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META-ANALYSIS

Impact of mechanical bowel preparation in elective colorectal surgery: A meta-analysis

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Abstract

AIM

To analyse the effect of mechanical bowel preparation *vs* no mechanical bowel preparation on outcome in patients undergoing elective colorectal surgery.

METHODS

Meta-analysis of randomised controlled trials and observational studies comparing adult patients receiving mechanical bowel preparation with those receiving no mechanical bowel preparation, subdivided into those receiving a single rectal enema and those who received no preparation at all prior to elective colorectal surgery.

RESULTS

A total of 36 studies (23 randomised controlled trials and 13 observational studies) including 21568 patients undergoing elective colorectal surgery were included. When all studies were considered, mechanical bowel preparation was not associated with any significant difference in anastomotic leak rates (OR = 0.90, 95%CI: 0.74 to 1.10, P = 0.32), surgical site infection (OR = 0.99, 95%CI: 0.80 to 1.24, P = 0.96), intraabdominal collection (OR = 0.86, 95%CI: 0.63 to 1.17, P = 0.34), mortality (OR = 0.85, 95%CI: 0.57 to 1.27, P = 0.43), reoperation (OR = 0.91, 95%CI: 0.75 to 1.12, P = 0.38) or hospital length of stay (overall mean difference 0.11 d, 95%CI: -0.51 to 0.73, P = 0.72), when compared with no mechanical bowel preparation, nor when evidence from just randomized controlled

trials was analysed. A sub-analysis of mechanical bowel preparation vs absolutely no preparation or a single rectal enema similarly revealed no differences in clinical outcome measures.

CONCLUSION

In the most comprehensive meta-analysis of mechanical bowel preparation in elective colorectal surgery to date, this study has suggested that the use of mechanical bowel preparation does not affect the incidence of postoperative complications when compared with no preparation. Hence, mechanical bowel preparation should not be administered routinely prior to elective colorectal surgery.

Key words: Bowel preparation; Mechanical; Antibiotics; Morbidity; Mortality; Surgery; Outcome complications; Meta-analysis

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Core tip: At present there is no evidence that bowel preparation makes a difference to clinical outcomes in either colonic or rectal surgery, in terms of anastomotic leak rates, surgical site infection, intra-abdominal collection, mortality, reoperation or hospital length of stay. Given its potential adverse effects and patient dissatisfaction rates, it should not be administered routinely to patients undergoing elective colorectal surgery.

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INTRODUCTION

Mechanical bowel preparation (MBP) for colorectal surgery has been surgical dogma for decades, despite increasing evidence from the 1990s refuting its benefits^[1,2]. The rationale behind the administration of MBP is that it reduces fecal bulk and, therefore, bacterial colonisation, thereby reducing the risk of postoperative complications such as anastomotic leakage and wound infection^[3], as well as to facilitate dissection and allow endoscopic evaluation. Opponents argue that in the 21st century, with rational use of oral and intravenous prophylactic antibiotics there is no longer a place for MBP, that it may cause marked fluid and electrolyte imbalance in the preoperative period, and that evidence has shown that the gut microbial flora load is not reduced grossly by bowel preparation^[4]. There is also concern that bowel preparation liquefies feces, thereby increasing the risk of spillage and contamination intraoperatively^[5]. Its use remains controversial, particularly within the context of an enhanced recovery after surgery (ERAS) program setting^[6,7].</sup>

Meta-analyses^[8-12] have been published on MBP in elective colorectal surgery showing mixed results, with most studies demonstrating no difference in infective complications between patients receiving MBP or control treatment, although control treatment varied significantly between the use of a rectal enema or absolutely no preparation. Similar results have been found in gynaecological^[13,14] and urological^[15,16] surgery where studies have shown no benefits in visualisation, bowel handling or complication rates between patients treated with bowel preparation and those given no bowel preparation. As a result of this inconclusive evidence, several studies have established that practice varies significantly between countries, and even surgeons in the same institution^[17,18]. Further impediments to the issue are that no consensus has yet been reached regarding the optimal method of bowel cleansing. Various agents such as polyethylene glycol (PEG), sodium phosphate, mannitol, milk of magnesia, liquid paraffin and senna have been used to achieve bowel cleansing.

Infective complications are amongst the leading causes of morbidity and mortality in patients undergoing colorectal surgery^[19]. However, MBP is not without its own complications and the process is both time-consuming and unpleasant for patients^[20]. It has been shown to cause clinically significant dehydration^[21] and electrolyte disturbances, particularly hypocalcaemia and hypokalaemia to which the elderly are especially vulnerable^[22-24]. Patient satisfaction is poor for undergoing bowel preparation prior to surgery and colonoscopy, and this may necessitate an additional day preoperatively in hospital, particularly for frail elderly patients.

In the United Kingdom, the National Institution of Health and Clinical Excellence (NICE) does not recommend using MBP routinely to reduce the risk of surgical site infection (SSI)^[25] and the ERAS[®] Society guidelines on perioperative care of patients undergoing colonic resection^[6] also recommend against using preoperative bowel preparation. However, for rectal^[7] resection the recommendation, albeit weak, is to use MBP for patients undergoing anterior resection with diverting stomas. In recent years further evidence has emerged from large database studies using the National Surgical Quality Improvement (NSQIP) database in America^[26-29] showing reduced rates of anastomotic leakage, intra-abdominal abscess formation and wound infection when patients were given MBP with intraluminal antibiotics pre-operatively.

We have assessed this expanding body of evidence in this new comprehensive meta-analysis encompassing both randomised controlled trials and observational studies. We sought to address deficiencies in previous studies by including all levels of evidence, separating those in which patients received a single rectal enema *vs* full or no preparation, and including the recently



published large database studies.

Our aims for this meta-analysis were: (1) To analyse the effect of MBP *vs* no preparation or rectal enema alone on postoperative infective complications in patients undergoing elective colorectal surgery; (2) To examine the differences in results between evidence obtained from randomised controlled trials and observational studies; and (3) To determine what effect, if any, bowel preparation had on postoperative complications in rectal surgery.

MATERIALS AND METHODS

Search Strategy

We performed an electronic search of the PubMed database and the Cochrane Central Register of Controlled Trials to identify studies comparing outcomes in patients undergoing elective colorectal surgery treated with MBP vs either no preparation or a single rectal enema (last search on 1st May 2017). We used the search terms "(bowel preparation OR bowel cleansing OR bowel cleaning) AND (surgery OR preoperative)". Further sources were obtained by a manual search of the bibliography of the papers obtained to ensure the search was as comprehensive as possible. We did not apply language restriction or time limitations. Two independent researchers (KER and HJ-E) reviewed the abstracts for inclusion. Where there was a difference of opinion on the inclusion of papers, the opinion of the senior author was sought (DNL). We performed this meta-analysis according to the Preferred Reporting Items for Systematic Reviews and Metaanalyses (PRISMA)^[30] and Guidelines for Meta-Analyses and Systematic Review of Observational Studies (MOOSE) statements^[31].

Selection of articles

We reviewed full text articles for suitability after excluding studies on the basis of title and abstract. Our inclusion criteria specified that studies must have a minimum of two comparator groups and were either designed as randomised controlled trials or observational studies. Publications comparing preoperative MBP with no preparation or a single rectal enema were included and comparisons with other forms of bowel preparation (*e.g.* intraoperative colonic lavage) were excluded. Only studies on adult patients undergoing elective colorectal surgery were included. We included studies on laparoscopic and open surgical procedures but excluded endoscopic studies. Relevant outcome measures were anastomotic leak, SSI, intra-abdominal abscess, mortality, reoperation and hospital length of stay.

Duplication of results was a particular hazard encountered when selecting which of the studies to include that extracted information from the NSQIP database^[26-29,32-36]. The papers were scrutinised for their enrollment dates. There was overlap in these dates and after correspondence with the authors, it was apparent that there was considerable overlap in the data sets used. Hence, we selected the largest study for inclusion with the greatest number of clinically relevant outcome measures^[29]. Two further studies^[37,38] had duplication of results and in this situation the larger of the two studies was included^[38]. One study^[39] was a subgroup analysis of patients undergoing anastomosis below the peritoneal reflection taken from a study which was already included^[40] in the meta-analysis so this was excluded from the main meta-analysis to prevent dual inclusion of patients. However, this subgroup was included in the separate analysis of rectal surgery. A further study^[41] reviewed as a full text article was retracted since its inclusion in the 2011 Cochrane Review^[10], so we chose to exclude this. One paper^[2] analysed in the Cochrane Review included pediatric patients and so has been excluded from our meta-analysis.

Data extraction

HJ-E extracted the data and they were verified independently by KER. Quantitative data relevant to the endpoints we selected were extracted. Several studies presented hospital length of stay results in formats other than mean and standard deviation. Where this occurred, the authors were contacted for the raw data in order to ascertain the mean and standard deviation necessary for creation of Forest plot. When the raw data were unavailable, mean and standard deviation were calculated using the technique described by Hozo *et al*^[42].

Risk of bias and completeness of reporting of individual studies

The risk of bias was assessed using the Cochrane Collaboration tool in RevMan 5.3^[43], which focuses upon random sequence generation (selection bias), allocation concealment (selection bias), blinding of participants and personnel (performance bias), blinding of outcome assessment (detection bias), incomplete outcome data (attrition bias) and selective reporting (reporting bias).

Statistical analysis

The analysis was performed using RevMan 5.3 software^[43]. Continuous variables were calculated as a mean difference and 95% confidence interval using an inverse variance random effects model. Dichotomous variables were analysed using the Mantel-Haenszel random effects model to quote the risk ratio (RR) and 95% confidence interval. These analyses were used to construct forest plots, with statistical significance taken to be a *P* value of < 0.05 on two tailed testing. A predetermined subgroup analysis was performed for the impact of MBP in rectal surgery specifically using the same methodology. Study inconsistency and heterogeneity were assessed using the I^2 statistic^[44].

