

Published in final edited form as:

Ann Surg. 2019 July ; 270(1): 43–58. doi:10.1097/SLA.0000000000003145.

The Role of Oral Antibiotic Preparation in Elective Colorectal Surgery: A Meta-analysis

Katie E. Rollins, MRCS^{*}, Hannah Javanmard-Emamghissi, MRCS^{*}, Austin G. Acheson, DM, FRCS^{*}, and Dileep N. Lobo, DM, FRCS, FACS, FRCPE^{*,†}

^{*}Gastrointestinal Surgery, Nottingham Digestive Diseases Centre and National Institute for Health Research (NIHR) Nottingham Biomedical Research Centre, Nottingham University Hospitals NHS Trust and University of Nottingham, Queen's Medical Centre, Nottingham, UK

[†]MRC/ARUK Centre for Musculoskeletal Ageing Research, School of Life Sciences, University of Nottingham, Queen's Medical Centre, Nottingham, UK

Abstract

Objectives—To compare the impact of the use of oral antibiotics (OAB) with or without mechanical bowel preparation (MBP) on outcome in elective colorectal surgery.

Summary Background Data—Meta-analyses have demonstrated that MBP does not impact upon postoperative morbidity or mortality, and as such it should not be prescribed routinely. However, recent evidence from large retrospective cohort and database studies has suggested that there may be a role for combined OAB and MBP, or OAB alone in the prevention of surgical site infection (SSI).

Methods—A meta-analysis of randomized controlled trials and cohort studies including adult patients undergoing elective colorectal surgery, receiving OAB with or without MBP was performed. The outcome measures examined were SSI, anastomotic leak, 30-day mortality, overall morbidity, development of ileus, reoperation and *Clostridium difficile* infection.

Results—A total of 40 studies with 69,517 patients (28 randomized controlled trials, n = 6437 and 12 cohort studies, n = 63,080) were included. The combination of MBP+OAB versus MBP alone was associated with a significant reduction in SSI [risk ratio (RR) 0.51, 95% confidence interval (CI) 0.46–0.56, $P < 0.00001$, $I^2 = 13\%$], anastomotic leak (RR 0.62, 95% CI 0.55–0.70, $P < 0.00001$, $I^2 = 0\%$), 30-day mortality (RR 0.58, 95% CI 0.44–0.76, $P < 0.0001$, $I^2 = 0\%$), overall morbidity (RR 0.67, 95% CI 0.63–0.71, $P < 0.00001$, $I^2 = 0\%$), and development of ileus (RR 0.72, 95% CI 0.52–0.98, $P = 0.04$, $I^2 = 36\%$), with no difference in *Clostridium difficile* infection

This is an open access article distributed under the Creative Commons Attribution License 4.0 (CCBY), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited. (<http://creativecommons.org/licenses/by/4.0/>)

Reprints: Dileep N. Lobo, DM, FRCS, FACS, FRCPE, Gastrointestinal Surgery, Nottingham Digestive Diseases Centre, Nottingham University Hospitals NHS Trust and University of Nottingham, E Floor, West Block, Queen's Medical Centre, Nottingham NG7 2UH, UK. Dileep.Lobo@nottingham.ac.uk.

DNL has received unrestricted research funding for BBraun and speakers' honoraria from BBraun, Fresenius Kabi, Baxter Healthcare, and Shire for unrelated work. None of the other authors reports a conflict of interest.

This paper has been accepted for presentation to the Society for Academic and Research Surgery, London, January 2019 and a conference abstract will be published in the *British Journal of Surgery*.

rates. When a combination of MBP+OAB was compared with OAB alone, no significant difference was seen in SSI or anastomotic leak rates, but there was a significant reduction in 30-day mortality, and incidence of postoperative ileus with the combination. There is minimal literature available on the comparison between combined MBP+OAB versus no preparation, OAB alone versus no preparation, and OAB versus MBP.

Conclusions—Current evidence suggests a potentially significant role for OAB preparation, either in combination with MBP or alone, in the prevention of postoperative complications in elective colorectal surgery. Further high-quality evidence is required to differentiate between the benefits of combined MBP+OAB or OAB alone.

Keywords

anastomotic leak; colorectal; mechanical bowel preparation; oral antibiotics; surgery; surgical site infection

Surgical site infection (SSI) is a major burden for patients undergoing elective colorectal surgery. It adds significantly to the cost of health care, and administration of preoperative bowel preparation has been proposed to reduce the incidence of SSI. The role of mechanical bowel preparation (MBP) with polyethylene glycol or sodium phosphate has been studied in randomized controlled trials (RCTs), with perceived benefits including ease of manipulation of the bowel, reduced spillage and resultant contamination, reduced luminal pressure, and lesser bacterial load. However, a recent metaanalysis¹ of 36 RCTs and cohort studies, and an earlier one² of 14 RCTs found that that the administration of MBP did not impact upon postoperative morbidity or mortality. This, in combination with high rates of patient dissatisfaction and fluid and electrolyte disturbances, has led to the conclusion that MBP should not be prescribed routinely. This is reflected in Guidelines from the Enhanced Recovery After Surgery Society,^{3,4} the National Institute of Health and Care Excellence,⁵ and the American Society for Enhanced Recovery,⁶ all of which suggest that MBP should not be administered routinely. However, although the American Society for Enhanced Recovery guidelines suggest that MBP should not be given in isolation, they recommend routine use of an isosmotic bowel preparation and combined oral antibiotic prior to elective colorectal surgery.⁶

The use of oral antibiotic (OAB) prophylaxis, in the form of nonabsorbable luminal antibiotics, was first proposed in 1971 by Rosenberg et al⁷ in a RCT of 150 patients undergoing large bowel surgery receiving MBP alone, or MBP in combination with phthalylsulphathiazole or phthalylsulphathiazole and neomycin. The combination of MBP +OAB was associated with a significant reduction in SSI (23% vs. 40%), anastomotic leak rates (24% vs. 52%), and sepsis rates (37.3% vs. 64.4%).⁶ Although several studies provided evidence for the role of oral antibiotics in elective colorectal surgery, the regimens included large volume preparations,^{8–10} prolonged preoperative hospital admission, and in the setting of prolonged preoperative starvation protocols, dehydration, and electrolyte disturbances were commonplace.^{11,12} Decreased compliance and inconsistent bowel cleansing resulted in a reduced intervention effect and, this, combined with reduced preoperative admission times, resulted in the practice of combined MBP+OAB dwindling in favor of more restrictive MBP regimens alone. However, recently there has been resurgent

interest in the use of OAB in colorectal surgery,^{13,14} particularly in light of a large number of retrospective cohort and database studies, many of which originated from the American College of Surgeons National Surgical Quality Improvement Program (ACS NSQIP) targeted colectomy database.^{15–20} Evidence for the role of OAB has been summarized in several narrative reviews^{21,22} as well as meta-analyses,^{23–25} which have supported a reduction in SSI associated with combined MBP, OAB, and parenteral antibiotics over MBP and parenteral antibiotics alone. However, the most recent of these studies have been flawed in their inclusion of multiple studies based on the NSQIP database which have large degrees of cross-over of the same study population and have mostly focused upon SSI alone rather than other postoperative outcomes. In addition, recent studies^{18,26} have suggested that OAB alone may provide equivalent prophylaxis in terms of SSI and anastomotic leak rates when compared with a combined regimen of MBP+OAB.

