outcomes

All trials reported the following secondary outcomes:

- mortality;
- surgical re-intervention;
- radiological anastomotic dehiscence;
- wound infection (not standardised across trials).

The participants assigned to the drainage group compared with the participants assigned to non-drainage group showed:

- anastomotic dehiscence: 4% (18/454) participants compared to 4% (18/454) participants;
- mortality: 3.5% (16/454) participants compared to 6.6% (30/454) participants;
- surgical intervention: 6.8% (31/454) participants compared to 6.2% (28/454) participants;
- radiological anastomotic dehiscence: 2.4% (11/454) participants compared to 2.9% (13/454) participants;
- wound infection: 4% (18/454) participants compared to 4.8% (22/454) participants.

Excluded studies

The review excluded six trials for the following reasons:

- Brown 2001, Hoffmann 1987, and Johnson 1989 were quasi-randomised. These studies stated "drawing of sealed envelopes" as the method of sequence generation for randomisation. We judged this as pseudo-randomisation as the investigators were taking from a receptacle. There was no description of random sequence generation and the authors did not provide further information, for example sequentially numbered envelopes. If future assignments can be anticipated, either by predicting them (number of envelopes left in receptacle) or knowledge of envelope contents (previous envelope used again in same receptacle), then selection bias can arise due to the selective enrolment and non-enrolment of participants into the study (Section 8.9.1; *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011)).
- Sagar 1993 and Sagar 1995 included people having emergency surgery;
- Hagmuller 1990 awaiting classification. Review authors awaiting paper for detailed analysis (requested from German authors).

Risk of bias in included studies

See [Characteristics of included studies](#) table.

Allocation

All included trials were at low risk of selection bias related to sequence generation as they used random number tables. None of the included trials described the allocation concealment method used and were at unclear risk of selection bias related to allocation concealment.

Blinding

We considered that blinding of participants and outcome observers would not influence the objective primary or secondary outcomes. It would not be possible to blind the personnel involved in surgical interventions. It would have been challenging to blind study participants. It would have been possible to blind outcome assessors following drain removal. None of the included trials stated whether participants and outcome assessors were blinded.

Incomplete outcome data

All trials reported their exclusions and losses to follow-up. The studies were at low risk of attrition bias. The reported drop-out rates were low for all three studies, 1/100 (1%) (Mennigen 1989), and lost to follow-up rates were 2/317 (less than 1%) (Merad 1998), and 2/492 (less than 1%). An assumption was made that the five participants who dropped out were unlikely to bias the results (e.g. reasons for the drop-out unlikely to be related to true outcome measures).

Selective reporting

We were unable to identify published protocols for the included studies. The included RCTs reported all outcomes that were pre-stated in the methods section. All included RCTs reported adverse events.

Other potential sources of bias

No studies reported substantial baseline differences in the treatment and control groups and in sample size calculation.

Effects of interventions

See: [Summary of findings for the main comparison](#) Intra-abdominal drainage compared with no drainage for elective colorectal surgery

Primary outcome

Anastomotic dehiscence (30-day postoperative)

There was no statistically significant difference in anastomotic dehiscence in participants treated with intra-abdominal drainage compared with no treatment (RR 1.40, 95% CI 0.45 to 4.40; 2 RCTs; 809 participants) ([Analysis 1.1](#)). There was a low degree of heterogeneity ($I^2 = 0\%$); however, this has little value due to the low number of included trials. When we carried out the pre-defined subgroup analysis, anastomotic dehiscence was not affected by the level of colorectal anastomosis or drain type ([Analysis 1.1](#); [Analysis 2.1](#)). We were unable to carry out the pre-defined sensitivity analysis, as all included trials were associated with a high risk of bias.

Secondary outcomes

Mortality (30-day postoperative)

All trials reported 30-day mortality. There was no statistically significant difference in anastomotic dehiscence in participants treated with intra-abdominal drainage compared with no treatment (RR 0.77, 95% CI 0.41 to 1.45; 3 RCTs; 908 participants) ([Analysis 1.2](#)). There was a low degree of heterogeneity ($I^2 = 0\%$).

Surgical re-intervention (30-day postoperative)

All trials reported surgical re-intervention. There was no statistically significant difference in anastomotic dehiscence in participants treated with intra-abdominal drainage compared with no treatment (RR 1.11, 95% CI 0.67 to 1.82; 3 RCTs; 908 participants) ([Analysis 1.3](#)). There was a low degree of heterogeneity ($I^2 = 29\%$).

Radiological anastomotic dehiscence (30-day postoperative)

All trials reported radiological anastomotic dehiscence. AL was assessed clinically and with radiological imaging. There was no statistically significant difference in anastomotic dehiscence in participants treated with intra-abdominal drainage compared with no treatment (RR 0.85, 95% CI 0.39 to 1.83; 3 RCTs; 908 participants) ([Analysis 1.4](#)). There was a low degree of heterogeneity ($I^2 = 0\%$).

Wound infection (30-day postoperative)

All trials reported wound infection. Trials did not use a validated classification system to define wound infection. The definition of wound infection was based on clinical signs and elevated laboratory blood markers for infection. There was no statistically significant difference in wound infection in participants treated with intra-abdominal drainage compared with no treatment (RR 0.82, 95% CI 0.45 to 1.51; 3 RCTs; 908 participants) ([Analysis 1.5](#)). There was a low degree of heterogeneity ($I^2 = 0\%$).