Protocol registration

The protocol for this meta-analysis was registered



Ref.	Random sequence generation	Allocation concealment	Blinding of participants and personnel	Blinding of outcome assessment	Incomplete outcome data	Selective reporting
Ji et al ^[76]	NA	NA	NA	NA	NA	NA
Chan et al ^[77]	NA	NA	NA	NA	NA	NA
Hu et al ^[64]	?	?	?	?	?	?
Bhattacharjee et al ^[65]	+	?	-	?	?	?
Allaix et al ^[74]	NA	NA	NA	NA	NA	NA
Kiran et al ^[29]	NA	NA	NA	NA	NA	NA
Yamada <i>et al</i> ^[66]	NA	NA	NA	NA	NA	NA
Otchy et al ^[67]	NA	NA	NA	NA	NA	NA
Kim et al ^[75]	NA	NA	NA	NA	NA	NA
Tahirkheli <i>et al</i> ^[62]	+	?	?	?	-	-
Sasaki <i>et al</i> ^[61]	+	?	?	?	?	?
Bertani et al ^[45]	+	+	?	?	+	+
Roig et al ^[68]	NA	NA	NA	NA	NA	NA
Bretagnol et al ^[46]	+	+	+	+	-	+
Pitot et al ^[69]	NA	NA	NA	NA	NA	NA
Alcantara Moral et al ^[47]	+	+	?	?	?	+
Miron <i>et al</i> ^[70]	NA	NA	NA	NA	NA	NA
Pena-Soria et al ^[48]	+	+	+	+	-	+
Leiro <i>et al</i> ^[59]	+	+	?	?	?	+
Contant et al ^[40]	+	+	- (2)	- (2)	-	+
Bretagnol et al ^[71]	NA	NA	NA	NA	NA	NA
Jung et al ^[49]	+	+	+	+	-	?
Veenhof <i>et al</i> ^[72]	NA	NA	NA	NA	NA	NA
Ali et al ^[63]	?	?	?	?	?	?
Jung et al ^[50]	+	+	+	+	-	?
Platell et al ^[51]	+	+	+	+	-	-
Fa-Si-Oen et al ^[52]	+	+	?	?	+	+
Bucher <i>et al</i> ^[53]	+	+	+	+	+	+
Ram et al ^[54]	+	- (1)	?	?	?	+
Zmora et al ^[37]	+	+	?	?	_	+
Young Tabusso <i>et al</i> ^[55]	?	?	- (2)	- (2)	?	?
Miettinen <i>et al</i> ^[56]	+	+	?	?	+	+
Memon et al ^[73]	NA	NA	NA	NA	NA	NA
Fillmann <i>et al</i> ^[60]	+	+	+	+	+	+
Burke <i>et al</i> ^[57]	?	?	+	+	-	_
Brownson <i>et al</i> ^[58]	?	?	?	?	?	?

NA: Not applicable (observational study); +: Low risk of bias; -: High risk of bias; (1): Allocation concealment utilized identification number of patient (odd or even); (2): Not blinded.

with the PROSPERO database (www.crd.york.ac.uk/ prospero) - registration number CRD42015025279.

RESULTS

From 1594 studies identified from the original search, 97 were reviewed as full text articles. Of these, 36 comprising 23^[37,40,45-65] randomised controlled trials and 13 observational studies^[29,66-77] were eligible for inclusion (Figure 1). The risk of bias of the randomised controlled trials included in this study was moderate (Table 1).

Patient demographics

Overall, 21568 patients were included in the metaanalysis, of whom 6166 had no bowel preparation of any sort, 2739 had a solitary rectal enema and 12663 underwent full MBP as per local policy. Of these, 6277 patients were included in randomised controlled trials and 15291 in observational studies. Demographic details are summarised in Table 2 and of details of interventions (bowel preparation and perioperative antibiotics) in Table 3.

Anastomotic leak

All studies except one^[75] included data on the primary outcome measure of this meta-analysis, the incidence of anastomotic leak (Figure 2). When MBP was compared with no MBP (including no preparation at all and those who underwent a single rectal enema), there was no difference in the incidence of anastomotic leak (OR = 0.90, 95%CI: 0.74 to 1.10, P = 0.32). When MBP vs absolutely no MBP was analysed^[29,40,46,48-50,52,54-65,68,70,71,73], this made no difference to anastomotic leak rates (OR 0.94, 95% CI 0.70 to 1.25, P = 0.67), nor when MBP was compared with a single rectal enema^[37,45,47,51,53,66,67,69,72,74,76,77] (OR = 0.92, 95%CI: 0.70 to 1.20, P = 0.52).

When randomised controlled trials alone were included in the analysis^[37,40,45-65] (Supplementary Figure 1A), the use of MBP *vs* no MBP did not affect the incidence of anastomotic leak (OR = 1.02, 95%CI: 0.75



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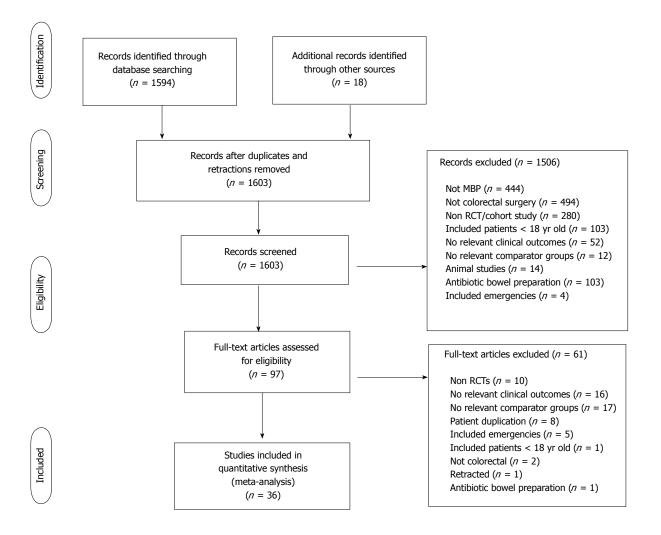


Figure 1 PRISMA diagram showing identification of relevant studies from initial search, PRISMA: Preferred reporting Items for systematic reviews and metaanalyses

to 1.40, P = 0.90), nor when MBP vs absolutely no MBP^[40,46,48-50,52,54-65] or MBP vs single rectal enema^{[37,45,47,} ^{51,53]}were considered. When observational studies alone were analysed^[66-73,76,77] (Supplementary Figure 1B), the use of MBP vs no MBP did significantly affect the incidence of anastomotic leak (OR = 0.76, 95%CI: 0.63 to 0.91, P = 0.003), although this was not significant when MBP vs single rectal enema^[66,67,69,72,74,76,77] and MBP vs absolutely no MBP^[29,68,70,71,73] were considered separately.

SSI

Data on the incidence of SSI were presented in a total of 19780 patients in 32 studies^[29,37,40,45-61,64-70,72-75,77] (Figure 3). There was no difference in the incidence of SSI in those who did vs those who did not undergo MBP (OR = 0.99, 95%CI: 0.80 to 1.24, P = 0.96), nor in those who had MBP vs those receiving a single rectal enema^[37,45,47,51,53,66,67,69,72,74,77] (OR = 1.00, 95%CI: 0.57 to 1.76, P = 1.00) or those who had MBP vs those receiving absolutely no preparation^[29,40,46,48-50,52,54-61,64,65,68,70,73,75] (OR = 0.98, 95%CI: 0.78 to 1.24, P = 0.87).

When data obtained from 21 randomised controlled trials^[37,40,43,45-61,64,65] alone with a total of 5971 patients were included (Supplementary Figure 2A), the use of MBP vs no MBP did not impact upon the incidence of SSI (OR = 1.16, 95%CI: 0.96 to 1.39, P = 0.12), nor when MBP vs single rectal enema[37,45,47,51,53] or MBP vs absolutely no preparation^[40,43,46,48-50,52,54-61,64,65] were considered. When just observational studies were included^[29,66-70,72-75,77] (11 studies, 13809 patients; Supplementary Figure 2B), patients who received MBP had a significantly reduced incidence of SSI than those who did not receive MBP (OR = 0.64, 95%CI: 0.55 to 0.75, P < 0.0001), with similar results seen in those who received MBP vs absolutely no MBP^[29,68,70,73,75], although no difference was seen between those who received full MBP vs a single rectal enema^[66,67,69,72,74,77].

Intra-abdominal collection

A total of 29 studies^[29,37,40,45,46,48,49,51,53-56,58,59,61,62,64-75,77] on 19327 patients included data on postoperative intraabdominal collections (Figure 4). The administration of