The aims of this meta-analysis of RCTs and observational cohort studies in patients undergoing elective colorectal surgery were to:

- Compare the impact of OAB with or without MBP in elective colorectal surgery in terms of SSI, anastomotic leak, 30-day mortality, overall morbidity, development of ileus, reoperations, and *Clostridium difficile* infection.
- Compare evidence derived from RCTs and cohort studies.
- Compare the role of administration of OAB with and without MBP in the setting of laparoscopic versus open surgery.

Methods

Search Strategy

The PubMed, Google Scholar, MEDLINE, and the Cochrane Library databases were searched to identify studies evaluating the effect of OAB in adults undergoing elective colorectal surgery published between January 1, 1981 and May 30, 2018. This date restriction was imposed as recommendations that parenteral antibiotics should be administered routinely for prophylaxis against SSI in colorectal surgery were made in 1981²⁷ and it was felt that all studies considering the role of oral antibiotic prophylaxis should include parenteral antibiotic prophylaxis, to reflect current perioperative care. The search terms used were: (oral antibiotic OR oral antibacterial) AND (colon OR rectal OR colorectal) AND surgery. The bibliographies of all studies which met the inclusion criteria, and previous systematic reviews and meta-analyses on the subject were reviewed to ensure study inclusion was as complete as possible. Non-English-language papers were translated for inclusion. The meta-analysis was conducted in accordance with the PRISMA statement.²⁸

Selection of Articles

Articles were screened for suitability on the basis of title and abstract by 2 independent researchers (K.E.R. and H.J.-E.). Studies were eligible for inclusion if they examined the role of OAB preparation with or without MBP, compared with either MBP alone, OAB alone, or no preparation in adult patients due to undergo elective colorectal surgery, with at

least 1 relevant clinical outcome reported. The type of colorectal surgery performed in terms of type of resection or laparoscopic versus open, the presence or absence of rectal enema administration, or the indication for surgery were not discriminants. Studies were excluded if they did not consider any relevant clinical outcomes, included emergency procedures, or duplicated study populations from other included studies. From the large number of ACS NSQIP studies published 15–20, 26, 29–40 (Supplementary Table 1, <http://links.lww.com/SLA/B542>), only the largest study by Midura et al³¹ was included to avoid the risk of duplication of patient populations within the analysis. Similarly, 3 publications^{41–43} originated from the Michigan Surgical Quality Collaborative Colectomy Best Practices Project. When these were reviewed, 2 studies^{41,42} considered the same comparison of preparations (MBP+OAB vs no preparation), and as such only the more comprehensive study including a larger number of clinical outcomes was included.⁴¹ The third study from the Michigan Surgical Quality Collaborative database⁴³ examined a different preparation combination, thus this was included in the meta-analysis. Finally, the national Veterans Affairs Surgical Quality Improvement Program was the basis for 2 studies^{44,45} on the same regimen comparison, thus only the largest study was included within the meta-analysis.⁴⁵ One study⁴⁶ included a small proportion of patients undergoing emergency colorectal resection within the cohort (311 of a total population of 2240), so any outcomes that included this study were analyzed both with and without it included to discern any difference in results.

Data Extraction

Data were extracted by 2 independent researchers (K.E.R. and H.J.-E.) and any discrepancies were resolved by a senior author (D.N.L.). The primary outcome measure was SSI, with secondary outcome measures including anastomotic leak, 30-day mortality, overall morbidity, development of ileus, reoperation, and *Clostridium difficile* infection. Data were also collected on patient demographics (age, sex), surgical variables (type of resection, open vs. laparoscopic, underlying disease necessitating resection), and details of the preparation used, in terms of parenteral and oral antibiotics as well as MBP. Several studies stated that MBP was not used in patients with obstructing masses, which is mirrored in standard clinical practice, thus these papers were included in the meta-analysis.

The risk of bias was assessed for the RCTs included using the Cochrane Collaboration tool within the RevMan software⁴⁷ which considers 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

Data were entered into RevMan 5.3 software.⁴⁷ Dichotomous variables were calculated as risk ratios (RR) with a 95% confidence interval using the Mantel–Haenszel random effects model. From this, forest plots were derived, with a *P* value of less than 0.05 on 2-tailed testing representing a statistically significant difference. Data from RCTs and cohort studies were included separately within each forest plot, with a summative analysis of all the evidence performed in addition. Inconsistency and heterogeneity between studies were

estimated using the I^2 statistic; 48 25% represented low heterogeneity, 25% to 50% represented moderate, and >50% high heterogeneity.

Protocol Registration

The protocol for this meta-analysis was registered with the PROSPERO database (www.crd.york.ac.uk/prospere)—registration number CRD42018098950.

Results

From the 520 studies identified in the initial search, 40 studies^{31,41,43,45,46,49–83} on 69,517 participants were included (Supplementary Figure 1, <http://links.lww.com/SLA/B542>). Of these 28 were RCTs with 6437 participants^{49–53,55–59,61–67,69–73,75,76,78–80,83} and 12 were cohort (case control) studies with 63,080 participants.^{31,41,43,45,46,54,60,68,74,77,81,82} The risk of bias in the RCTs included was variable, with poor levels of documentation particularly surrounding randomization methods, allocation concealment, and blinding in the earlier studies (Table 1). Six studies^{57,58,62,64–66} administered different parenteral antibiotic regimens depending upon whether the patient was receiving MBP+OAB or MBP alone, which may provide significant source of bias in terms of SSI prevention. In addition, 1 study⁷³ included 2 differing parenteral antibiotic regimens, both in combination with MBP, versus OAB, MBP and parenteral antibiotics. As both of the parenteral antibiotic regimens were considered eligible for inclusion, these were grouped together to form the MBP alone group. In terms of oral antibiotics, 2 studies administered OAB preparation only on the day of surgery; one⁶⁴ gave ciprofloxacin 1 g 1 hour preoperatively and the other⁷⁴ ciprofloxacin 750 mg 1 to 3 hour preoperatively. A subgroup of another study⁵¹ received only 1 dose of OAB the day before surgery, with the remainder receiving 3 doses. These 3 studies may, therefore, have an attenuated the intervention effect from the OAB administered.