DISCUSSION

The purpose of this review was to compare outcomes after elective colorectal surgery with and without the use of prophylactic anastomotic drains.

Meta-analysis of three RCTs showed no statistically significant difference in the incidence of clinically or radiologically determined AL associated with the use of drains ([Mennigen 1989](#); [Merad 1998](#); [Merad 1999](#)). There was no detectable difference in mortality between the drainage and non-drainage groups ([Mennigen 1989](#); [Merad 1998](#); [Merad 1999](#)). Re-operation rates were consistent between the two groups, suggesting that in the event of AL, a drain placed at the time of surgery may not have a therapeutic benefit or change the outcome. However, the power of the analyses in this review was low, therefore, no firm conclusions can be made based on current evidence ([Mennigen 1989](#); [Merad 1998](#); [Merad 1999](#)).

The incidence of AL in the two drainage groups was also not affected by whether the anastomosis was intra- or extra-peritoneal, there was no evidence of a greater benefit in drains covering low rectal anastomoses ([Merad 1999](#)). This subgroup analysis was limited in its conclusion due to limited participant numbers and a very low event rate of AL (4%); low participant numbers makes it difficult for analyses to detect differences between groups that may be present. Many surgeons routinely place drains after mid- to low-rectal resections with a view to draining pelvic haematomas postoperatively rather than detecting or preventing ALs. Further high-quality large RCTs are required to address this topic in greater detail.

There may be several reasons why drains placed at surgery appear to be of no benefit. Following surgery, drains will often become blocked, eradicating their therapeutic potential ([Averbach 1995](#)). Drains may also become displaced from their original placement site. Furthermore, the studies employed in this meta-analysis used various types of drain. The differing properties of the drainage systems used could have affected the results and needs to be investigated further. The reasoning against the use of drains in colorectal surgery includes concerns about potential negative effects of drains. Indeed, the routine use of drains is not recommended in colorectal enhanced recovery programmes, which have become widespread in recent years, due to concerns regarding increased pain, difficulty with mobilisation and subsequent complications ([Lassen 2009](#)).

The conclusions of this review were limited by the heterogeneous nature of the operations performed in the included studies, which included both intra- and extra-peritoneal anastomoses. The lack of recruitment of sufficient sample sizes in the included studies means that the conclusions may be limited due to inadequate power and, therefore, a type II error. In particular, the studies included are relatively dated and there was no new evidence for this review since 2004. The available evidence was insufficient to draw firm conclusions regarding the benefit or harm from routine drainage in colorectal resections as the two main results from the pooled data were contradictory and inconsistent. Further trials are required, not only to determine the safety and efficacy of drains, but also their role in diagnosing anastomotic leakage, which has been one of the purported benefits of routine drain placement. In addition, trials are needed to investigate the effects of drainage in the emergency colorectal surgical setting which is not addressed in this review.

Summary of main results

There was insufficient evidence to support or refute the routine use of abdominal drainage for elective colorectal anastomosis. There were no statistical significant differences in the primary outcome, postoperative anastomotic dehiscence, or secondary outcomes including mortality, surgical re-intervention, radiological anastomotic dehiscence and wound infection.

Overall completeness and applicability of evidence

The trials recruited participants considered at low anaesthetic and surgical risk, therefore, the applicability of this review to inform practice in high-risk people undergoing surgery is limited. There was insufficient evidence to draw any conclusions regarding differences in efficacy for people with benign disease compared with people with malignancy. All trials recruited participants within high resource settings, which limits the applicability of this research to inform practice within low resource settings. All included studies reported the primary and secondary outcomes but there was substantial clinical heterogeneity. Clinically, colorectal surgery has moved towards minimally invasive laparoscopic operations and enhanced recovery pathways. The studies included did not address the effectiveness of drains in current colorectal surgical practice.

Quality of the evidence

The quality of the evidence for effectiveness outcomes was low, denoting that further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate. The methodological bias was low in the included trials as methods of randomisation were described, and it was unclear if blinding was likely to affect the objective primary and secondary outcome measures. There was sufficient evidence to reach preliminary conclusions regarding primary and secondary outcomes. Lack of power calculations in the included studies means that the conclusions drawn may be subject to type II errors or due to problems in participant recruitment to the trial.

Potential biases in the review process

The strengths of this systematic review include its robust search strategy (guided and developed by the Colorectal Cancer Review Group) and the methodological design and statistical analysis (developed with reference to the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011)). The protocol was peer-reviewed, published and freely available. We applied no limitations such as a publication type, language or date restrictions. Two review authors independently conducted study selection, risk of bias assessment and data collection. We resolved disagreements by discussion or by a third review author (RN or PN). All review authors have stated that there are no conflicts of interest. Systematic reviews are not without limitations. The comprehensive search strategy identified only three trials, making extrapolation difficult. Notably, two of the studies are from the same research group, which may be a potential source of bias. The studies included are relatively dated in surgical practice and do not account for advances in laparoscopic surgical approaches. This may limit the applicability of the conclusions to current practice.

Agreements and disagreements with other studies or reviews

The results from this systematic review were in agreement with previous systematic reviews on this topic, which also concluded that there is insufficient evidence to support the use of routine drainage in elective colorectal surgery (Karliczek 2006; Urbach 1999; Jesus 2004). We found no new evidence for this review compared to the first published version of the systematic review in 2004 (Jesus 2004).