\mathbf{k} k	Table 2 Baseline patie	ent demog	Baseline patient demographics for all studies included	udies ind	luded							
	Ref.	Year published	Study methodology	Study	numbers	Male: Fema	lle gender	Indication for surgery	Location	Primary anastomosis	Laparoscopi	c approach
				MBP, n		MBP	No MBP				MBP, n	No MBP, n
	Ji <i>et al</i> ^[76]	2017	Observational	538	831	Unknown	Unknown	Cancer	Rectum	Υ	Unknown	Unknown
	$\operatorname{Chan} et al^{[m]}$	2016	Observational	159	97	85:74	55:42	Cancer	Colon and rectum	Υ	159	67
	Hu et al ^[64]	2017	RCT	76	72	Unknown	Unknown	Cancer	Colon and rectum	Υ	Unknown	Unknown
	Bhattacharjee et al ^[65]	2015	RCT	38	33	21:17	20:13	Cancer, inflammatory bowel disease, volvulus,	Colon and rectum	Y	0	0
	A 11 - 11 - 11 - 11 74	3 100		202	000	361.045	700.004	tuberculosis		>	000	
	AllarX et al	C102	Observational	907	678	361:345	432:39/	Cancer, agenoma, diverticulitis, reversal of	Colon and rectum	Y	828	907
	[67]1- T74	304 5	Chanter	24.62	2000	2000-00116	1111.1105	Hartmann's procedure, rectal prolapse	materia Proceeding	14	CF F F	1780
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Niran $et at$	C102	Observational	0140	0677	9115:USUS	C011:1111		Colon and rectum	z >	4440 70	69CT
align and align	$O_{tchyr} = a^{l[6]}$	2014	Observational	761	00T	30.47	30-40	Cancer dittarticular diseases IBD radal nuclanea	Colon and rectium	- >	16 27	18 18
		E T 07	In to the to the top of top of the top of top o	8		ECO	DE-CC	ischemic colitis, volvulus, colovaginal fistula		4	5	ê
al2113RCT484822.022.21Cancer diverticular disease, IBD is chemic collisColon and rectunYunbown2011RCT1311365.369.3569.35Cancer diverticular disease, IBD is chemic collisY122010Observational396970.3Rectal disease, IBD is chemic collisColon and rectunY2312010Observational3999UhbownUhbownCancer diverticular disease, IBD is chemic collisY242010Observational69127312887.3Cancer diverticular disease, IBD is chemic collisY232010Observational6099UhbownUhbownCancer diverticular disease, IBD is chemic collisY2320112018Observational613332.2Bengin and meturnY14201606999131.3Cancer diverticular disease, IBD is chemic collisY2320172018Observational613332.3Bengin and meturnY10201706663033.3ServerCancer diverticular disease, IBD is choose and rectunY10201706663033.4Cancer diverticular disease, IBD is choose and rectunY10201706663033.4Cancer diverticular disease, IBD is choose and rectunY10201706663033.4	$\operatorname{Kim} et al^{[5]}$	2014	Observational	1363	1112	502:694	669:610	Unknown	Colon and rectum	X	209	472
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Tahirkheli <i>et al</i> ^[62]	2013	RCT	48	48	28:20	24:24	Cancer, diverticular disease, IBD, ischemic colitis	Colon and rectum	Y	unknown	unknown
	Sasaki et al ^[61]	2012	RCT	38	41	17:21	24:17	Cancer	Colon only	Y	29	19
$ \begin{array}{{ccccccccccccccccccccccccccccccccccc$	Bertani <i>et al</i> ^[45]	2011	RCT	114	115	65:49	60:55	Cancer	Colon and rectum	Х	55	51
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Roig et al ^[68]	2010	Observational	39	69	Unknown	Unknown	Cancer, diverticular disease, IBD, FAP	Colon and rectum	γ	12	20
	Bretagnol et al ^[46]	2010	RCT	89	89	56:33	46:43	Rectal cancer	Rectum only	Υ	73	74
$ \left[t_{\rm eff} \left[\begin{array}{cccccccccccccccccccccccccccccccccccc$	Pitot <i>et al</i> ^[69]	2009	Observational	59	127	31:28	53:74	Cancer, diverticular disease, IBD	Colon only	γ	26	30
41 2008Cheservational6039UnknownUnknownUnknownUnknownUnknownYUnknown2008RCT656435/230032/2338/2334/2397/20Colon and rectumYUnknown2007RCT67068437/33334/3390Cancer, IBDColon onlyYUnknown2007Observational615239/2334/33937/340Cancer, IBDColon onlyYUnknown2007Observational787128/334/359Cancer, RibnColon onlyYUnknown2007Observational787128/333/45Not specified cancerRectur onlyYUnknown2007Observational787128/333/45Not specified cancerColon onlyYUnknown2007RCT147147UnknownUnknownCancer, diverticular disease, ademonaColon and rectumYUnknown2006RCT147147UnknownUnknownCancer, diverticular disease, ademonaYUnknown2008RCT147147UnknownCancer, diverticular disease, ademonaYUnknown2008RCT147147UnknownCancer, diverticular disease, ademonaYUnknown2008RCT147147UnknownCancer, diverticular disease, ademonaYUnknown2008RCT143243<	Alcantara Moral <i>et al</i> ^[47]	2009	RCT	70	69	41:28	48:22		Left colon and rectum	Y	12	15
***2008RCT656433:25:0033:22Cancer, IBDColon and rectumYUhknown12007RCT67645532:3934:33Marganat colorectal pathologyColon and rectumYUhknown2007RCT665730:38331:340Cancer, IBDColon and rectumYNone2007Observational615242:1932:20Rectal cancerRectal cancerRectum onlyYNone2007Observational615242:1932:40Cancer, silverticular disease, adenomaColon and rectumYUnknown2007Observational615242:1932:40Cancer, silverticular disease, adenomaYUnknown2007RCT20RCT147UnknownUnknownColon and rectumYUnknown2008RCT147147UnknownUnknownCancer, diverticular disease, adenomaYUnknown2008RCT147147UnknownUnknownColon and rectumYUnknown2008RCT147147UnknownUnknownColon and rectumYUnknown2008RCT147147UnknownUnknownColon and rectumYUnknown2008RCT147147UnknownUnknownColon and rectumYUnknown2008RCT143144Cancer, diverticular disease, BIDCo	Miron <i>et al</i> ^[70]	2008	Observational	60	39	Unknown	Unknown	Unknown	Colon and rectum	Y	Unknown	Unknown
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Pena-Soria <i>et al</i> ^[48]	2008	RCT	65	64	35:29:00	33:22	Cancer, IBD	Colon and rectum	Υ	Unknown	Unknown
	Leiro et al ^[59]	2008	RCT	64	65	39:25	38:27	Benign and malignant colorectal pathology	Colon and rectum	Z	Unknown	Unknown
	Contant <i>et al</i> ^[40]	2007	RCT	670	684	337:333	345:339	Cancer, IBD	Colon and rectum	Х	None	None
	Bretagnol et al ^[71]	2007	Observational	61	52	42:19	32:20	Rectal cancer	Rectum only	Y	Unknown	27
	Jung et al ^[49]	2007	RCT	686	657	306:380	317:340	Cancers, diverticular disease, adenoma	Colon only	Υ	None	None
207RCT109101UnknownUnknownUnknownUnknownYUnknown206RCT2717UnknownUnknownUnknownUnknownNoneYNone206RCT147UnknownUnknownUnknownCancer, diverticular diseaseRetum onlyYNone206RCT12512556:69Cancer, diverticular disease, adenomaColon and rectumNNUnknown205RCT12534:41Cancer, diverticular disease, reversal of Hartman'sLeft colon and rectumYNone206RCT16416599:65102:63Cancer, diverticular disease, reversal of Hartman'sLeft colon and rectumYNone207RCT187193103:8494:99Cancer, diverticular disease, reversal of Hartman'sLeft colon and rectumYUnknown208RCT187193103:8494:99Cancer, diverticular disease, IBDColon and rectumYUnknown200RCT187193103:8494:99Cancer, diverticular disease, IBDColon and rectumYUnknown200RCT187193103:8494:99Cancer, diverticular disease, IBDColon and rectumYUnknown200RCT187133123:63Cancer, diverticular disease, IBDColon and rectumYUnknown200RCT2423122:67Cancer, diverticular disease, IBD <td< td=""><td>Veenhof et al^[72]</td><td>2007</td><td>Observational</td><td>78</td><td>71</td><td>28:43</td><td>33:45</td><td>Not specified</td><td>Colon and rectum</td><td>Υ</td><td>Unknown</td><td>Unknown</td></td<>	Veenhof et al ^[72]	2007	Observational	78	71	28:43	33:45	Not specified	Colon and rectum	Υ	Unknown	Unknown
206RCT2717UnknownCancer, adenoma and diverticular diseaseRectum onlyYNone2005RCT147UnknownUnknownUnknownCancer, daenoma and diverticular diseaseRectum onlyYNone2005RCT12558.6756.69Cancer, diverticular disease, adenomaColon and rectumNVNone2005RCT12538.6756.69Cancer, diverticular disease, adenoma, colon and rectumYNone2005RCT12538.6756.69Cancer, diverticular disease, adenoma, elf colon and rectumYNone2005RCT16416599.55102.63Cancer, diverticular disease, IBDColon and rectumYUnknown2002RCT13812968.7062.67Cancer, diverticular disease, IBDColon and rectumYUnknown2002RCT13812968.7062.67Cancer, diverticular disease, IBDColon and rectumYUnknown2002RCT13812968.7062.67Cancer, BD, diverticular disease, IBDColon and rectumYUnknown2001RCT13812968.7062.67Cancer, BD, diverticular disease, IBDColon and rectumYUnknown2002RCT13812968.7062.67Cancer, BD, diverticular disease, IBDColon and rectumYUnknown2003RCT3030UnknownUnknownColon and	Ali et al ^[63]	2007	RCT	109	101	Unknown	Unknown	Unknown	Colon and rectum	Х	Unknown	Unknown
2006RCT147147UnknownUnknownUnknownUnknownNUnknown2005RCT12558:6756:69Cancer, diverticular disease, adenomaColon onlyYNone2005RCT787547:3134:41Cancer, diverticular disease, reversal of Hartman/sLeft colon and rectumYNone2005RCT18719599:65102:63Cancer, diverticular disease, reversal of Hartman/sLeft colon and rectumYNone2003RCT187193192:63Cancer, diverticular disease, BDColon and rectumYUnknown2003RCT187193103:8494:99Cancer, diverticular disease, BDColon and rectumYUnknown2003RCT187193103:8494:99Cancer, diverticular disease, BDColon and rectumYUnknown2003RCT18712994:99Cancer, diverticular disease, BDColon and rectumYUnknown2000RCT18712968:7062:67Cancer, diverticular disease, BDColon and rectumYUnknown2000RCT18712968:7062:67Cancer, diverticular disease, BDColon and rectumYUnknown2001RCT18712968:7062:67Cancer, diverticular disease, BDAlon and rectumYUnknown2002RCT3030UnknownCancer, diverticular disease, BD, ische	Jung et al ^[50]	2006	RCT	27	17	Unknown	Unknown	Cancer, adenoma and diverticular disease	Rectum only	Х	None	None
2005RCT12512558.6756.69Cancer, diverticular diseaseColon onlyYNone2005RCT787547.3134.41Cancer, diverticular disease, reversal of Hartman/'sLeft colon and rectumY202005RCT16416599.65102.63Cancer, diverticular disease, IBDColon and rectumY202003RCT187193103.8494.99Cancer, diverticular disease, IBDColon and rectumYUnknown2002RCT13812968.7062.67Cancer, diverticular disease, IBDColon and rectumYUnknown2002RCT13812968.7062.67Cancer, diverticular disease, IBDColon and rectumYUnknown2003RCT13812968.7062.67Cancer, diverticular disease, IBDColon and rectumYUnknown2004RCT13812968.7062.67Cancer, diverticular disease, IBDColon and rectumYUnknown2005RCT13812963.7062.67Cancer, diverticular disease, IBDColon and rectumYUnknown2007RCT30UnknownInknownColon and rectumYUnknown2008RCT30UnknownInknownColon and rectumYUnknown1995RCT8233UnknownInknownInknownInknownInknownInknown1992 <t< td=""><td>Platell <i>et al</i>^[51]</td><td>2006</td><td>RCT</td><td>147</td><td>147</td><td>Unknown</td><td>Unknown</td><td>Cancer, IBD, diverticular disease, adenoma</td><td>Colon and rectum</td><td>Z</td><td>Unknown</td><td>Unknown</td></t<>	Platell <i>et al</i> ^[51]	2006	RCT	147	147	Unknown	Unknown	Cancer, IBD, diverticular disease, adenoma	Colon and rectum	Z	Unknown	Unknown
2005RCT787547.3134.41Cancer, diverticular disease, reversal of Hartmann'sLeft colon and rectumY202005RCT16416599.65102.63Cancer, diverticular disease, BDColon and rectumYUnknown2003RCT187193103.8494.99Cancer, diverticular disease, BDColon and rectumYUnknown2003RCT187193103.8494.99Cancer, diverticular disease, BDColon and rectumYUnknown2002RCT242312:129:14UnknownColon and rectumYUnknown2000RCT13812962:67Cancer, IBD, diverticular disease, IBD, diverticular disease, IBD, and rectumYUnknown2001RCT13812962:67Cancer, IBD, diverticular disease, IBD, adenoma, lipona91% primaryNone2001RCT30UnknownNoneObservational617532:2944:31Cancer, diverticular disease, IBD, adenoma, liponaYUnknown1997Observational617532:2944:31Cancer, diverticular disease, IBD, ischemic colitisYUnknown1997RCT30UnknownUnknownCancer, diverticular disease, IBD, ischemic colitisYUnknown1994RCT8593UnknownUnknownCancer, diverticular disease, IBD, ischemic colitisYUnknown1992RCT8693 <td< td=""><td>Fa-Si-Oen <i>et al</i>^[52]</td><td>2005</td><td>RCT</td><td>125</td><td>125</td><td>58:67</td><td></td><td>Cancer, diverticular disease</td><td>Colon only</td><td>γ</td><td>None</td><td>None</td></td<>	Fa-Si-Oen <i>et al</i> ^[52]	2005	RCT	125	125	58:67		Cancer, diverticular disease	Colon only	γ	None	None
2005RCT16416599:65102:63Cancer diverticular disease, BDColon and rectumYUnknown2003RCT187193103:8494:99Cancer, diverticular disease, BDColon and rectumYUnknown2002RCT187193103:8494:99Cancer, diverticular disease, BDColon and rectumYUnknown2002RCT242312:129:14UnknownColon and rectumYUnknown2000RCT13812968:7062:67Cancer, IBD, diverticular disease, IBD, diverticular disease, IBD, and rectumYUnknown1997Observational617532:2944:31Cancer, diverticular disease, IBD, adenoma, lipomaPoth armsPoth arms1997Observational617532:2944:31Cancer, diverticular disease, IBD, ischemic colitisColon and rectumYUnknown1997RCT30UnknownUnknownCancer, diverticular disease, IBD, ischemic colitisColon and rectumYUnknown1994RCT8293UnknownUnknownCancer, diverticular disease, IBD, ischemic colitisYUnknown1992RCT8693UnknownUnknownCancer, diverticular disease, IBD, ischemic colitisYUnknown1994RCT8693UnknownUnknownCancer, diverticular disease, IBD, ischemic colitisYUnknown1994RCT8693 <td>Bucher <i>et al</i>^[53]</td> <td>2005</td> <td>RCT</td> <td>78</td> <td>75</td> <td>47:31</td> <td></td> <td>Cancer, diverticular disease, reversal of Hartmann's</td> <td>Left colon and rectum</td> <td>Υ</td> <td>20</td> <td>22</td>	Bucher <i>et al</i> ^[53]	2005	RCT	78	75	47:31		Cancer, diverticular disease, reversal of Hartmann's	Left colon and rectum	Υ	20	22
2003RCI10410599:30102:35Carnetr, cuverticular disease, IBUColon and rectumTUnknown2003RCT187193103:8494:99Cancer, diverticular disease, IBUColon and rectumYUnknown2002RCT187193103:8494:99Cancer, diverticular disease, IBDColon and rectumYUnknown2003RCT13812968:7062:67Cancer, IBD, diverticular disease, IBDColon and rectumYUnknown2000RCT13812968:7062:67Cancer, IBD, diverticular disease, IBD, diverticular disease, IBD, and rectumYUnknown1977Observational617532:2944:31Cancer, diverticular disease, IBD, adenoma, lipomaLeft colon and rectumYUnknown1997Observational617532:2944:31Cancer, diverticular disease, IBD, ischemic colitisColon and rectumYUnknown1997RCT30UnknownUnknownCancer, diverticular disease, IBD, ischemic colitisColon and rectumYUnknown1994RCT8593UnknownUnknownCancer, diverticular disease, IBD, ischemic colitisYUnknown1992RCT8693UnknownUnknownCancer, diverticular disease, IBD, ischemic colitisYUnknown1992RCT8693UnknownUnknownCancer, diverticular disease, IBD, ischemic colitisYUnknown	[5] [7]	1000	EC C		L T		07 001	procedure, adenoma, endometriosis		;	1 11	1 1
2003RCT187193103:8494:99Cancer, diverticular disease, IBDColon and rectumYUnknown2002RCT242312:129:14UnknownYUnknownYUnknown2000RCT13812968:7062:67Cancer, IBD, diverticular diseaseColon and rectumYUnknown2000RCT13812968:7062:67Cancer, IBD, diverticular diseaseColon and rectum91% primaryNone1977Observational617532:2944:31Cancer, diverticular disease, IBD, adenoma, lipomaLeft colon and rectumYUnknown1997RCT3030UnknownCancer, diverticular disease, IBD, ischemic colitisColon and rectumYUnknown1994RCT8252:3043:44Cancer, diverticular disease, IBD, ischemic colitisColon and rectumYUnknown1992RCT8693UnknownUnknownCancer, diverticular disease, IBD, ischemic colitisColon and rectumYUnknown1992RCT8693UnknownUnknownCancer, diverticular disease, IBDColon and rectumYUnknown	Kam et al	CUU2	IN	10 1	COT	C0:66	C0:701	Cancer, uiverncular uisease, IDD	Colon and rectum	I	UNKNOWN	UNKNOWN
2002 RCT 24 23 12:12 9:14 Unknown Colon and rectum Y Unknown 2000 RCT 138 129 68:70 62:67 Cancer, IBD, diverticular disease Colon and rectum 91% primary None 2000 RCT 138 129 68:70 62:67 Cancer, IBD, diverticular disease Colon and rectum 91% primary None 1997 Observational 61 75 32:29 44:31 Cancer, diverticular disease, IBD, ischemic colitis Colon and rectum Y Unknown 1995 RCT 30 Unknown Unknown Cancer, diverticular disease, IBD, ischemic colitis Colon and rectum Y Unknown 1994 RCT 82 93 Unknown Unknown Cancer, diverticular disease, IBD, ischemic colitis Colon and rectum Y Unknown 1992 RCT 86 93 Unknown Cancer, diverticular disease, IBD Left colon and rectum Y Unknown	Zmora <i>et al</i> ^{$[37]$}	2003	RCT	187	193	103:84	94:99	Cancer, diverticular disease, IBD	Colon and rectum	Х	Unknown	Unknown
2000 RCT 138 129 68.70 62.67 Cancer, IBD, diverticular disease Colon and rectum 91% primary None 1997 Observational 61 75 32.29 44:31 Cancer, diverticular disease, IBD, adenoma, lipoma Left colon and rectum Y Unknown 1995 RCT 30 30 Unknown Unknown Cancer, diverticular disease, IBD, ischemic colitis Colon and rectum Y Unknown 1994 RCT 82 87 52:30 43:44 Cancer, diverticular disease, IBD, ischemic colitis Colon and rectum Y Unknown 1992 RCT 86 93 Unknown Unknown Cancer, diverticular disease, IBD Left colon and rectum Y Unknown	Young Tabusso et al ^[30]	2002	RCT	24	23	12:12	9:14	Unknown	Colon and rectum	Υ	Unknown	Unknown
1997 Observational 61 75 32.29 44:31 Cancer, diverticular disease, IBD, adenoma, lipoma Left colon and rectum Y Unknown 1997 Observational 61 75 32.29 44:31 Cancer, diverticular disease, IBD, adenoma, lipoma Left colon and rectum Y Unknown 1994 RCT 82 87 52:30 43:44 Cancer, diverticular disease, IBD, ischemic colitis Colon and rectum Y Unknown 1992 RCT 86 93 Unknown Unknown Cancer and other Colon and rectum Y Unknown	Miettinen <i>et al</i> ^[56]	2000	RCT	138	129	68:70	62:67	Cancer, IBD, diverticular disease	Colon and rectum	91% primary	None	None
1997Observational617532.2944:31Cancer, diverticular disease, IBD, adenoma, lipomaLeft colon and rectumYUnknown1995RCT30UnknownUnknownCancer, diverticular disease, IBD, ischemic colitisColon and rectumNUnknown1994RCT828752.3043:44Cancer, diverticular disease, IBDLeft colon and rectumYUnknown1992RCT8693UnknownUnknownCancer and otherColon and rectumYUnknown										both arms		
1995 RCT 30 30 Unknown Unknown Cancer, diverticular disease, IBD, ischemic colitis Colon and rectum N Unknown 1994 RCT 82 87 52:30 43:44 Cancer, diverticular disease, IBD Left colon and rectum Y Unknown 1992 RCT 86 93 Unknown Unknown Cancer and other Colon and rectum Y Unknown	Memon et al ^[73]	1997	Observational	61	75	32:29			Left colon and rectum	Y	Unknown	Unknown
1994 RCT 82 87 52:30 43:44 Cancer, diverticular disease, IBD Left colon and rectum Y Unknown 1992 RCT 86 93 Unknown Unknown Cancer and other Colon and rectum Y Unknown	Fillmann <i>et al</i> ^[60]	1995	RCT	30	30	Unknown	Unknown	Cancer, diverticular disease, IBD, ischemic colitis	Colon and rectum	z	Unknown	Unknown
1992 RCT 86 93 Unknown Unknown Cancer and other Colon and rectum Y Unknown	Burke <i>et al</i> ^[57]	1994	RCT	82	87	52:30	43:44	Cancer, diverticular disease, IBD	Left colon and rectum	Υ	Unknown	Unknown
	Brownson <i>et al</i> ^[58]	1992	RCT	86	93	Unknown	Unknown	Cancer and other	Colon and rectum	Y	Unknown	Unknown