Patient Demographics

Two studies^{53,55} focused on surgery using laparoscopic techniques, 21 on open surgery alone,^{46,50,52,57,58,61,62,64–74,76,78,80} with 9 studies^{41,43,49,54,60,75,77,81,82} mixing both open and laparoscopic techniques and the remaining 8 studies not providing this information.^{31,45,51,56,59,63,79,83} The most recent publication³¹ included patients undergoing robotic surgery. The indication for surgery was colorectal cancer in 8 studies,^{46,54,55,59,61,75,78,81} inflammatory bowel disease in 2,^{67,80} with the remaining including a mixture of benign and malignant pathologies. Patient demographics and surgical variables as well as the details of MBP, OAB, and parenteral antibiotics administered are detailed in Table 2.

Surgical Site Infection (SSI)

MBP+OAB Versus MBP—The comparison between MBP+OAB versus MBP alone was performed in 35 studies; 26 RCTs^{49–53,55–59,61–67,69,70,72,73,75,76,78–80} and 9 cohort studies^{31,43,45,54,60,68,74,77,81} with a total of 47,610 patients. When all studies were considered (Fig. 1), the combination of MBP+OAB was associated with a significant reduction in SSI versus MBP alone (RR 0.51, 95% CI 0.46–0.56, $P < 0.00001$, $I^2 = 13\%$).

The results remained consistent when just RCT studies were examined (5378 patients; RR 0.57, 95% CI 0.48–0.68, $P < 0.00001$, $I^2 = 12\%$), as well as cohort studies (42,232 patients; RR 0.48, 95% CI 0.44–0.51, $P < 0.00001$, $I^2 = 0\%$).

MBP+OAB Versus OAB—The analysis of MBP+OAB versus OAB alone was considered by 4 studies; 2 RCTs^{71,83} and 2 cohort studies^{31,45} including 23,483 patients (Fig. 2). Overall, the combination of MBP+OAB was not associated with any difference in the incidence of SSI versus OAB alone (RR 0.98, 95% CI 0.64–1.50, $P = 0.92$), with high heterogeneity ($I^2 = 77\%$). When RCTs alone were considered, again no difference was seen (RR 1.36, 95% CI 0.78–2.35, $P = 0.28$, $I^2 = 0\%$), as with cohort studies (RR 0.83, 95% CI 0.48–1.43, $P = 0.51$, $I^2 = 90\%$).

MBP+OAB Versus No Preparation—No RCTs considered the comparison between combined MBP+OAB and no preparation, with evidence arising from just 4 cohort studies (36,642 patients).^{31,41,45,46} The combination of MBP+OAB was associated with a significant reduction in SSI (RR 0.54, 95% CI 0.43–0.68, $P < 0.00001$, $I^2 = 82\%$) when compared with no preparation.

OAB Alone Versus No Preparation—No RCTs focused upon the comparison between OAB alone versus no preparation, with evidence arising from 16,390 patients included in 2 cohort studies.^{31,45} OAB alone reduced the incidence of SSI versus no preparation (RR 0.56, 95% CI 0.38–0.83, $P = 0.004$, $I^2 = 81\%$).

OAB Versus MBP—Two studies^{31,45} considered the incidence of SSI with OAB alone versus MBP alone, with OAB associated with a reduction in SSI rates. However, this did not reach statistical significance (RR 0.57, 95% CI 0.31–1.05, $P = 0.07$, $I^2 = 93\%$).

Anastomotic Leak

MBP+OAB Versus MBP—Rates of anastomotic leak in those receiving combined MBP+OAB versus MBP alone were compared in 22 studies (Fig. 3); 17 RCTs^{49–53,55,56,58,61,63,64,66,69,70,75,76,78} and 5 cohort studies.^{31,68,74,77,81} Only 2 RCTs^{49,52} included data regarding the management of the anastomotic leak, with none of the 124 patients receiving combined MBP+OAB requiring return to theater for anastomotic leakage compared with 2 of 127 patients receiving MBP alone. Overall, the combination of MBP+OAB was associated with a significant reduction in anastomotic leak rates (RR 0.62, 95% CI 0.55–0.70, $P < 0.00001$, $I^2 = 0\%$), and when evidence from cohort studies alone was considered (RR 0.45, 95% CI 0.25–0.80, $P = 0.007$, $I^2 = 22\%$), but no significant difference was seen when RCTs were analyzed (RR 0.69, 95% CI 0.43–1.11, $P = 0.13$, $I^2 = 0\%$). Six studies^{51,53,55,68,77,81} included data on the use of a diverting stoma, with 133 patients of 1028 in the combined MBP+OAB group and 99 patients of 862 in the MBP alone group undergoing a protective stoma formation.

MBP+OAB Versus OAB—The combination of MBP+OAB versus OAB alone was considered by 3 studies; 2 RCTs^{71,83} and 1 cohort study,³¹ with no difference observed in anastomotic leak rates when all studies (RR 0.79, 95% CI 0.59–1.05, $P = 0.11$, $I^2 = 0\%$), or

just RCTs (RR 1.39, 95% CI 0.47–4.10, $P = 0.55$, $I^2 = 0\%$) were considered (Supplementary Figure 2, <http://links.lww.com/SLA/B542>). No data were available on return to theater rates related to anastomotic leaks.

MBP+OAB Versus No Preparation—The comparison between MBP+OAB versus no preparation in terms of anastomotic leak was considered by just 2 cohort studies, 31,46 with combined MBP+OAB being associated with a significant reduction in anastomotic leak rates (RR 0.52, 95% CI 0.45–0.59, $P < 0.00001$, $I^2 = 0\%$). No data were available on return to theater rates secondary to anastomotic leaks or diverting stoma rates.

Other Comparisons—The comparison of anastomotic leak rates between OAB alone versus no preparation and OAB versus MBP was each only considered by 1 cohort study, 31 and as such meta-analysis was not feasible.