The findings of this review agree with additional systematic reviews on the routine use of drains for abdominal surgery including gastrectomy, liver resection, cholecystectomy and incisional hernia repair (Gurusamy 2007a; Gurusamy 2007b; Gurusamy 2013; Wang 2011). There remains a paucity of high-quality RCTs in this field and large-scale RCTs are required to provide robust scientific evidence in current surgical practice.

AUTHORS' CONCLUSIONS

Implications for practice

There remains no firm evidence for the prophylactic use of drains in preventing anastomotic leak after colorectal surgery, although the review also does not reveal any significant contraindication to the use of drains. The conclusions of this review were based on evidence from three studies all performed in the 1980s and 1990s with no new studies since the previous systematic review in 2004 (Jesus 2004). The applicability of the conclusions based on the included studies to current minimally invasive colorectal surgery with enhanced recovery programmes is unclear and remains a significant limitation of the review.

Implications for research

Uncertainty about the effects of drains in colorectal surgery remains. A large-scale randomised controlled trial would help to resolve these uncertainties. This trial should be performed in the context of a modern enhanced recovery programme following elective laparoscopic resection, with the drainage group having a standardised passive closed drain. All participants would need to be randomised to drainage or no drainage with both groups receiving enhanced recovery programmes. Blinding participants and clinicians to the intervention would not be possible. The endpoints should be clearly defined, and in addition to the endpoints used in this review, include measurements of postoperative pain, and specific drain-related complications such as herniation, wound infection and quality of life scores. A prospective protocol with consensus outcome definitions should be published. In particular, radiological anastomotic leak as an outcome measure requires a consensus opinion between three independent radiologists to confirm an anastomotic leak to prevent subjective reporting. Given the relatively high leak rates after low rectal anastomosis, this should form a separate subgroup for analysis. In addition, separate trials should be conducted to investigate the effects of drains in emergency colorectal surgery.

ACKNOWLEDGEMENTS

We would like to acknowledge Jesus EC, Karliczek A, Matos D, Castro AA and Atallah AN, authors of the previous Cochrane review (Jesus 2004). Thanks to the Cochrane Colorectal Cancer Group (Henning Keinke Andersen and Marija Barbateskovic) for their

invaluable support, editors and peer referees, and the Cochrane

Copy Edit Support (Anne Lawson) for careful copy editing of this updated review.

REFERENCES

References to studies included in this review

Mennigen 1989 {published data only}

Mennigen R, Kusche J, Troidl H. Prophylaktische drainage von kolonanastomosen. *Coloproctology* 1989;**11**(2):76-80.

Merad 1998 {published data only}

Merad F, Yahchouchi E, Hay JM, Fingerhut A, Laborde Y, Langlois-Zantain O, for the French Associations for Surgical Research. Prophylactic abdominal drainage after elective colonic resection and suprapromontory anastomosis. *Archives of Surgery* 1998;**133**(3):309-14. [MEDLINE: 1998176925]

Merad 1999 {unpublished data only}

Merad F, Hay JM, Fingerhut A, Yahchouchi E, Laborde Y, Pelissier E, et al. Is prophylactic pelvic drainage useful after elective rectal or anal anastomosis? A multicenter controlled randomized trial. *Surgery* 1999;**125**:529-35.

References to studies excluded from this review

Brown 2001 {published data only}

Brown SR, Seow-Choen F, Eu KW, Heah SM, Tang CL. A prospective randomised study of drains in infra-peritoneal rectal anastomoses. *Techniques in Coloproctology* 2001;**5**:89-92.

Hoffmann 1987 {published data only}

Hoffmann J, Shokouh-Amiri MH, Damm P, Jersen R. A prospective, controlled study of prophylactic drainage after colonic anastomoses. *Diseases of the Colon and Rectum* 1987;**30**(6):449-52. [MEDLINE: 1987245965]

Johnson 1989 {published data only}

Johnson CD, Lamont PM, Orr N, Lennox M. Is a drain necessary after colonic anastomosis?. *Journal of the Royal Society of Medicine* 1989;**82**(11):661-4. [MEDLINE: 1990079903]

Sagar 1993 {published data only}

Sagar PM, Couse N, Kerin N, May J, MacFie J. Randomized trial of drainage of colorectal anastomosis. *British Journal of Surgery* 1993;**80**:769-71.

Sagar 1995 {published data only}

Sagar PM, Hartley MN, MacFie J, Mancey-Jones B, Sedman P, May J. Randomized trial of pelvic drainage after rectal resection. *Diseases of the Colon and Rectum* 1995;**38**(3):254-8. [MEDLINE: 1995188656]

References to studies awaiting assessment

Hagmuller 1990 {published data only}

Hagmuller E, Lorenz D, Werthmann K, Trede M. Uses and risks of drainage following elective colon resection. A prospective, randomized and controlled clinical study. *Chirurgie* 1990;**61**(4):266-71. [MEDLINE: 1990269140]

Additional references

Adams 2013

Adams K, Papagrigoriadis S. Little consensus in either definition or diagnosis of a lower gastro-intestinal anastomotic leak amongst colorectal surgeons. *International Journal of Colorectal Disease* 2013;**28**(7):967-71.