	MBP	N	No MBP			Odds ratio	Odds ratio
Study or subgroup	Events		Events	Total		M-H, random, 95%CI	M-H, random, 95%CI
MBP vs rectal en		Total	LVCING	Total	meight		
Allaix 2015	24	706	30	829	8.7%	0.94 [0.54, 1.62]	1
Bertani 2011	9	114	9	115	3.6%	1.01 [0.39, 2.64]	-
Bucher 2005	5	78	1	75	0.8%	5.07 [0.58, 44.45]	-
Chan 2016	1	159	1	97	0.5%	0.61 [0.04, 9.83]	-
Ji 2017	42	538	77	831	12.8%	0.83 [0.56, 1.23]	-
Moral 2009	5	70	4	69	2.0%	1.25 [0.32, 4.87]	-
Otchy 2014	1	86	2	79	0.7%	0.45 [0.04, 5.09]	
Pitot 2009	2	59	6	127	1.4%	0.71 [0.14, 3.62]	
Platell 2006	3	147	7	147	1.9%	0.42 [0.11, 1.64]	-
Veenhof 2007	1	78	4	71	0.8%	0.22 [0.02, 1.99]	-
Yamada 2014	8			106	1.5%		-
	8 7	152 187	2 4			2.89 [0.60, 13.88]	-
Zmora 2003	/		4	193	2.3%	1.84 [0.53, 6.38]	-
Subtotal (95%CI)	100	2374	147	2739	36.9%	0.92 [0.70, 1.20]]
Total events	108	0 52 4	147	0.57	$1. \tau^2 = 00/$		
Heterogeneity: Tau ²				' = 0.57	; I = 0%)	
Test for overall effect	: <i>∠</i> = 0.65 (P = 0.52)				
MBP <i>vs</i> No ME	3P						
Ali 2007	6	109	1	101	0.8%	5.83 [0.69, 49.25]	• • • • • • •
Bhattacharjee 2015	4	38	2	33	1.2%	1.82 [0.31, 10.66]	· · · · · · · · · · · · · · · · · · ·
Bretagnol 2007	9	61	8	52	3.2%	0.95 [0.34, 2.68]	
Bretagnol 2010	6	89	14	89	3.4%	0.39 [0.14, 1.06]	
Brownson 1992	8	86	1	93	0.9%	9.44 [1.15, 77.10]	· · · · · · · · · · · · · · · · · · ·
Burke 1994	3	82	4	87	1.6%	0.79 [0.17, 3.63]	
Contant 2007	32	670	37	684	10.1%	0.88 [0.54, 1.43]	
Fa-Si-Oen 2005	7	125	6	125	2.8%	1.18 [0.38, 3.61]	
Fillmann 1995	0	30	1	30	0.4%	0.32 [0.01, 8.24]	·
Hu 2017	1	76	0	72	0.4%	2.88 [0.12, 71.87]	
Jung 2006	3	27	0	17	0.4%	5.00 [0.24, 103.07]	
Jung 2007	13	686	17	657	5.7%	0.73 [0.35, 1.51]	
Kiran 2015	184	6146	104	2296	18.5%	0.65 [0.51, 0.83]	
Leiro 2008	3	64	9	65	2.0%	0.31 [0.08, 1.19]	
Memon 1997	5	61	2	75	1.3%	3.26 [0.61, 17.42]	
Miettinen 2000	5	138	3	129	1.7%	1.58 [0.37, 6.74]	
Miron 2008	3	60	1	39	0.7%	2.00 [0.20, 19.95]	
Pena-Soria 2008	4	65	3	64	1.6%	1.33 [0.29, 6.21]	
Ram 2005	1	164	2	165	0.7%	0.50 [0.04, 5.57]	
Roig 2010	4	39	7	69	2.1%	1.01 [0.28, 3.70]	
Sasaki 2012	1	38	3	41	0.7%	0.34 [0.03, 3.44]	·
Tahirkheli 2013	8	48	6	48	2.7%	1.40 [0.45, 4.39]	
Young Tabusso 2002	5	24	0	23	0.4%	13.26 [0.69, 254.97]	
Subtotal (95%CI)	5	8926	Ū		63.1%	0.94 [0.70, 1.25]	
Subtotal (55 /001)		0520		5051	001170	0101 [01/07 1120]	
Total events	315		231				
Heterogeneity: Tau ² :		29.20 d		P = 0.14). $I^2 = 250$	0/6	
Test for overall effect				- 0.14), 1 – 25	70	
	. 2 – 0.45 (- 0.07	,				
Total (95%CI)		11300		7793	100.0%	0.90 [0.74, 1.10]	
	472	11500	270		100.0 /0	0.50 [0.74, 1.10]	
Total events	423	20.00	378		n). n ² - f	10/	
Heterogeneity: Tau ² =				$\nu = 0.23$	$S_{I}; I^{-} = 14$	1%	0.01 0.1 1 10 100
Test for overall effect				0.00	72 001		Favours MBP Favours No MBP
Test for subgroup diff	rerences: χ^2	= 0.01, 0	лг = 1 (<i>P</i>	= 0.90	$I^{-} = 0\%$		