30-day Mortality

MBP+OAB Versus MBP—Seventeen studies (35,633 patients) examined 30-day mortality rates between those receiving MBP+OAB versus MBP alone; 14 RCTs 49,50,52,55,58,59,62,64–66,70,72,76,79 and 3 cohort studies 31,68,74 (Fig. 4). Overall, the combination of MBP+OAB was associated with a significant reduction in 30-day mortality versus MBP alone (RR 0.58, 95% CI 0.44–0.76, $P < 0.0001$, $I^2 = 0\%$). This was also the case when evidence arising from cohort studies alone was considered (RR 0.56, 95% CI 0.42–0.76, $P = 0.0002$, $I^2 = 0\%$), but not when RCTs alone were examined (RR 0.66, 95% CI 0.35–1.25, $P = 0.20$, $I^2 = 0\%$).

MBP+OAB Versus OAB—Three studies (2 RCTs 71,83 and 1 cohort study 31) including 19,360 patients considered 30-day mortality in those receiving MBP+OAB versus OAB alone (Supplementary Figure 3, <http://links.lww.com/SLA/B542>), with the combination being associated with a significant reduction in 30-day mortality in all studies (RR 0.58, 95% CI 0.34–0.97, $P = 0.04$, $I^2 = 0\%$). However, no difference was observed in RCTs (RR 1.02, 95% CI 0.30–3.50, $P = 0.97$, $I^2 = 0\%$).

MBP+OAB Versus No Preparation—Just 2 cohort studies 31,46 including 29,350 patients considered the impact of MBP+OAB versus no preparation on 30-day mortality. The combination of MBP+OAB was associated with a significant reduction in 30-day mortality (RR 0.36, 95% CI 0.17–0.76, $P = 0.008$, $I^2 = 46\%$).

Other Comparisons—Comparison of 30-day mortality between those receiving OAB versus no preparation and OAB versus MBP included just a single cohort study, 31 thus meta-analysis was not conducted.

Overall Morbidity

Only studies comparing MBP+OAB versus MBP alone were considered in terms of overall morbidity rates due to a paucity of data available for all other comparisons. When all 6 studies 31,61,62,66,68,76 (32,568 patients) were compared, the combination of MBP+OAB was associated a significant reduction in overall morbidity (RR 0.67, 95% CI 0.63–0.71, $P <$

0.00001, $I^2 = 0\%$), as well as when evidence from cohort studies alone^{31,68} was considered (RR 0.67, 95% CI 0.63–0.71, $P < 0.00001$, $I^2 = 0\%$). However, with RCTs alone,^{61,62,66,76} there was no difference in overall morbidity between preparation methods (RR 0.71, 95% CI 0.41–1.24, $P = 0.23$, $I^2 = 9\%$).

Development of Ileus

MBP+OAB Versus MBP—Five studies^{31,43,51,53,54} were included in the comparison of MBP+OAB versus MBP; 2 RCTs^{51,53} (879 patients) and 3 cohort studies (33,119 patients).^{31,43,54} Only 1 study⁴³ provided a definition of ileus, with the other 4 studies^{31,43,53,54} not providing a definition. Overall, the combination of MBP+OAB was associated a significant reduction in the incidence of postoperative ileus (RR 0.72, 95% CI 0.52–0.98, $P = 0.04$, $I^2 = 36\%$). However, no difference was seen when just RCTs were considered (RR 0.62, 95% CI 0.14–2.67, $P = 0.52$, $I^2 = 50\%$) or cohort studies alone (RR 0.68, 95% CI 0.45–1.03, $P = 0.07$, $I^2 = 53\%$).

MBP+OAB Versus OAB—Three studies^{31,71,83} were included in the comparison between MBP+OAB versus OAB; 2 RCTs^{71,83} and 1 cohort study.³¹ None of these studies provided a definition for ileus. Overall, the combination of MBP+OAB was associated with a significant reduction in the incidence of postoperative ileus (RR 0.83, 95% CI 0.73–0.95, $P = 0.008$, $I^2 = 0\%$), mostly determined by the large single cohort study.³¹ However, no difference was seen when RCTs were considered (RR 1.25, 95% CI 0.68–2.33, $P = 0.47$, $I^2 = 0\%$).

MBP+OAB Versus No Preparation—No RCTs considered the comparison between MBP+OAB versus no preparation, with evidence arising from 2 cohort studies only.^{31,41} Only 1 study⁴¹ provided a definition of ileus. This demonstrated that the combination of MBP+OAB was associated with a significant reduction in ileus (RR 0.72, 95% CI 0.68–0.77, $P < 0.00001$, $I^2 = 0\%$).

Other Comparisons—The comparison in reoperation rates between OAB alone versus no preparation and OAB versus MBP were each only considered by 1 cohort study,³¹ thus meta-analysis was not performed.

Reoperation

Insufficient data were available for any of the planned analyses on reoperation rates, with 2 studies including data comparing MBP+OAB versus MBP (1 RCT⁴⁹ and 1 cohort study³¹), and just 2 studies comparing MBP+OAB versus OAB alone (again 1 RCT⁷¹ and 1 cohort study).³¹ Thus, no meta-analysis was performed. The comparisons of reoperation rates between MBP+OAB versus no preparation, OAB alone versus no preparation and OAB versus MBP were each only considered by 1 cohort study,³¹ and as such meta-analysis was not performed. However, the largest cohort study³¹ showed a significant reduction ($P < 0.001$) in reoperation rates with combined MBP+OAB (3.2%) compared with OAB alone (4.7%), MBP alone (4.2%), and no preparation (4.5%).

***Clostridium difficile* Infection**

MBP+OAB Versus MBP—Data on *Clostridium difficile* infection were sufficient only for the comparison between MBP+OAB versus MBP alone, with data from 14 studies, including 10 RCTs^{53,55,61,62,65,67,69,75,78,80} and 4 cohort studies.^{43,54,68,82} No difference in *C difficile* infection rates were seen when all evidence was considered (RR 0.94, 95% CI 0.55–1.61, $P = 0.81$, $I^2 = 37\%$), nor when just RCT studies or cohort studies alone were analyzed (RR 0.79, 95% CI 0.21–2.96, $P = 0.72$, $I^2 = 10\%$ and RR 0.97, 95% CI 0.54–1.75, $P = 0.92$, $I^2 = 64\%$, respectively).