Alberts 2003

Alberts JCJ, Parvaiz A, Moran BJ. Predicting risk and diminishing the consequences of anastomotic dehiscence following rectal resection. *Colorectal Disease* 2003;**5**:478-82.

Almeida 2012

Almeida AB, Faria G, Moreira H, Pinto-de-Sousa J, Correia-da-Silva P, Maia JC. Elevated serum C-reactive protein as a predictive factor for anastomotic leakage in colorectal surgery. *International Journal of Surgery* 2012;**10**(2):87-91.

Averbach 1995

Averbach AM, Sugarbaker PH. The use of drains in elective surgery for colorectal cancer: always, never or selectively?. *Tumori* 1995;**81**(3 Suppl):89-97.

Bakker 2014

Bakker IS, Grossmann I, Henneman D, Havenga K, Wiggers T. Risk factors for anastomotic leakage and leak-related mortality after colonic cancer surgery in a nationwide audit. *British Journal of Surgery* 2014;**101**(4):424-32.

Boccola 2011

Boccola MA, Buettner PG, Rozen WM, Siu SK, Stevenson AR, Stitz R, et al. Risk factors and outcomes for anastomotic leakage in colorectal surgery: a single-institution analysis of 1576 patients. *World Journal of Surgery* 2011;**35**(1):186-95.

Borowski 2010

Borowski DW, Bradburn DM, Mills SJ, Bharathan B, Wilson RG, Ratcliffe AA, et al. Volume-outcome analysis of colorectal cancer-related outcomes. *British Journal of Surgery* 2010;**97**:1416-30.

Cini 2013

Cini C, Wolthuis A, D'Hoore A. Peritoneal fluid cytokines and matrix-metalloproteinases as early markers of anastomotic leakage in colorectal anastomosis. A literature review and meta-analysis. *Colorectal Disease* 2013;**15**(9):1070-7.

Cochrane Bias Methods Group

Cochrane Bias Methods Group. Assessing risk of bias in included studies, 2013. bmj.cochrane.org/assessing-risk-bias-included-studies (accessed 23 March 2016).

Davis 2013

Davis B, Rivadeneira DE. Complications of colorectal anastomoses: leaks, strictures, and bleeding. *Surgical Clinics of North America* 2013;**93**(1):61-87.

GRADEpro [Computer program]

McMaster University. GRADEpro. McMaster University, 2015.

Gurusamy 2007a

Gurusamy KS, Samraj K, Davidson BR. Routine abdominal drainage for uncomplicated liver resection. *Cochrane Database of Systematic Reviews* 2007, Issue 3. [DOI: [10.1002/14651858.CD006232.pub2](https://doi.org/10.1002/14651858.CD006232.pub2)]

Gurusamy 2007b

Gurusamy KS, Samraj K, Mullerat P, Davidson BR. Routine abdominal drainage for uncomplicated laparoscopic cholecystectomy. *Cochrane Database of Systematic Reviews* 2007, Issue 3. [DOI: [10.1002/14651858.CD006004.pub3](https://doi.org/10.1002/14651858.CD006004.pub3)]

Gurusamy 2013

Gurusamy KS, Allen VB. Wound drains after incisional hernia repair. *Cochrane Database of Systematic Reviews* 2013, Issue 12. [DOI: [10.1002/14651858.CD005570.pub4](https://doi.org/10.1002/14651858.CD005570.pub4)]

Higgins 2011

Higgins JPT, Green S (editors). *Cochrane Handbook for Systematic Reviews of Intervention* Version 5.1.0 [updated March 2011]. The Cochrane Collaboration, 2011. Available from www.cochrane-handbook.org.

Kao 2012

Kao LS, Millas SG. Predicting the risk of anastomotic leakage in left-sided colorectal surgery using a Colon Leakage Score. *Journal of Surgical Research* 2012;**173**(2):246-8.

Karliczek 2006

Karliczek A, Jesus EC, Matos D, Castro AA, Atallah AN, Wiggers T. Drainage or nondrainage in elective colorectal anastomosis: a systematic review and meta-analysis. *Colorectal Disease* 2006;**8**(4):259-65.

Klein 2012

Klein M, Gögenur I, Rosenberg J. Postoperative use of non-steroidal anti-inflammatory drugs in patients with anastomotic leakage requiring reoperation after colorectal resection: cohort study based on prospective data. *BMJ* 2012;**26**:345.

Kornmann 2013

Kornmann VN, Treskes N, Hoonhout LH, Bollen TL, van Ramshorst B, Boerma D. Systematic review on the value of CT scanning in the diagnosis of anastomotic leakage after colorectal surgery. *International Journal of Colorectal Disease* 2013;**28**(4):437-45.

Kube 2010

Kube R, Mroczkowski P, Granowski D, Benedix F, Sahm M, Schmidt U, et al. Quality assurance in primary colorectal carcinoma [Qualitätssicherung Kolon/Rektum-Karzinome (Primärtumor)]. *European Journal of Surgical Oncology* 2010;**36**(2):120-4.

Lassen 2009

Lassen K, Soop M, Nygren J, Cox PB, Hendry PO, Spies C, et al. Consensus review of optimal perioperative care in colorectal

surgery: Enhanced Recovery After Surgery (ERAS) Group recommendations. *Archives of Surgery* 2009;**144**(10):961-9.