Figure 2 Forest plot comparing overall anastomotic leak rate for patients receiving mechanical bowel preparation vs either a single rectal enema (top) or absolutely no preparation (bottom). A Mantel-Haenszel random effects model was used to perform the meta-analysis and odds ratios are quoted including 95% confidence intervals. MBP: Mechanical bowel preparation.

MBP *vs* no MBP did not impact upon the incidence of intra-abdominal collection (OR = 0.86, 95%CI: 0.63 to 1.17, *P* = 0.34), nor when full MBP *vs* single rectal enema^[37,45,47,51,53,66,67,69,72,74,77] (OR = 0.83, 95%CI: 0.45 to 1.51, *P* = 0.54) or MBP *vs* absolutely no preparation at all were considered^[29,40,46,48-50,52,54-61,64,65,68,70,73,75] (OR = 0.92, 95%CI: 0.62 to 1.34, *P* = 0.65).

When randomised controlled trials alone were

considered^[37,40,45, 46,48,49,51,53-56, 58,59,61,62,64,65] (Supplementary Figure 3A), no differences were seen in the incidence of intra-abdominal collection between any of the groups (OR = 1.17, 95%CI: 0.66 to 2.10, P = 0.59). However, when observational studies were analysed^[29,66-75,77] (Supplementary Figure 3B), the incidence of intra-abdominal collection was significantly reduced in those who had MBP *vs* those who did not (OR

Table 3 Nature of	f the bowel preparation used	in studies included in the meta-a	malysis
Ref.	Details of MBP	Details of no MBP	Antibiotics given
Allaix et al ^[74]	PEG	Enema before left sided operations	As per local policy
Kiran et al ^[29]	As per local policy	Unclear	As per local policy
Yamada <i>et al</i> ^[66]	PEG	Glycerin Enema	Flomoxef at induction and 3 hourly intra op
Otchy et al ^[67]	PEG	Colonic resections- no MBP	Ertapenem 1 g or levofloxacin/metronidazole 500 mg 1 h
5		Rectal resections- single enema	post op then continued for 24 h post op
Kim et al ^[75]	As per local policy	Unclear	As per local policy
Tahirkheli <i>et al</i> ^[62]	Saline	No preparation	Oral ciprofloxacin plus unspecified intravenous antibiotics for 24 h post op
Sasaki <i>et al</i> ^[61]	PEG and sodium picosulphate	No preparation	Antibiotic regime not specified
Bertani <i>et al</i> ^[45]	PEG and a single enema	Single enema only	Cefotixin given at induction, 4, 12 and 24 h. Ceftriaxone and metronidazole given for 5 d post op if heavy contamination
Roig et al ^[68]	Mono and di sodium phosphate	No prep	Antibiotic regime not specified
Bretagnol <i>et al</i> ^[46]	Senna plus povidone-iodine	No prep	ceftriaxone and metronidazole at induction and every 2
	enema		hours intra op
Pitot et al ^[69]	PEG	Rectal resections had single enema	Antibiotic regime not specified
Alcantara Moral <i>et</i> al ^[47]	Sodium phosphate or PEG	Two preoperative enemas	Neomycin and metronidazole 1 d pre op, ceftriaxone and metronidazole at induction
Miron <i>et al</i> ^[70]	PEG and sodium sulphate	No preparation	Antibiotic regime not specified
Pena-Soria <i>et al</i> ^[48]	PEG and standard enema	No preparation	Gentamicin and metronidazole 30 min pre op and 8 hourly post op
Leiro et al ^[59]	Sodium di or monobasic phosphate or PEG	No preparation	Ciprofloxacin and metronidazole 500 mg pre op
Contant et al ^[40]	PEG and bisocodyl/ sodium phosphate	No preparation	Antibiotic regime not specified
Bretagnol et al ^[71]	Senna plus povidone-iodine enema	No preparation	Ceftriaxone and metronidazole at induction and every 2 h intra op
Jung et al ^[49]	As per local policy	No preparation	Trimethoprim + metronidazole or cef and met or dozy and met
Veenhof et al ^[72]	PEG	Single enema	Antibiotic regime not specified
Ali et al ^[63]	Saline	No preparation	Antibiotic regime not specified
Jung et al ^[50]	PEG or sodium phosphate	No preparation	Oral sulphamethoxazole-trimethoprim and metronidazole, cephalsporin and metronidazole, doxycycline and
TT	77.0		metronidazole
Platell <i>et al</i> ^[51]	PEG	Phosphate enema	Timentin or gentamycin and metronidazole at induction
Fa-Si-Oen <i>et al</i> ^[52]	PEG	No preparation	Ceftriaxone and metronidazole or gentamycin and metronidazole at induction
Bucher et al ^[53]	PEG	Rectal resections had single saline enema	Ceftriaxone and metronidazole at induction and 24 h post op
Ram et al ^[54]	Monobasic and dibasic sodium phosphate	No preparation	Ceftriaxone and metronidazole 1 h pre op and 48 post op
Zmora et al ^[37]	PEG	Rectal resections had a single phosphate enema	Erythromycin and neomycin for 3 doses and then for 24 h
Young Tabusso et al ^[55]	PEG or saline/mannitol	No preparation	Antibiotic regime not specified
Miettinen <i>et al</i> ^[56]	PEG	No preparation	Ceftriaxone and metronidazole at induction
1001	Phosphate enema, picolax, PEG, saline lavage	No preparation	Antibiotic regime not specified
Fillmann <i>et al</i> ^[60]	Mannitol	No preparation	Metronidazole and gentamicin 1 h pre op then for 48 h
Burke <i>et al</i> ^[57]	sodium picosulphate	No preparation	Ceftriaxone 1 g, metronidazole at induction and 8 and 16 h

MBP: Mechanical bowel preparation; PEG: Polyethylene glycol.

= 0.67, 95%CI: 0.53 to 0.85, P = 0.0008). A significant reduction in the incidence of intra-abdominal collection was seen in the subgroup of patients who underwent MBP *vs* absolutely no preparation^[29,68,70,71,73,75] (OR = 0.65, 95%CI: 0.54 to 0.78, P < 0.0001), however no difference was seen in those undergoing MBP *vs* a single rectal enema^[66,67,69, 72,74,77] (OR = 0.80, 95%CI: 0.34 to 1.88, P = 0.60).