Laparoscopic Versus Open Procedures

Nineteen RCTs^{50,52,57,58,61–67,69,70,72–74,76,79,80} provided data on SSI rates in patients undergoing open elective colorectal procedures between patients receiving combined MBP+OAB versus MBP alone, and 2 RCTs^{53,55} provided data on laparoscopic procedures alone. The remaining studies included either both open and laparoscopic procedures which could not be separated for analysis or did not state the surgical approach. No other comparison between preparations was considered due to a paucity of data. The combination of MBP+OAB versus MBP alone was associated with a significant reduction in SSI rates in patients undergoing an open resection (RR 0.55, 95% CI 0.44–0.69, $P < 0.00001$, $I^2 = 5\%$); however, no significant difference was seen in patients undergoing a laparoscopic procedure (RR 0.74, 95% CI 0.43–1.29, $P = 0.29$, $I^2 = 50\%$), although it should be borne in mind that this evidence was based upon 2 studies (1090 patients).

When anastomotic leak rates were compared between MBP+OAB versus MBP alone, divided by open and laparoscopic procedures, data could be analyzed from 9 RCTs^{50,52,58,61,64,66,69,70,76} in the open group and 2 RCTs^{53,55} in the laparoscopic group. There was no significant difference in anastomotic leak rates in either the open or laparoscopic groups (RR 0.69, 95% CI 0.30–1.60, $P = 0.39$, $I^2 = 13\%$ and RR 0.68, 95% CI 0.28–1.65, $P = 0.39$, $I^2 = 0\%$, respectively).

Discussion

Main Findings

This meta-analysis has provided evidence to suggest that MBP+OAB should be given serious consideration in patients undergoing elective colorectal surgery to reduce the risk of SSI. In addition, it has shown that the combination of MBP+OAB is associated with significant reductions in anastomotic leak rates, 30-day mortality, overall morbidity, and the incidence of postoperative ileus, without increasing the risk of developing *C difficile* infection (Table 3). Its findings are in contradiction with previous meta-analyses^{1,2} that did not account for the role of luminal antibiotics and showed that MBP on its own was of no benefit when compared with no bowel preparation or rectal enemas alone.

However, as only 9.3% (6437 patients) of the 69,517 patients included were studied in the context of RCTs, the results must be interpreted with some caution. Hence, when evidence arising from RCTs alone was considered, the combination of MBP+OAB was associated with a significant reduction in SSI alone. The evidence for the combination of MBP+OAB to

reduce SSI rates is, thus, strong. European data reporting the results of colorectal surgery in the context of Enhanced Recovery After Surgery protocols where mechanical bowel preparation is not used routinely, have shown SSI rates of >10%,^{84,85} whereas the US NSQIP studies have shown that SSI rates are approximately 3% with a combination of MBP +OAB, 6% with MBP alone and 7% with no preparation.³¹

When the combination of MBP+OAB was compared with OAB alone, a significant reduction in 30-day mortality and incidence of postoperative ileus was seen, but no difference was seen between the 2 preparations in RCTs alone. There are no RCTs focusing on the combinations of MBP+OAB versus no preparation, OAB alone versus no preparation or OAB alone versus MBP alone. However, evidence from cohort studies suggests that the combination of MBP+OAB versus no preparation is associated with a significant reduction in SSI, anastomotic leak, 30-day mortality, and postoperative ileus. For OAB versus no preparation, the only significant reduction was in SSI rates, and for OAB versus MBP there was no significant difference in any of the clinical outcome measures. When a planned subgroup analysis of patients undergoing open versus laparoscopic surgery was undertaken, the combination of MBP+OAB versus MBP alone was associated with a significant reduction in SSI rates in patients undergoing open procedures, but not in those undergoing laparoscopic procedures.

Strengths and Weaknesses

The main weakness of this meta-analysis is the inclusion of both RCTs and cohort studies. While this lowers the overall quality of evidence, the decision to include cohort studies and large database studies was made as a large proportion of the recent evidence supporting the potential role of OAB or combined MBP+OAB has arisen from such studies. However, every analysis was conducted separately using evidence from RCT and cohort studies alone, as well as a summative analysis, to provide a more robust interpretation of the data.

The role of parenteral antibiotic prophylaxis is considered a standard of care in current practice, with evidence published in 1981²⁷ providing evidence for its benefit in terms of infection prevention and overall mortality and dictating that no further placebo or no intervention trials should be conducted. Definitive support was provided in a Cochrane Review⁸⁶ demonstrating a significant reduction in SSI in patients receiving parenteral antibiotic prophylaxis versus those receiving no antibiotics or placebo (RR 0.34, 95% CI 0.28–0.41, $P < 0.0001$).

The practice of mechanical bowel preparation has changed significantly since the early 1980s. The regimen of Lazorthes et al⁶² included admission 3 days prior to surgery and administration of a low-residue diet and standard mechanical procedures such as enemas and magnesium sulphate purges. In contrast, more modern regimens are typically administered the day before surgery and are less invasive. This is particularly important in the setting of prolonged starvation protocols in vogue prior to the more modern ones, as they resulted in increased preoperative dehydration and electrolyte disturbances which are known to have adverse effects on postoperative complications. It should, however, be considered that each study level comparison between preparation types should have been exposed to the same level of bias, thus making the results more comparable. The OAB agent, dosing, and timing

as well as the parenteral antibiotic details were also inconsistent between studies, with insufficient data from each differing combination to perform a meaningful analysis. Several included just 1 preoperative dose of OAB, or differing parenteral antibiotic regimens depending upon which preparation regimen the patient received which exerts a potential significant bias. In addition, because of limited data, we have been unable to discern conclusively whether the reduction in morbidity is a result of OAB on their own or in combination with MBP.

The definition of anastomotic leak was not stipulated for inclusion within this meta-analysis, with the data from each individual study included, irrespective of whether this was based upon clinical or radiological diagnosis of anastomotic leak. However, the definition of leak was consistent within individual studies, thus the data from each study were comparable, attenuating this potential weakness.

Interpretation of the Data in Context of Other Recent Studies

A recent meta-analysis²⁵ included 23 RCTs and 8 cohort studies published between 1980 and 2015. However, multiple cohort studies arising from the NSQIP database were included within this study,²⁵ and this probably represents multiple reporting of the same patient datasets. This study²⁵ reported a significant reduction in SSI rates in patients included within cohort studies receiving MBP, OAB, and IV antibiotics versus those receiving MBP and IV antibiotics alone (RR 0.48, 95% CI 0.44–0.52, $P=0.00001$, $I^2=45\%$). However, 4 of the 5 studies included within this analysis arose from the ACS NSQIP database. Bellows et al²³ previously performed a meta-analysis on the role of oral nonabsorbable and intravenous antibiotics versus intravenous antibiotics alone in colorectal surgery, focusing on SSI. This study included 16 RCTs encompassing 2669 patients published between 1980 and 2011, with all studies including MBP within the protocol. This meta-analysis found that the combination of oral and IV antibiotics versus IV antibiotics alone was associated with a significant reduction in wound infection rates (RR 0.57, 95% CI 0.43–0.76, $P=0.0002$, $I^2=19\%$), but no significant difference in anastomotic leak rates (RR 0.63, 95% CI 0.28–1.41, $P=0.3$, $I^2=0\%$). The findings of the currently reported meta-analysis coincide with the results of these previous meta-analyses.