McDermott 2015

McDermott FD, Heeney A, Kelly ME, Steele RJ, Carlson GL, Winter DC. Systematic review of preoperative, intraoperative and postoperative risk factors for colorectal anastomotic leaks. *British Journal of Surgery* 2015;**102**(5):462-79.

Mirnezami 2011

Mirnezami A, Mirnezami R, Chandrakumaran K, Sasapu K, Sagar P, Finan P. Increased local recurrence and reduced survival from colorectal cancer following anastomotic leak: systematic review and meta-analysis. *Annals of Surgery* 2011;**253**(5):890-9.

Montedori 2010

Montedori A, Cirocchi R, Farinella E, Sciannameo F, Abraha I. Covering ileo- or colostomy in anterior resection for rectal carcinoma. *Cochrane Database of Systematic Reviews* 2010, Issue 5. [DOI: [10.1002/14651858.CD006878.pub2](https://doi.org/10.1002/14651858.CD006878.pub2)]

Sajid 2012

Sajid MS, Rafay M, Siddiqui MR, Baig MK. Single layer versus double layer suture anastomosis of the gastrointestinal tract. *Cochrane Database of Systematic Reviews* 2012, Issue 1. [DOI: [10.1002/14651858.CD005477.pub4](https://doi.org/10.1002/14651858.CD005477.pub4)]

Slieker 2013

Slieker JC, Daams F, Mulder IM, Jeekel J, Lange JF. Systematic review of the technique of colorectal anastomosis. *Journal of the American Medical Association of Surgery* 2013;**148**(2):190-201.

Tsujinaka 2011

Tsujinaka S, Konishi F. Drain vs no drain after colorectal surgery. *Indian Journal of Surgical Oncology* 2011;**2**(1):3-8.

Urbach 1999

Urbach DR, Kennedy ED, Cohen MM. Colon and rectal anastomoses do not require routine drainage: a systematic review and meta-analysis. *Annals of Surgery* 1999;**229**(2):174-80.

Vakalopoulos 2013

Vakalopoulos KA, Daams F, Wu Z, Timmermans L, Jeekel JJ, Kleinrensink GJ, et al. Tissue adhesives in gastrointestinal anastomosis: a systematic review. *Journal of Surgical Research* 2013;**180**(2):290-300.

Vignali 1997

Vignali A, Fazio VW, Lavery IC, Milsom JW, Church JM, Hull TL, et al. Factors associated with the occurrence of leaks in stapled rectal anastomoses: a review of 1014 patients. *Journal of the American College of Surgeons* 1997;**185**(2):113-21.

Vignali 2000

Vignali A, Gianotti L, Braga M, Radaelli G, Malvezzi L, Di Carlo V. Altered microperfusion at the rectal stump is predictive for rectal anastomotic leak. *Diseases of the Colon and Rectum* 2000;**43**(1):76-82.

Wang 2011

Wang Z, Chen J, Su K, Dong Z. Abdominal drainage versus no drainage post gastrectomy for gastric cancer. *Cochrane Database of Systematic Reviews* 2011, Issue 8. [DOI: [10.1002/14651858.CD008788.pub3](https://doi.org/10.1002/14651858.CD008788.pub3)]

References to other published versions of this review

Jesus 2004

Jesus EC, Karliczek A, Matos D, Castro AA, Atallah AN. Prophylactic anastomotic drainage for colorectal surgery. *Cochrane Database of Systematic Reviews* 2004, Issue 2. [DOI: [10.1002/14651858.CD002100.pub2](https://doi.org/10.1002/14651858.CD002100.pub2)]

CHARACTERISTICS OF STUDIES

Characteristics of included studies [ordered by study ID]

Mennigen 1989

Methods	Sequence generation: random number tables Allocation concealment: not described Blinded: not described
Participants	Inclusion criteria: elective resection, intra-abdominal anastomosis Exclusion criteria: obstruction, perforation or sepsis present at time of performing anastomosis, i.e. emergency operation Type of disease: 77/100 carcinoma, 6/100 adenoma, 7/100 diverticular disease, 9/100 inflammatory colon disease, 1/100 type of disease not reported (for 1 drop-out) Type of anastomosis: 59/100 left hemicolectomy, 33/100 right hemicolectomy, 7/100 subtotal colectomy, 1/100 not described Number of participants: 100 Age range: 15-87 years Gender: 49 male, 50 female Place of study: Germany (single centre) Time of study: June 1984 to November 1986
Interventions	Treatment: silicone drain Control: no drain
Outcomes	Primary outcome: anastomotic dehiscence Secondary outcomes: mortality, surgical re-intervention, radiological anastomotic dehiscence and wound infection
Notes	Publication: journal article

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Random number tables
Allocation concealment (selection bias)	Unclear risk	No description

Mennigen 1989 *(Continued)*

Blinding (performance bias and detection bias) All outcomes	Unclear risk	Unclear
Blinding of participants and personnel (performance bias) All outcomes	High risk	No description
Blinding of outcome assessment (detection bias) All outcomes	High risk	No description
Incomplete outcome data (attrition bias) All outcomes	Low risk	Drop-out 1/100 (1%)
Selective reporting (reporting bias)	Low risk	Reported all outcomes that were pre-stated in the methods section
Other bias	Unclear risk	No studies reported substantial baseline differences in the treatment and control groups. We found no other potential sources of bias within the included studies Sample size calculation not described. Source of bias