Hospital length of stay

Hospital length of stay (LOS) was reported in 20

studies^[40,45,46,49,51-56,61,63,67-69,71-74,77] including 7381 patients (Figure 5), with the use of MBP *vs* not (including those who received a single rectal enema) resulting in no significant difference in hospital length of stay (overall mean difference 0.11 d, 95%CI: -0.51 to 0.73, P = 0.72). This was mirrored when just randomised controlled trials were examined^[40,45,46,49,51-56,61,63] (Supplementary Figure 4A; overall mean difference 0.22 d, 95%CI: -0.44 to 0.88, P = 0.52) and when just observational studies were included^[67-69,71-74,77] (Supplementary Figure 4B; overall mean difference

	MBP		No M	RP		Odds ratio	Odds ratio
Study or subgroup	Events		Events	Total	Weight	M-H, random, 95%CI	M-H, random, 95%CI
MBP vs rectal en						,	
Allaix 2015	4	706	12	829	2.8%	0.39 [0.12, 1.21]	
Bertani 2011	7	114	14	115	3.7%	0.47 [0.18, 1.22]	
Bucher 2005	10	78	3	75	2.2%	2.53 [0.93, 13.37]	
Chan 2016	6	159	4	97	2.3%	0.91 [0.25, 3.32]	
Moral 2009	8	70	4	69	2.4%	2.10 [0.60, 7.32]	
Otchy 2014	5	86	5	79	2.3%	0.91 [0.25, 3.28]	
Pitot 2009	1	59	4	127	0.9%	0.53 [0.06, 4.85]	
Platell 2006	19	147	21	147	5.6%	0.89 [0.46, 1.74]	
Veenhof 2007	1	78	7	71	1.0%	0.12 [0.01, 0.99]	
Yamada 2014	5	152	2	106	1.5%	1.77 [0.34, 9.29]	
			2			13.17 [1.69, 102.30]	
Zmora 2003	12	187	1	193	1.0% 25.8%		
Subtotal (95%CI)	70	1836	77	1908	25.0%	1.00 [0.57, 1.76]	\bullet
Total events	78	20 72	77	D 0 07	οι. <i>τ</i> ² Γο	0/	
Heterogeneity: Tau ²				r = 0.02	2; 1 = 52	170	
Test for overall effect	:∠ = 0.00 (P = 1.00)				
MBP <i>vs</i> No MI			-	~~	2 00/		
Bhattacharjee 2015	11	38	6	33	2.8%	1.83 [0.59, 5.67]	
Bretagnol 2007	3	89	1	89	0.9%	3.07 [0.31, 30.09]	
Brownson 1992	5	86	7	93	2.6%	0.76 [0.23, 2.49]	
Burke 1994	4	82	3	87	1.8%	1.44 [0.31, 6.62]	
Contant 2007	90	670	96	684	9.3%	0.95 [0.70, 1.30]	- - -
Fa-si-oen 2005	9	125	7	125	3.3%	1.31 [0.47, 3.63]	
Fillmann 1995	1	30	2	30	0.8%	0.48 [0.04, 5.63]	
Hu 2017	9	76	2	72	1.7%	4.70 [0.98, 22.56]	
Jung 2006	4	27	1	17	0.9%	2.78 [0.28, 27.27]	
Jung 2007	54	686	42	657	8.0%	1.25 [0.82, 1.90]	
Kim 2014	52	1363	73	1112	8.6%	0.56 [0.39, 0.81]	_ _
kiran 2015	349	6146	190	2296	10.5%	0.67 [0.56, 0.80]	+
Leiro 2008	10	64	10	65	3.6%	1.02 [0.39, 2.64]	
Memon 1997	4	61	10	75	2.6%	0.46 [0.14, 1.53]	
Miettinen 2000	5	138	3	129	1.9%	1.58 [0.37, 6.74]	
Miron 2008	9	60	7	39	3.0%	0.81 [0.27, 2.38]	
Pena-Soria 2008	16	65	11	64	4.2%	1.57 [0.67, 3.72]	
Ram 2005	16	164	10	165	4.4%	1.68 [0.74, 3.81]	
Roig 2010	5	39	13	69	2.9%	0.63 [0.21, 1.93]	
Sasaki 2012	0	38	0	41		Not estimable	
Young Tabusso 2002		24	0	23	0.5%	5.22 [0.24, 114.87]	
Subtotal (95%CI)		10071		5965	74.2%	0.98 [0.78, 1.24]	▲
Total events	658		494				Ţ
Heterogeneity: Tau ²		= 32,88. d		P = 0.02): $I^2 = 42^{\circ}$	%	
Test for overall effect				0.02	// IZ		
	_ 0.17	,	,				
Total (95%CI)		11907		7873	100.0%	0.99 [0.80, 1.24]	•
Heterogeneity: Tau ²	$= 0.11: \gamma^2 =$		df = 30.6				Ţ
Test for overall effect				0.00	- // -		
Test for subgroup dif				r = 0.95	$J^2 = 0\%$		
		0.00,		5.55	, 70		
							0.01 0.1 1 10
							Favours MBP Favours No MBP
							FAVOUIS MIDE FAVOUIS NO MIDE

Figure 3 Forest plot comparing overall surgical site infection rates for patients receiving mechanical bowel preparation vs either a single rectal enema (top) or absolutely no preparation (bottom). A Mantel-Haenszel random effects model was used to perform the meta-analysis and odds ratios are quoted including 95% confidence intervals. MBP: Mechanical bowel preparation.

-0.12 d, 95%CI: -1.48 to 1.25, *P* = 0.87).

Mortality S

Mortality was reported in 25 studies^{[29,37,40,45-49,51-54,56,57,59,60,}

 $^{65,66,68,69,71-74,77]}$ that included 16657 patients (Figure 6). The time point this outcome measure was measured was variable between studies, with the majority taken at 30 d $^{[29,37,45-49,51,53,60,65,69,71,73,77]}$, two taken at first outpatient clinic quoted to be approximately two weeks following hospital discharge $^{[40]}$ or four weeks following

surgery^[66], one at two months^[56] and one at three months^[52], with six papers not stating when mortality was taken from^[54,57,59,68,72,74]. No difference was seen with the use of full MBP, single rectal enema or no preparation at all.

A similar result was seen, with no significant differences, when this comparison was made using only randomised controlled trials^[37,40,45-49,51-54,56,57,59,60,65] (Supplementary Figure 5A). However, in observational studies^[29,66,68,69,71-74,77], MBP was associated with a

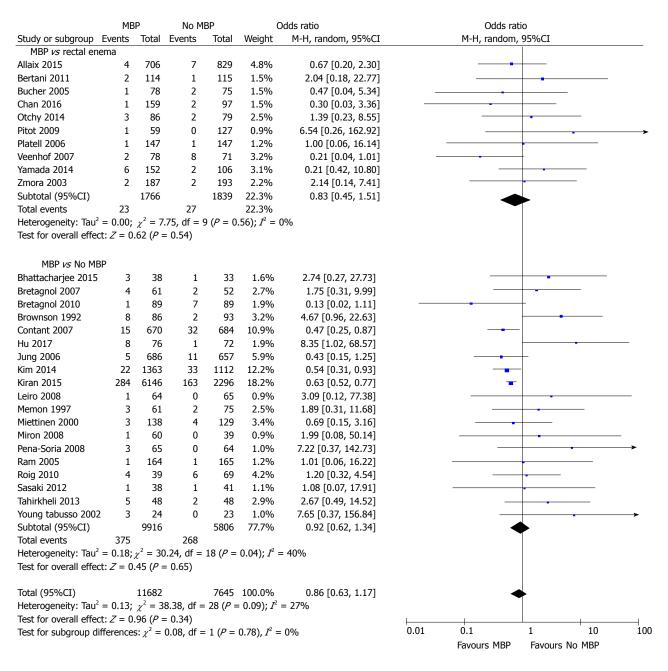


Figure 4 Forest plot comparing overall intra-abdominal collection rates for patients receiving mechanical bowel preparation vs either a single rectal enema (top) or absolutely no preparation (bottom). A Mantel-Haenszel random effects model was used to perform the meta-analysis and odds ratios are quoted including 95% confidence intervals. MBP: Mechanical bowel preparation.

significant reduction in mortality (OR = 0.50, 95%CI: 0.34 to 0.74, *P* = 0.0005) (Supplementary Figure 5B). A significant reduction in the incidence of intraabdominal collection was seen in the subgroup of patients in observational studies who underwent MBP *vs* absolutely no preparation^[29,68,71,73] (OR = 0.42, 95%CI: 0.27 to 0.56, *P* < 0.0001). However, no difference was seen in these undergoing MBP *vs* a single rectal enema^[66,69,72,74,77] (OR = 0.99, 95%CI: 0.41 to 2.41, *P* = 0.98).

Reoperation

A total of 20 studies on 16742 patients $^{[29,40,46,49,51-57,59,65,68,69,71,72,74,76,77]}$ examined the impact of MBP upon

reoperation rates (Figure 7). Overall the use of MBP *vs* no MBP did not impact upon requirement for reoperation^[29,40,46,49,51-57, 59,65,68,69,71,72,74,76,77] (OR = 0.91, 95%CI: 0.75 to 1.12, P = 0.38), nor when MBP *vs* a single rectal enema^[51,53,69,72,74,76,77] (OR = 0.82, 95%CI: 0.42 to 1.60, P = 0.56) or MBP *vs* absolutely no preparation^[29,40,46,49,52,54-57,59,65,68,71] (OR = 0.85, 95%CI: 0.72 to 1.01, P = 0.06) were compared.

When only randomised controlled trials were examined^[40,46,49,51-57,59,65] (Supplementary Figure 6A), again no difference was seen by the use of MBP, a single rectal enema or absolutely no preparation. When observational studies were examined^[29,68,69,71,72,74,76,77] (Supplementary Figure 6B) overall MPB resulted in no

Table 4 Effect of bowel preparation on outcome in patients undergoing rectal surgery										
	Number of participants (MBP vs No MBP)	Odds ratio (95%CI), MBP vs No MBP	P value							
Anastomotic leak	2351 (1042 vs 1309)	0.86 (0.64 to 1.15)	0.30							
Surgical site infection	965 (513 vs 452)	1.22 (0.82 to 1.81)	0.33							
Intra-abdominal collection	921 (486 vs 435)	0.54 (0.21 to 1.38)	0.20							
Mortality	813 (419 vs 394)	0.73 (0.29 to 1.82)	0.50							
Re-operation	1660 (688 vs 392)	1.57 (1.02 to 2.43)	0.04							

Data from^[39,45,46,50,56,57,59,71,75-77]. MBP: Mechanical bowel preparation.