Conclusion

The present meta-analysis is the largest and most comprehensive to date examining the role of bowel preparation prior to colorectal surgery, and supports a potentially significant benefit for OAB preparation, either in combination with MBP or alone, in the prevention of postoperative complications. While evidence arising from large retrospective cohort and database studies suggests a strong positive benefit, these are tempered when evidence arising from RCTs alone is considered. However, the evidence presented would suggest a benefit from OAB preparation in terms of SSI, which represents a major source of morbidity and increased healthcare costs. Further high-quality evidence is required to differentiate between the benefits of combined MBP+OAB or OAB alone in this setting before more definitive recommendations can be made.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

This work was supported by the Medical Research Council [grant number MR/K00414X/1]; and Arthritis Research UK [grant number 19891].

References

1. Rollins KE, Javanmard-Emamghissi H, Lobo DN. Impact of mechanical bowel preparation in elective colorectal surgery: a meta-analysis. *World J Gastroenterol*. 2018; 24:519–536. [PubMed: 29398873]
2. Slim K, Vicaut E, Launay-Savary MV, et al. Updated systematic review and meta-analysis of randomized clinical trials on the role of mechanical bowel preparation before colorectal surgery. *Ann Surg*. 2009; 249:203–209. [PubMed: 19212171]
3. Gustafsson UO, Scott MJ, Schwenk W, et al. Guidelines for perioperative care in elective colonic surgery: Enhanced Recovery After Surgery (ERAS(R)) Society recommendations. *Clin Nutr*. 2012; 31:783–800. [PubMed: 23099039]
4. Nygren J, Thacker J, Carli F, et al. Guidelines for perioperative care in elective rectal/pelvic surgery: Enhanced Recovery After Surgery (ERAS(R)) Society recommendations. *Clin Nutr*. 2012; 31:801–816. [PubMed: 23062720]
5. National Institute for Health and Care Excellence. Surgical site infections: prevention and treatment—Clinical guideline (CG74). London: National Institute for Health and Care Excellence; 2008. (Updated 2017) Available at: <https://www.nice.org.uk/guidance/cg74/resources/surgical-site-infections-prevention-and-treatment-pdf-975628422853> [Accessed April 15, 2018]
6. Holubar SD, Hedrick T, Gupta R, et al. American Society for Enhanced Recovery (ASER) and Perioperative Quality Initiative (POQI) joint consensus statement on prevention of postoperative infection within an enhanced recovery pathway for elective colorectal surgery. *Perioper Med (Lond)*. 2017; 6:4. [PubMed: 28270910]
7. Rosenberg IL, Graham NG, De Dombal FT, et al. Preparation of the intestine in patients undergoing major large-bowel surgery, mainly for neoplasms of the colon and rectum. *Br J Surg*. 1971; 58:266–269. [PubMed: 5108232]
8. Washington JA 2nd, Dearing WH, Judd ES, et al. Effect of preoperative antibiotic regimen on development of infection after intestinal surgery: Prospective, randomized, double-blind study. *Ann Surg*. 1974; 180:567–572. [PubMed: 4606495]
9. Clarke JS, Condon RE, Bartlett JG, et al. Preoperative oral antibiotics reduce septic complications of colon operations: results of prospective, randomized, double-blind clinical study. *Ann Surg*. 1977; 186:251–259. [PubMed: 889372]
10. Nichols RL, Condon RE, DiSanto AR. Preoperative bowel preparation. Erythromycin base serum and fecal levels following oral administration. *Arch Surg*. 1977; 112:1493–1496. [PubMed: 931637]
11. Holte K, Nielsen KG, Madsen JL, et al. Physiologic effects of bowel preparation. *Dis Colon Rectum*. 2004; 47:1397–1402. [PubMed: 15484356]
12. Shapira Z, Feldman L, Lavy R, et al. Bowel preparation: comparing metabolic and electrolyte changes when using sodium phosphate/polyethylene glycol. *Int J Surg*. 2010; 8:356–358. [PubMed: 20457286]
13. Cirocco WC. The fatal flaw of outcome studies comparing colorectal operations with and without mechanical bowel preparation: the absence of oral antibiotics! *Dis Colon Rectum*. 2016; 59:e421. [PubMed: 27384100]
14. Zelhart MD, Hauch AT, Slakey DP, et al. Preoperative antibiotic colon preparation: have we had the answer all along? *J Am Coll Surg*. 2014; 219:1070–1077. [PubMed: 25260679]