Merad 1998

Methods	Sequence generation: random number tables Allocation concealment: not described Blinded: not described
Participants	Inclusion criteria: elective resection, intra-abdominal anastomosis Exclusion criteria: emergency operation, infection at time of inclusion, resection without anastomosis or with pelvic anastomosis Type of disease: 223/317 carcinoma, 13/317 sigmoid diverticular disease, 34/317 benign tumour, 18/317 Crohn's disease, 27/317 other, 2/317 (information not provided in the study) Type of anastomosis: 222/317 ileocolica, 95/317 colocolica Methods of anastomosis: 124/317 manual, 193/317 stapled Number of eligible participants: 319 Gender: 135 male and 184 female Number of participants randomised: 317 (two participants excluded due to protocol violations); 156 patients were randomized to the abdominal drainage group and 161 to the no abdominal drainage group. Age range: 21-95 years Place of study: France (15 centres) Time of study: 1990-1995

Merad 1998 (Continued)

Interventions	Treatment: drain (82/156 suction - multi-perforated tubular 14F polyvinylchloride catheter; 74/156 non-suction - silicone multi-tubular 10 mm) Control: no drain
Outcomes	Primary outcome: anastomotic dehiscence Secondary outcomes: mortality, surgical re-intervention, radiological anastomotic dehiscence and wound infection
Notes	Publication: journal article

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Random number tables
Allocation concealment (selection bias)	Unclear risk	No description
Blinding (performance bias and detection bias) All outcomes	Unclear risk	Unclear
Blinding of participants and personnel (performance bias) All outcomes	High risk	No description
Blinding of outcome assessment (detection bias) All outcomes	High risk	No description
Incomplete outcome data (attrition bias) All outcomes	Low risk	Lost to follow-up 2/317 (< 1%)
Selective reporting (reporting bias)	Low risk	Reported all outcomes that were pre-stated in the methods section
Other bias	Unclear risk	No studies reported substantial baseline differences in the treatment and control groups Sample size calculation not described. Source of bias

Merad 1999

Methods	Sequence generation: random number tables Allocation concealment: not described Blinded: not described
Participants	Inclusion criteria: elective resection with pelvic anastomosis

Merad 1999 (Continued)

Exclusion criteria: intra-abdominal anastomosis, infection, inadequate haemostasis, emergency operation

Type of disease: carcinoma: 304/492, sigmoid diverticular disease: 123/492, benign tumour: 23/492, Crohn's disease: 13/492, other: 26/492, 3/492 (information not provided in the study)

Type of anastomosis: 360/492 suprapерitoneal rectum, 115/492 intraperitoneal rectum, 17/492 anus

Methods of anastomosis: 360/492 manual, 132/492 stapled

Number of participants: 492

Age range: 24-98 years

Gender: 248 male, 244 female

Place of study: France (18 centres)

Time of study: September 1990 to June 1995

Interventions	Treatment: multi-perforated polyvinyl chloride F14 suction drains Control: no drain
Outcomes	Primary outcome: anastomotic dehiscence Secondary outcomes: mortality, surgical re-intervention, radiological anastomotic dehiscence and wound infection
Notes	Publication: journal article

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Random number tables
Allocation concealment (selection bias)	High risk	No description
Blinding (performance bias and detection bias) All outcomes	Unclear risk	Unclear
Blinding of participants and personnel (performance bias) All outcomes	High risk	No description
Blinding of outcome assessment (detection bias) All outcomes	High risk	No description
Incomplete outcome data (attrition bias) All outcomes	Low risk	Lost to follow-up 2/492 (< 1%)
Selective reporting (reporting bias)	Low risk	Reported all outcomes that were pre-stated in the methods section

Merad 1999 *(Continued)*

Other bias	Unclear risk	No reported substantial baseline differences in the treatment and control groups
		Sample size calculation not described. Source of bias

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

Study	Reason for exclusion
Brown 2001	Quasi-randomisation
Hoffmann 1987	Quasi-randomisation
Johnson 1989	Quasi-randomisation
Sagar 1993	Participants undergoing emergency surgery
Sagar 1995	Participants undergoing emergency surgery

Characteristics of studies awaiting assessment *[ordered by study ID]*
Hagmuller 1990

Methods	Randomised prospective clinical study
Participants	Prophylactic drainage for elective resection of the colon (intra-peritoneal)
Interventions	Capillar drainage (easy-flow-drainage) (n = 60) vs. no drainage(n = 53)
Outcomes	Anastomotic leakage; impaired wound healing; re-laparotomy
Notes	Unable to source original paper from German authors; awaiting assessment

n: number of participants.