			MBP	No M	BP			Mean difference	Mean difference
Study or subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV,random, 95%CI	IV, random, 95%CI
MBP vs rectal en	ema								
Allaix 2015	8.64	6.24	706	8.01	6.08	829	7.8%	0.63 [0.01, 1.25]	
Bertani 2011	7	3.2	114	6	3.3	115	7.3%	1.00 [0.16, 1.84]	_ _
Bucher 2005	14.9	13.1	78	9.9	3.8	75	2.8%	5.00 [1.97, 8.03]	
Chan 2016	5.6	7.1	159	6.8	7.3	97	4.9%	-1.20 [-3.02, 0.62]	
Otchy 2014	6	4.3	86	5	4.3	79	6.1%	1.00 [-0.31, 2.31]	
Pitot 2009	8	16	59	5	19.8	127	1.2%	3.00 [-2.34, 8.34]	
Platell 2006	9	3.1	147	9.4	4.3	147	7.2%	-0.40 [-1.26, 0.46]	
Veenhof 2007	8	4.4	78	8	4.4	71	5.9%	0.00 [-1.41, 1.41]	
Subtotal (95%CI)			1427			1540	43.1%	0.56 [-0.19, 1.30]	
Heterogeneity: Tau ²	= 0.62;	$\chi^2 = 1$	9.44, df	^f = 7 (<i>P</i>	= 0.00)7); <i>I</i> ² =	= 64%		
Test for overall effect	t: Z = 1	45 (<i>P</i> :	= 0.15)						
MBP <i>vs</i> No MB	P								
Ali 2007	10	3.8	109	15	13.1	101	3.3%	-5.00 [-7.65, -2.35]	
Bretagnol 2007	12	11.25	61	10	8.5	52	2.2%	2.00 [-1.65, 5.65]	
Bretagnol 2010	14	9	89	16	12	89	2.7%	-2.00 [-5.12, 1.12]	
Contant 2007	10	4.4	670	10	3.7	684	8.1%	0.00 [-0.43, 0.43]	
Fa-Si-Oen 2005	10	36.5	125	9	8.5	125	0.8%	1.00 [-5.57, 7.57]	
Jung 2006	8.6	7	686	8.8	6.9	657	7.5%	-0.20 [-0.94, 0.54]	
Memon 1997	13.25	0.8	61	16.17	1.75	75	8.1%	-1.92 [-2.36, -1.48]	-
Miettinen 2000	4.8	1.6	138	5	2	129	8.1%	-0.20 [-0.64, 0.24]	
Ram 2005	8.2	5.1	164	8	2.7	165	7.2%	0.20 [-0.68, 1.08]	
Roig 2010	9.1	6.2	39	9.2	8.7	69	3.1%	-0.10 [-2.93, 2.73]	
Sasaki 2012	19.9	24.25	38	15.5	18.25	41	0.4%	4.40 [-5.12, 13.92]	`
Young tabusso 2002	14	3	24	11	2.3	23	5.6%	3.00 [1.48, 4.52]	
Subtotal (95%CI)			2204			2210	56.9%	-0.28 [-1.12, 0.56]	
Heterogeneity: Tau ²	= 1.26;	$\chi^2 = 84$	1.50, df	= 11 (P	< 0.0	0001);	$I^2 = 87\%$)	
Test for overall effec				``		,,			
			,						
Total (95%CI)			3631			3750	100.0%	0.11 [-0.51, 0.73]	•
Heterogeneity: Tau ²	= 1.20:	$\gamma^2 = 1$	24.12. 0	df = 19	(<i>P</i> < 0				
Test for overall effec							,,	-	-4 -2 0 2 4
Test for subgroup di		•	,	f = 1 (P)	= 0.1	5), $I^2 =$	52.5%		Favours MBP Favours No MBP
		<i>r</i>	, u		0.1	-,,-	/ / /		

Figure 5 Forest plot comparing overall hospital length of stay for patients receiving mechanical bowel preparation vs either a single rectal enema (top) or absolutely no preparation (bottom). An inverse-variance random effects model was used to perform the meta-analysis and mean differences are quoted including 95% confidence intervals. MBP: Mechanical bowel preparation.

significant reduction in the reoperation rate vs those who did not have bowel preparation but may have had a rectal enema (OR = 0.86, 95%CI: 0.64 to 1.15, P = 0.30), as well as when those who has a single rectal enema (OR = 0.82, 95%CI: 0.44 to 1.52, P = 0.52), however a significant difference was seen when MBP was compared with patients who received absolutely no preparation (OR = 0.78, 95%CI: 0.63 to 0.97, P = 0.02).

Rectal surgery

A total of 11 studies^[39,45,46,50,56,57,59,71,75-77] included either only patients who were undergoing rectal or surgery, or outcome measures for the subgroup of patients who had undergone rectal surgery. Ten studies compared MBP with no MBP, with just one study comparing MBP with a single rectal enema^[45]. All studies except one^[77] included data on anastomotic leak rates, finding MBP not to be associated with any difference in incidence (OR = 0.86, 95%CI: 0.64 to 1.15, P = 0.30). Only seven studies^[39,45,46,50,71,75,77] included data on SSI, which also demonstrated no significant difference (OR = 1.22, 95%CI: 0.82 to 1.81, P = 0.33). Intraabdominal collection and mortality data were similarly only available for five^[39,45,46,71,77] and four studies^[39,45,46,71] respectively, neither of which were associated with the use of MBP (OR = 0.54, 95%CI: 0.21 to 1.38, P =

	M	BP	No M	IBP	0	dds ratio		Odds ratio
Study or subgroup	Events	Total	Events	Total	Weight M	-H, random, 95%CI		M-H, random, 95%CI
MBP vs rectal en	ema							
Allaix 2015	5	706	6	829	8.0%	0.98 [0.30, 3.22	2]	
Bertani 2011	0	114	0	115		Not estimat	e	
Bucher 2005	0	78	0	75		Not estimate	e	
Chan 2016	1	159	0	97	1.5%	1.85 [0.07, 45.75	5]	
Moral 2009	2	70	0	69	1.6%	5.07 [0.24, 107.62	2]	
Pitot 2009	1	59	1	127	1.9%	2.17 [0.13, 35.34	ŀĴ	
Platell 2006	4	147	1	147	2.9%	4.08 [0.45, 36.98	5]	
Veenhof 2007	2	78	3	71	4.1%	0.60 [0.10, 3.68	3]	
Yamada 2014	0	152	0	106		Not estimat	-	
Zmora 2003	3	187	3	193	5.0%	1.03 [0.21, 5.18	8]	
Subtotal (95%CI)		1750		1839	25.0%	1.27 [0.62, 2.61	.]	
Total events	18		14				-	
Heterogeneity: Tau ²	$= 0.00; \chi^2 =$	= 3.00, d	lf = 6 (<i>P</i>	= 0.81);	$I^2 = 0\%$			
Test for overall effect								
MBP <i>vs</i> rectal N	1BP							
Bhattacharjee 2015	1	38	1	33	1.9%	0.86 [0.05, 14.39]		
Bretagnol 2007	0	61	1	52	1.4%	0.28 [0.01, 7.00]		
Bretagnol 2010	1	89	0	89	1.4%	3.03 [0.12, 75.48]		
Burke 1994	2	82	0	87	1.6%	5.43 [0.26, 114.92]		_
Contant 2007	20	670	26	684	17.7%	0.78 [0.43, 1.41]		
Fa-Si-Oen 2005	2	125	1	125	2.5%	2.02 [0.18, 22.52]		
Fillman 1995	0	30	0	30		Not estimate		
Jung 2006	6	686	6	657	8.6%	0.96 [0.31, 2.98]		
Kiran 2015	31	6146	37	2296	20.5%	0.31 [0.19, 0.50]		
Leiro 2008	1	64	2	65	2.4%	0.50 [0.04, 5.66]		
Memon 1997	2	61	0	75	1.6%	6.34 [0.30, 134.68]		
Miettinen 2000	0	138	0	129		Not estimate		
Pena-Soria 2008	3	65	4	64	5.4%	0.73 [0.16, 3.38]		_
Ram 2005	2	164	2	165	3.6%	1.01 [0.14, 7.23]		
Roig 2010	4	39	5	69	6.5%	1.46 [0.37, 5.80]		
Subtotal (95%CI)		8458		4620	75.0%	0.77 [0.47, 1.25]		
Total events	75		85					-
Heterogeneity: Tau ²	= 0.19; χ ² =	= 17.30, d	df = 12 (A	P = 0.14); <i>I</i> ² = 31%			
Test for overall effect	:: <i>Z</i> = 1.06 ((P = 0.29)))					
Total (95%CI)		10208		6449	100.0%	0.85 [0.57, 1.27]		◆
Total events	93		99					
Heterogeneity: Tau ²	= 0.14; χ^2 =	= 24.15,	df = 19 (P = 0.19	9); I ² = 21%	, 0		
Test for overall effect								
Test for subgroup dif	ferences: χ^2	² = 1.29,	df = 1 (A	? = 0.26)), $I^2 = 22.79$	ю		
							0.01	0.1 1 10 100
								Favours MBP Favours No MBP

Figure 6 Forest plot comparing overall mortality rates for patients receiving mechanical bowel preparation vs either a single rectal enema (top) or absolutely no preparation (bottom). A Mantel-Haenszel random effects model was used to perform the meta-analysis and odds ratios are quoted including 95% confidence intervals. MBP: Mechanical bowel preparation.

0.20; and OR = 0.73, 95%CI: 0.29 to 1.82, P = 0.50, respectively). The results in patients undergoing rectal surgery are summarized in Table 4.

DISCUSSION

This meta-analysis of 23 randomised controlled trials and 13 observational studies has demonstrated that, overall, the use of MBP *vs* either absolutely no bowel preparation or a single rectal enema was not associated with a statistically significant difference in the incidence of anastomotic leak, SSI, intra-abdominal collection, mortality, reoperation or total hospital length of stay. When just randomised controlled trial evidence was analysed, there was, again, no significant difference by preparation method in any clinical outcome measure. Finally, when observational studies were analysed, the use of full preparation was associated overall with a reduced incidence of anastomotic leak, SSI, intra-abdominal collection and mortality rates, with these results mirrored in patients receiving MBP *vs* absolutely no preparation, but no significant differences in those receiving MBP *vs* a single rectal enema. When a separate subgroup of just rectal surgery was considered, MBP was not associated with a statistically significant difference in anastomotic leak rates, SSI, intra-abdominal collection or mortality, irrespective of whether patients not receiving MBP were given a single

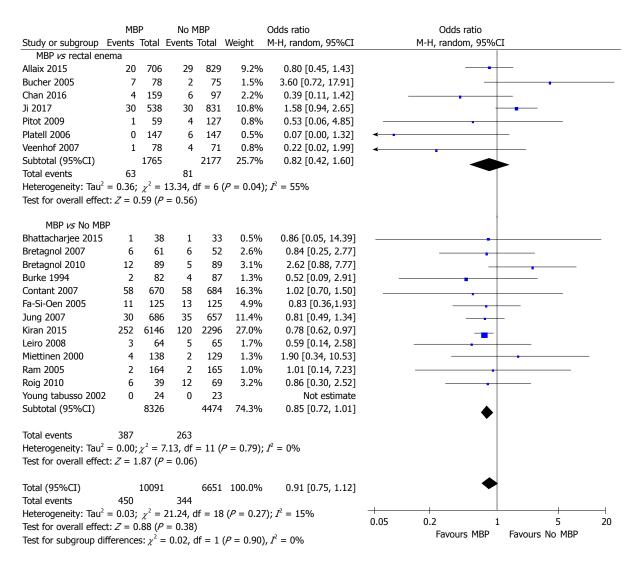


Figure 7 Forest plot comparing overall reoperation rates for patients receiving mechanical bowel preparation vs either a single rectal enema (top) or absolutely no preparation (bottom). A Mantel-Haenszel random effects model was used to perform the meta-analysis and odds ratios are quoted including 95% confidence intervals. MBP: Mechanical bowel preparation.

rectal enema.

Strengths of study

This study represents the most comprehensive examination of the role of MBP prior to elective colorectal surgery to date. As part of the study plan, the decision was made to include observational studies as well as randomised controlled trials. However, in order to ensure that inclusion of studies of less rigorous methodology did not exert an undue bias, a predetermined analysis of studies of both methodologies was conducted. This revealed that the overall results and those from analysing just evidence from randomised controlled trials were much the same. However, when analysing evidence from observational studies, this resulted in a significant reduction in anastomotic leak, SSI, intraabdominal collection and mortality rates. The reasons for this difference in results is not clear from this study, but it is possible that selection bias may exert a confounding effect upon the results, and as such the use of MBP in selected patients as determined by the physician in

charge may be appropriate.