15. Scarborough JE, Mantyh CR, Sun Z, et al. Combined mechanical and oral antibiotic bowel preparation reduces incisional surgical site infection and anastomotic leak rates after elective colorectal resection: an analysis of colectomy-targeted ACS NSQIP. *Ann Surg.* 2015; 262:331–337. [PubMed: 26083870]
16. Klinger AL, Green H, Monlezun DJ, et al. The role of bowel preparation in colorectal surgery: results of the 2012–2015 ACS-NSQIP data. *Ann Surg.* 2017
17. Haskins IN, Fleshman JW, Amdur RL, et al. The impact of bowel preparation on the severity of anastomotic leak in colon cancer patients. *J Surg Oncol.* 2016; 114:810–813. [PubMed: 27634398]
18. Garfinkle R, Abou-Khalil J, Morin N, et al. Is there a role for oral antibiotic preparation alone before colorectal surgery? ACS-NSQIP analysis by coarsened exact matching. *Dis Colon Rectum.* 2017; 60:729–737. [PubMed: 28594723]
19. Dolejs SC, Guzman MJ, Fajardo AD, et al. Bowel preparation is associated with reduced morbidity in elderly patients undergoing elective colectomy. *J Gastrointest Surg.* 2017; 21:372–379. [PubMed: 27896654]
20. Kiran RP, Murray AC, Chiuzan C, et al. Combined preoperative mechanical bowel preparation with oral antibiotics significantly reduces surgical site infection, anastomotic leak, and ileus after colorectal surgery. *Ann Surg.* 2015; 262:416–425. [PubMed: 26258310]
21. Yost MT, Jolissaint JS, Fields AC, et al. Mechanical and oral antibiotic bowel preparation in the era of minimally invasive surgery and enhanced recovery. *J Laparoendosc Adv Surg Tech A.* 2018; 28:491–495. [PubMed: 29630437]
22. Badia JM, Arroyo-Garcia N. Mechanical bowel preparation and oral antibiotic prophylaxis in colorectal surgery: analysis of evidence and narrative review. *Cir Esp.* 2018; 96:317–325. [PubMed: 29773260]
23. Bellows CF, Mills KT, Kelly TN, et al. Combination of oral non-absorbable and intravenous antibiotics versus intravenous antibiotics alone in the prevention of surgical site infections after colorectal surgery: a meta-analysis of randomized controlled trials. *Tech Coloproctol.* 2011; 15:385–395. [PubMed: 21785981]
24. McSorley ST, Steele CW, McMahan AJ. Meta-analysis of oral antibiotics, in combination with preoperative intravenous antibiotics and mechanical bowel preparation the day before surgery, compared with intravenous antibiotics and mechanical bowel preparation alone to reduce surgical-site infections in elective colorectal surgery. *BJS Open.* 2018; 2:185–194. [PubMed: 30079387]
25. Koullouros M, Khan N, Aly EH. The role of oral antibiotics prophylaxis in prevention of surgical site infection in colorectal surgery. *Int J Colorectal Dis.* 2017; 32:1–18. [PubMed: 27778060]
26. Parthasarathy M, Greensmith M, Bowers D, et al. Risk factors for anastomotic leakage after colorectal resection: a retrospective analysis of 17 518 patients. *Colorectal Dis.* 2017; 19:288–298. [PubMed: 27474844]
27. Baum ML, Anish DS, Chalmers TC, et al. A survey of clinical trials of antibiotic prophylaxis in colon surgery: evidence against further use of no-treatment controls. *N Engl J Med.* 1981; 305:795–799. [PubMed: 7266633]
28. Moher D, Liberati A, Tetzlaff J, et al. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. *Int J Surg.* 2010; 8:336–341. [PubMed: 20171303]
29. Althumairi AA, Canner JK, Pawlik TM, et al. Benefits of bowel preparation beyond surgical site infection: a retrospective study. *Ann Surg.* 2016; 264:1051–1057. [PubMed: 26727098]
30. Connolly TM, Foppa C, Kazi E, et al. Impact of a surgical site infection reduction strategy after colorectal resection. *Colorectal Dis.* 2016; 18:910–918. [PubMed: 26456021]
31. Midura EF, Jung AD, Hanseman DJ, et al. Combination oral and mechanical bowel preparations decreases complications in both right and left colectomy. *Surgery.* 2018; 163:528–534. [PubMed: 29198768]
32. Moghadamyeghaneh Z, Hanna MH, Carmichael JC, et al. Nationwide analysis of outcomes of bowel preparation in colon surgery. *J Am Coll Surg.* 2015; 220:912–920. [PubMed: 25907871]
33. Moghadamyeghaneh Z, Hwang GS, Hanna MH, et al. Risk factors for prolonged ileus following colon surgery. *Surg Endosc.* 2016; 30:603–609. [PubMed: 26017914]

34. Morris MS, Graham LA, Chu DI, et al. Oral antibiotic bowel preparation significantly reduces surgical site infection rates and readmission rates in elective colorectal surgery. *Ann Surg.* 2015; 261:1034–1040. [PubMed: 25607761]
35. Ohman KA, Wan L, Guthrie T, et al. Combination of oral antibiotics and mechanical bowel preparation reduces surgical site infection in colorectal surgery. *J Am Coll Surg.* 2017; 225:465–471. [PubMed: 28690206]
36. Parthasarathy M, Bowers D, Groot-Wassink T. Do preoperative oral antibiotics increase *Clostridium difficile* infection rates? An analysis of 13 959 colectomy patients. *Colorectal Dis.* 2018; 20:520–528. [PubMed: 29045025]
37. Rencuzogullari A, Benlice C, Costedio M, et al. Nomogram-derived prediction of postoperative ileus after colectomy: an assessment from nationwide procedure-targeted cohort. *Am Surg.* 2017; 83:564–572. [PubMed: 28637557]
38. Rencuzogullari A, Benlice C, Valente M, et al. Predictors of anastomotic leak in elderly patients after colectomy: nomogram-based assessment from the American College of Surgeons National Surgical Quality Program Procedure-Targeted Cohort. *Dis Colon Rectum.* 2017; 60:527–536. [PubMed: 28383453]
39. Shwaartz C, Fields AC, Sobrero M, et al. Does bowel preparation for inflammatory bowel disease surgery matter? *Colorectal Dis.* 2017; 19:832–839. [PubMed: 28436176]
40. Tevis SE, Carchman EH, Foley EF, et al. Does anastomotic leak contribute to high failure-to-rescue rates? *Ann Surg.* 2016; 263:1148–1151. [PubMed: 26587851]
41. Kim EK, Sheetz KH, Bonn J, et al. A statewide colectomy experience: the role of full bowel preparation in preventing surgical site infection. *Ann Surg.* 2014; 259:310–314. [PubMed: 23979289]
42. Hendren S, Fritze D, Banerjee M, et al. Antibiotic choice is independently associated with risk of surgical site infection after colectomy: a population-based cohort study. *Ann Surg.* 2013; 257:469–475. [PubMed: 23059498]
43. Englesbe MJ, Brooks L, Kubus J, et al. A statewide assessment of surgical site infection following colectomy: the role of oral antibiotics. *Ann Surg.* 2010; 252:514–519. [PubMed: 20739852]
44. Toneva GD, Deierhoi RJ, Morris M, et al. Oral antibiotic bowel preparation reduces length of stay and readmissions after colorectal surgery. *J Am Coll Surg.* 2013; 216:756–762. [PubMed: 23521958]
45. Cannon JA, Altom LK, Deierhoi RJ, et al. Preoperative oral antibiotics reduce surgical site infection following elective colorectal resections. *Dis Colon Rectum.* 2012; 55:1160–1166. [PubMed: 23044677]
46. Mik M, Berut M, Trzcinski R, et al. Preoperative oral antibiotics reduce infections after colorectal cancer surgery. *Langenbecks Arch Surg.* 2016; 401:1153–1162. [PubMed: 27650707]
47. Review Manager (RevMan) [Computer program]—Version 5.3. The Nordic Cochrane Centre, The Cochrane Collaboration; Copenhagen: 2014. Available at <https://community.cochrane.org/help/tools-and-software/revman-5> [Accessed 20 June 2018]
48. Higgins JP, Thompson SG. Quantifying heterogeneity in a meta-analysis. *Stat Med.* 2002; 21:1539–1558. [PubMed: 12111919]
49. Anjum N, Ren J, Wang G, et al. A randomized control trial of preoperative oral antibiotics as adjunct therapy to systemic antibiotics for preventing surgical site infection in clean contaminated, contaminated, and dirty type of colorectal surgeries. *Dis Colon Rectum.* 2017; 60:1291–1298. [PubMed: 29112565]
50. Coppa GF, Eng K. Factors involved in antibiotic selection in elective colon and rectal surgery. *Surgery.* 1988; 104:853–858. [PubMed: 3055394]
51. Espin-Basany E, Sanchez-Garcia JL, Lopez-Cano M, et al. Prospective, randomised study on antibiotic prophylaxis in colorectal surgery. Is it really necessary to use oral antibiotics? *Int J Colorectal Dis.* 2005; 20:542–546. [PubMed: 15843938]
52. Hanel KC, King DW, McAllister ET, et al. Single-dose parenteral antibiotics as prophylaxis against wound infections in colonic operations. *Dis Colon Rectum.* 1980; 23:98–101. [PubMed: 7379664]