DATA AND ANALYSES
Comparison 1. All studies stratified by type of drain

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Clinical anastomotic dehiscence	2	809	Risk Ratio (M-H, Fixed, 95% CI)	1.40 [0.45, 4.40]
1.1 Active drainage	1	492	Risk Ratio (M-H, Fixed, 95% CI)	1.98 [0.50, 7.84]
1.2 Active and passive drainage	1	317	Risk Ratio (M-H, Fixed, 95% CI)	0.52 [0.05, 5.63]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
2 Mortality	3	908	Risk Ratio (M-H, Fixed, 95% CI)	0.77 [0.41, 1.45]
2.1 Passive drainage	1	99	Risk Ratio (M-H, Fixed, 95% CI)	0.47 [0.04, 5.02]
2.2 Active drainage	1	492	Risk Ratio (M-H, Fixed, 95% CI)	0.79 [0.32, 1.98]
2.3 Active and passive drainage	1	317	Risk Ratio (M-H, Fixed, 95% CI)	0.80 [0.31, 2.10]
3 Re-intervention	3	908	Risk Ratio (M-H, Fixed, 95% CI)	1.11 [0.67, 1.82]
3.1 Passive drainage	1	99	Risk Ratio (M-H, Fixed, 95% CI)	0.71 [0.17, 2.99]
3.2 Active drainage	1	492	Risk Ratio (M-H, Fixed, 95% CI)	1.65 [0.83, 3.31]
3.3 Active and passive drainage	1	317	Risk Ratio (M-H, Fixed, 95% CI)	0.69 [0.29, 1.64]
4 Radiological anastomotic dehiscence	2	809	Risk Ratio (M-H, Fixed, 95% CI)	0.85 [0.39, 1.83]
4.1 Passive drainage	0	0	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
4.2 Active drainage	1	492	Risk Ratio (M-H, Fixed, 95% CI)	0.91 [0.41, 2.02]
4.3 Active and passive drainage	1	317	Risk Ratio (M-H, Fixed, 95% CI)	0.34 [0.01, 8.38]
5 Wound infection	3	908	Risk Ratio (M-H, Fixed, 95% CI)	0.82 [0.45, 1.51]
5.1 Passive drainage	1	99	Risk Ratio (M-H, Fixed, 95% CI)	1.88 [0.18, 20.09]
5.2 Active drainage	1	492	Risk Ratio (M-H, Fixed, 95% CI)	0.71 [0.32, 1.56]
5.3 Active and passive drainage	1	317	Risk Ratio (M-H, Fixed, 95% CI)	0.88 [0.30, 2.57]

Analysis 1.1. Comparison 1 All studies stratified by type of drain, Outcome 1 Clinical anastomotic dehiscence.

Analysis 1.2. Comparison 1 All studies stratified by type of drain, Outcome 2 Mortality.

Analysis 1.3. Comparison 1 All studies stratified by type of drain, Outcome 3 Re-intervention.

Analysis 1.4. Comparison 1 All studies stratified by type of drain, Outcome 4 Radiological anastomotic dehiscence.

Analysis 1.5. Comparison 1 All studies stratified by type of drain, Outcome 5 Wound infection.

Comparison 2. All studies stratified by level of anastomosis

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Clinical anastomotic dehiscence	2	416	Risk Ratio (M-H, Fixed, 95% CI)	0.66 [0.11, 3.94]
1.1 Intra-peritoneal anastomosis	2	416	Risk Ratio (M-H, Fixed, 95% CI)	0.66 [0.11, 3.94]
2 Mortality	2	416	Risk Ratio (M-H, Fixed, 95% CI)	0.74 [0.30, 1.80]
2.1 Intra-peritoneal anastomosis	2	416	Risk Ratio (M-H, Fixed, 95% CI)	0.74 [0.30, 1.80]
3 Re-intervention	2	416	Risk Ratio (M-H, Fixed, 95% CI)	0.71 [0.31, 1.62]
3.1 Intra-peritoneal anastomosis	2	416	Risk Ratio (M-H, Fixed, 95% CI)	0.71 [0.31, 1.62]
4 Radiological anastomotic dehiscence	1	317	Risk Ratio (M-H, Fixed, 95% CI)	0.34 [0.01, 8.38]
4.1 Intra-peritoneal anastomosis	1	317	Risk Ratio (M-H, Fixed, 95% CI)	0.34 [0.01, 8.38]
5 Wound infection	2	416	Risk Ratio (M-H, Fixed, 95% CI)	1.01 [0.39, 2.66]
5.1 Intra-peritoneal anastomosis	2	416	Risk Ratio (M-H, Fixed, 95% CI)	1.01 [0.39, 2.66]

Analysis 2.1. Comparison 2 All studies stratified by level of anastomosis, Outcome 1 Clinical anastomotic dehiscence.

Analysis 2.2. Comparison 2 All studies stratified by level of anastomosis, Outcome 2 Mortality.

Analysis 2.3. Comparison 2 All studies stratified by level of anastomosis, Outcome 3 Re-intervention.

Analysis 2.4. Comparison 2 All studies stratified by level of anastomosis, Outcome 4 Radiological anastomotic dehiscence.

Analysis 2.5. Comparison 2 All studies stratified by level of anastomosis, Outcome 5 Wound infection.