With the exception of hospital length of stay ($I^2 = 85\%$), overall study heterogeneity was low to moderate (0%-34%) for all clinical outcome measures, suggesting the studies to be relatively homogeneous. The risk of bias for the randomised controlled trials included in the meta-analysis (Table 1) was relatively low.

Limitations of study

As the raw mean and standard deviation data were not available on the hospital LOS for all studies, despite several attempts at obtaining this directly from the authors, it was necessary to infer this from what was available (either median and range or interquartile range) using statistical techniques previously described^[42]. This is a valid technique which has been well described previously, but this may exert some degree of bias upon the results of the meta-analysis.

There was poor documentation within the studies included regarding the side effects of MBP including the incidence of electrolyte disturbance, fluid depletion



and requirement of resuscitation, and renal disturbance or failure, hence this was not included as an outcome within the meta-analysis.

Emerging evidence, much of which has been derived from the studies based upon NSQIP datasets have focused upon the combination between intraluminal antibiotics and MBP and have demonstrated a reduction in SSI rates. However, the data contained within the studies included within this meta-analysis has been scanty regarding the use of intraluminal antibiotics and as such it has not been possible to include this data within the meta-analysis. This may act as a potential confounder when considering the effect of MBP and clinical outcomes.

The studies contained predominantly mixed populations of colonic and rectal procedures, with inadequate documentation to differentiate results between the two, which may be particularly important in addressing the question regarding the use of a single rectal enema as bowel preparation. In addition, there was poor documentation regarding the nature of the anastomoses within the studies included, with a mixture of ileocolic, colon-colon and colorectal. The role of mechanical bowel preparation in various anastomosis types has not been well established. The majority of studies included a predominance of colonic procedures, with some focusing entirely on colonic rather than rectal surgery. Only a small subgroup analysis was available to analyse the impact of MBP in rectal surgery, from which it is very difficult to draw strong conclusions. Further studies are required to discern the importance of a preoperative enema in this setting. Similarly, the level of documentation in studies regarding laparoscopic vs open surgery was not sufficient in terms of correlation with clinical outcome measures to be able to discern the importance of MBP in this setting. Only one recent observational study has focused entirely on laparoscopic procedures^[74] which demonstrated no significant difference in the rates of intra-abdominal septic complications by the use of MBP, and prior to this evidence was purely based on several small studies^[38,78].

The nature of the MBP used was inconsistent between studies, and this may introduce a further bias^[79]. There was also poor documentation regarding antibiotic usage, particularly in the early studies. Much of the recent literature regarding preparation of the bowel has focused upon the use of oral luminal antibiotics in combination with MBP, with these studies suggesting a potential role for this therapy^[26,27]. A recent metaanalysis on this topic has demonstrated a significant reduction in the risk of SSI in patients undergoing elective colorectal surgery given oral systemic antibiotics with MBP *vs* systemic antibiotics and MBP^[80], thus representing a further weakness in the studies included in this meta-analysis.

Comparison with other studies

A recently published meta-analysis^[8] of 18 randomised controlled trials, 7 non-randomised comparative studies,

and 6 single-group cohorts compared the use of oral MBP with or without an enema vs no oral MBP with or without an enema. This study found that MBP vs no MBP was associated with no difference in the rates of all-cause mortality (OR = 1.17, 95%CI: 0.67 to 2.67), anastomotic leakage (OR = 1.08, 95%CI: 0.79 to 1.63), SSI (OR = 1.19, 95%CI: 0.56 to 2.63) as well as wound infections, peritonitis or intra-abdominal abscess or reoperation. This study however found considerable variance in the estimation of treatment effects, possibly due to the large range of study methodology included, which may mask a treatment effect seen.

This topic has been reviewed by the Cochrane Collaboration^[81-83], with the most recent review conducted in 2011^[10]. This included a total of 18 randomised controlled trials in elective colorectal surgery (5805 patients), and demonstrated no statistically significant evidence to support the use of MBP in either low anterior resection, rectal or colonic surgery in terms of anastomotic leakage or wound infection.

A previous meta-analysis has examined the role of MBP prior to proctectomy^[12] from eleven publications (1258 patients), although extractable data were only available in a limited number of studies for outcome measures other than anastomotic leakage rates. This study^[12] found no beneficial effect from MBP prior to proctectomy with regards to anastomotic leakage (OR = 1.144, 95%CI: 0.767 to 1.708, *P* = 0.509), SSI (OR = 0.946, 95%CI: 0.597 to 1.498, *P* = 0.812), intraabdominal collection (OR = 1.720, 95%CI: 0.527 to 5.615, *P* = 0.369) or postoperative mortality.

Health policy implications

Worldwide, elective colorectal surgery is performed frequently. Current opinion regarding the use of MBP prior to this surgery is inconsistent^[17,18], despite several previous meta-analyses which have suggested this is not useful in reducing postoperative complications^[9,10]. The use of MBP is not without cost implications, including the preparation itself and in elderly and frail patients, MBP may also necessitate an additional stay in hospital prior to surgery due to the risk of dehydration and electrolyte disturbance which is associated with considerable additional healthcare costs. This metaanalysis further reinforces that MBP is not associated with any difference in postoperative complication rates, mortality of hospital length of stay, particularly in elective colonic surgery, and as such should not be administered routinely.

In conclusion, this study represents the most comprehensive meta-analysis to date on MBP in elective colorectal surgery. It has demonstrated that MBP *vs* a single rectal enema or no bowel preparation at all is not associated with a statistically significant difference in any of the clinical outcome measures studied. Given the risks of electrolyte disturbance and patient dissatisfaction, as well as potentially significant levels of dehydration and requirement for pre-admission prior to surgery, MBP should no longer be considered a standard of care prior



to elective colorectal surgery.

ARTICLE HIGHLIGHTS

Research background

Mechanical bowel preparation for colorectal surgery has been surgical dogma for decades, despite increasing evidence from the 1990s refuting its benefits. The rationale behind the administration of mechanical bowel preparation is that it reduces fecal bulk and, therefore, bacterial colonisation, thereby reducing the risk of postoperative complications such as anastomotic leakage and wound infection, as well as facilitate dissection and allow endoscopic evaluation. Opponents argue that in the 21st century, with rational use of oral and intravenous prophylactic antibiotics there is no longer a place for mechanical bowel preparation, that it may cause marked fluid and electrolyte imbalance in the preoperative period. As a result of this inconclusive evidence, practice varies between countries and even surgeons in the same institution. We conducted a comprehensive meta-analysis encompassing both randomised controlled trials and observational studies. We sought to address deficiencies in previous studies by including all levels of evidence, separating those in which patients received a single rectal enema vs full or no preparation.

Research motivation

The main topics focused on by this meta-analysis are the role of mechanical bowel preparation *vs* no preparation or rectal enema alone on postoperative infective complications in patients undergoing elective colorectal surgery, as well as in patients undergoing purely rectal resection. This meta-analysis also sought to examine evidence from both randomized controlled trials and observational studies and compare the results of meta-analyses conducted from these evidence sources.

Research objectives

The aims for this meta-analysis were to analyse the effect of mechanical bowel preparation *vs* no preparation or rectal enema alone on postoperative infective complications in patients undergoing elective colorectal surgery, to examine the differences in results between evidence obtained from randomised controlled trials and observational studies, and to determine what effect, if any, bowel preparation had on postoperative complications in rectal surgery. These aims were all achieved by this meta-analysis.

Research methods

We performed an electronic search of the PubMed database and the Cochrane Central Register of Controlled Trials to identify studies comparing outcomes in patients undergoing elective colorectal surgery treated with mechanical bowel preparation vs either no preparation or a single rectal enema. We performed this meta-analysis according to the Preferred Reporting Items for Systematic Reviews and Meta-analyses (PRISMA) statement. We reviewed full text articles for suitability after excluding studies on the basis of title and abstract. Our inclusion criteria specified that studies must have a minimum of two comparator groups and were either designed as randomised controlled trials or observational studies. Relevant outcome measures were anastomotic leak, surgical site infection, intra-abdominal abscess, mortality, reoperation and hospital length of stay. The analysis was performed using RevMan 5.3 software. Continuous variables were calculated as a mean difference and 95% confidence interval using an inverse variance random effects model. Dichotomous variables were analysed using the Mantel-Haenszel random effects model to quote the risk ratio (RR) and 95% confidence interval. These analyses were used to construct forest plots, with statistical significance taken to be a P value of < 0.05 on two tailed testing. A predetermined subgroup analysis was performed for the impact of MBP in rectal surgery specifically using the same methodology.

Research results

This meta-analysis of 23 randomised controlled trials and 13 observational studies has demonstrated that, overall, the use of MBP vs either absolutely no bowel preparation or a single rectal enema was not associated with a statistically significant difference in the incidence of anastomotic leak, surgical site infection, intra-abdominal collection, mortality, reoperation or total hospital

length of stay. When just randomised controlled trial evidence was analysed, there was again no significant difference by preparation method in any clinical outcome measure. Finally, when observational studies were analysed, the use of full preparation was associated overall with a reduced incidence of anastomotic leak, surgical site infection, intra-abdominal collection and mortality rates, with these results mirrored in patients receiving MBP vs absolutely no preparation, but no significant differences in those receiving MBP vs a single rectal enema.

Research conclusions

This study represents the most comprehensive examination of the role of mechanical bowel preparation prior to elective colorectal surgery to date and has demonstrated that, overall, the use of MBP vs either absolutely no bowel preparation or a single rectal enema was not associated with a statistically significant difference in the incidence of anastomotic leak, surgical site infection, intra-abdominal collection, mortality, reoperation or total hospital length of stay. Given the risks of electrolyte disturbance and patient dissatisfaction as well as potentially significant levels of dehydration and requirement for pre-admission prior to surgery, mechanical bowel preparation should no longer be considered a standard of care prior to elective colorectal surgery.

Research perspectives

This study represents the most comprehensive meta-analysis to date on mechanical bowel preparation in elective colorectal surgery. It has demonstrated that mechanical bowel preparation vs a single rectal enema or no bowel preparation at all is associated with no difference in any of the clinical outcome measures studied. Mechanical bowel preparation should no longer be considered a standard of care prior to elective colorectal surgery. Emerging evidence, much of which has been derived from the studies based upon NSQIP datasets, has focused upon the combination between intraluminal antibiotics and mechanical bowel preparation and has demonstrated a reduction in SSI rates. However, the data contained within the studies included within this metaanalysis have been scanty regarding the use of intraluminal antibiotics and as such it has not been possible to include these data within the meta-analysis. Further work on this topic should focus upon the role of intraluminal antibiotics in the setting of elective colorectal surgery.

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