APPENDICES

Appendix 1. CENTRAL search strategy

Cochrane Central Register of Controlled Trials (CENTRAL) (2015, Issue 2)

- #1 MeSH descriptor: [Colorectal Surgery] explode all trees
- #2 MeSH descriptor: [Digestive System Surgical Procedures] explode all trees
- #3 MeSH descriptor: [Colorectal Neoplasms] explode all trees
- #4 MeSH descriptor: [Rectum] explode all trees
- #5 MeSH descriptor: [Colon] explode all trees
- #6 MeSH descriptor: [Abdomen] explode all trees
- #7 ((colorect* or colon* or rect* or anal or anus or intestin* or bowel* or abdom*) and (surger* or resec* or operat*)):ti,ab,kw
- #8 (#1 or #2 or #3 or #4 or #5 or #6 or #7)
- #9 MeSH descriptor: [Anastomosis, Surgical] explode all trees
- #10 (anastomo*):ti,ab,kw
- #11 (#9 or #10)
- #12 MeSH descriptor: [Drainage] explode all trees
- #13 (drain* or dren* or suction*):ti,ab,kw
- #14 (#12 or #13)
- #15 (#8 and #11 and #14)

Appendix 2. MEDLINE search strategy

Ovid MEDLINE (1950 to 6 February 2015)

1. exp Colorectal Surgery/
2. exp Digestive System Surgical Procedures/
3. exp Colorectal Neoplasms/
4. exp Rectum/

5. exp Colon/
6. exp Abdomen/
7. ((colorect* or colon* or rect* or anal or anus or intestin* or bowel* or abdom*) and (surger* or resec* or operat*)).mp.
8. 1 or 2 or 3 or 4 or 5 or 6 or 7
9. exp Anastomosis, Surgical/
10. anastomo*.mp.
11. 9 or 10
12. exp Drainage/
13. (drain* or dren* or suction*).mp.
14. 12 or 13
15. 8 and 11 and 14
16. randomized controlled trial.pt.
17. controlled clinical trial.pt.
18. randomized.ab.
19. placebo.ab.
20. clinical trial.sh.
21. randomly.ab.
22. trial.ti.
23. 16 or 17 or 18 or 19 or 20 or 21 or 22
24. humans.sh.
25. 23 and 24
26. 15 and 25

Appendix 3. EMBASE search strategy

Ovid EMBASE (1974 to 6 February 2015)

1. exp colorectal surgery/
2. exp abdominal surgery/
3. exp colon tumor/
4. exp rectum tumor/
5. exp rectum/
6. exp colon/
7. ((colorect* or colon* or rect* or anal or anus or intestin* or bowel* or abdom*) and (surger* or resec* or operat*)).mp.
8. 1 or 2 or 3 or 4 or 5 or 6 or 7
9. exp anastomosis/
10. anastomo*.mp.
11. 9 or 10
12. exp surgical drainage/
13. (drain* or dren* or suction*).mp.
14. 12 or 13
15. 8 and 11 and 14
16. CROSSOVER PROCEDURE.sh.
17. DOUBLE-BLIND PROCEDURE.sh.
18. SINGLE-BLIND PROCEDURE.sh.
19. (crossover* or cross over*).ti,ab.
20. placebo*.ti,ab.
21. (doubl* adj blind*).ti,ab.
22. allocat*.ti,ab.
23. trial.ti.
24. RANDOMIZED CONTROLLED TRIAL.sh.
25. random*.ti,ab.
26. 16 or 17 or 18 or 19 or 20 or 21 or 22 or 23 or 24 or 25
27. (exp animal/ or exp invertebrate/ or animal.hw. or nonhuman/) not (exp human/ or human cell/ or (human or humans or man or men or wom?n).ti.)
28. 26 not 27
29. 15 and 28

WHAT'S NEW

Date	Event	Description
16 March 2016	Amended	No new RCTs included in this update. Methodology reported according to required MECIR standards.

HISTORY

Protocol first published: Issue 2, 2000

Review first published: Issue 4, 2004

Date	Event	Description
22 December 2015	Feedback has been incorporated	Editorial comments incorporated
25 March 2015	New citation required but conclusions have not changed	Updated Prisma Flow Chart with new search results. 2 authors have verified no new studies for inclusion from new search results. Conclusions remain unchanged in this update.
6 April 2014	Amended	Updated discussion and references
27 July 2013	New search has been performed	Updated text and results
22 January 2004	New citation required and conclusions have changed	Substantive amendment

CONTRIBUTIONS OF AUTHORS

RR: Took the lead in writing the review, was involved in selecting trials for inclusion, performed independent data extraction and quality assessment of the included trials, and was responsible for statistical analysis and interpretation of the data.

JD: Involved in selecting trials for inclusion, performed independent data extraction and quality assessment of the included trials, and was responsible for statistical analysis and interpretation of the data.

SA: Involved in selecting trials and quality assessment of included trials.

PN: Commented on drafts of the review, and added clinical expertise to the discussion.

RN: Commented on drafts of the review, and added clinical expertise to the discussion.

DECLARATIONS OF INTEREST

None known.

SOURCES OF SUPPORT

Internal sources

- No sources of support supplied

External sources

- NIHR Cochrane Incentive Award, Reference Number 14/175/53, UK.

This project was supported by the National Institute for Health Research, via Cochrane Incentive Award funding to the Cochrane Colorectal Cancer Group. The views and opinions expressed therein are those of the authors and do not necessarily reflect those of the Systematic Reviews Programme, NIHR, NHS or the Department of Health.

INDEX TERMS

Medical Subject Headings (MeSH)

*Drainage; Anastomosis, Surgical [adverse effects]; Colon [*surgery]; Digestive System Surgical Procedures; Randomized Controlled Trials as Topic; Rectum [*surgery]; Surgical Wound Dehiscence [*prevention & control]; Surgical Wound Infection [*prevention & control]

MeSH check words

Humans