53. Hata H, Yamaguchi T, Hasegawa S, et al. Oral and parenteral versus parenteral antibiotic prophylaxis in elective laparoscopic colorectal surgery (JMTO PREV 07-01): a phase 3, multicenter, open-label, randomized trial. *Ann Surg.* 2016; 263:1085–1091. [PubMed: 26756752]
54. Ichimanda M, Etoh T, Nakajima K, et al. The efficacy of kanamycin plus metronidazole administration as an OABP against incisional surgical site infection in colorectal cancer surgery. *Nippon Daicho Komonbyo Gakkai Zasshi.* 2017; 70:214–221.
55. Ikeda A, Konishi T, Ueno M, et al. Randomized clinical trial of oral and intravenous versus intravenous antibiotic prophylaxis for laparoscopic colorectal resection. *Br J Surg.* 2016; 103:1608–1615. [PubMed: 27550722]
56. Ishida H, Yokoyama M, Nakada H, et al. Impact of oral antimicrobial prophylaxis on surgical site infection and methicillin-resistant *Staphylococcus aureus* infection after elective colorectal surgery. Results of a prospective randomized trial. *Surg Today.* 2001; 31:979–983. [PubMed: 11766085]
57. Kaiser AB, Herrington JL Jr, Jacobs JK, et al. Cefoxitin versus erythromycin, neomycin, and cefazolin in colorectal operations. Importance of the duration of the surgical procedure. *Ann Surg.* 1983; 198:525–530. [PubMed: 6354113]
58. Khubchandani IT, Karamchandani MC, Sheets JA, et al. Metronidazole vs. erythromycin, neomycin, and cefazolin in prophylaxis for colonic surgery. *Dis Colon Rectum.* 1989; 32:17–20. [PubMed: 2910656]
59. Kobayashi M, Mohri Y, Tonouchi H, et al. Randomized clinical trial comparing intravenous antimicrobial prophylaxis alone with oral and intravenous antimicrobial prophylaxis for the prevention of a surgical site infection in colorectal cancer surgery. *Surg Today.* 2007; 37:383–388. [PubMed: 17468819]
60. Konishi T, Watanabe T, Kishimoto J, et al. Elective colon and rectal surgery differ in risk factors for wound infection: results of prospective surveillance. *Ann Surg.* 2006; 244:758–763. [PubMed: 17060769]
61. Lau WY, Chu KW, Poon GP, et al. Prophylactic antibiotics in elective colorectal surgery. *Br J Surg.* 1988; 75:782–785. [PubMed: 3167527]
62. Lazorthes F, Legrand G, Monrozier X, et al. Comparison between oral and systemic antibiotics and their combined use for the prevention of complications in colorectal surgery. *Dis Colon Rectum.* 1982; 25:309–311. [PubMed: 7044724]
63. Lewis RT. Oral versus systemic antibiotic prophylaxis in elective colon surgery: a randomized study and meta-analysis send a message from the 1990s. *Can J Surg.* 2002; 45:173–180. [PubMed: 12067168]
64. McArdle CS, Morran CG, Pettit L, et al. Value of oral antibiotic prophylaxis in colorectal surgery. *Br J Surg.* 1995; 82:1046–1048. [PubMed: 7648148]
65. Monrozier X, Lazorthes F, Fretigny E, et al. Evaluation of systemic antibiotic preventive treatment in colorectal surgery. *J Chir (Paris).* 1983; 120:393–396. [PubMed: 6619217]
66. Nohr M, Andersen JC, Juul-Jensen KE. Prophylactic single-dose fosfomycin and metronidazole compared with neomycin, bacitracin, metronidazole and ampicillin in elective colorectal operations. *Acta Chir Scand.* 1990; 156:223–230. [PubMed: 2186587]
67. Oshima T, Takesue Y, Ikeuchi H, et al. Preoperative oral antibiotics and intravenous antimicrobial prophylaxis reduce the incidence of surgical site infections in patients with ulcerative colitis undergoing IPAA. *Dis Colon Rectum.* 2013; 56:1149–1155. [PubMed: 24022532]
68. Ozdemir S, Gulpinar K, Ozis SE, et al. The effects of preoperative oral antibiotic use on the development of surgical site infection after elective colorectal resections: a retrospective cohort analysis in consecutively operated 90 patients. *Int J Surg.* 2016; 33(pt A):102–108. [PubMed: 27463886]
69. Peruzzo L, Savio S, De Lalla F. Systemic versus systemic plus oral chemoprophylaxis in elective colorectal surgery. *Chimioterapia.* 1987; 6:601–603. [PubMed: 3334642]
70. Playforth MJ, Smith GM, Evans M, et al. Antimicrobial bowel preparation. Oral, parenteral, or both? *Dis Colon Rectum.* 1988; 31:90–93. [PubMed: 3276469]
71. Ram E, Sherman Y, Weil R, et al. Is mechanical bowel preparation mandatory for elective colon surgery? A prospective randomized study. *Arch Surg.* 2005; 140:285–288. [PubMed: 15781794]

72. Reddy BS, Macfie J, Gatt M, et al. Randomized clinical trial of effect of synbiotics, neomycin and mechanical bowel preparation on intestinal barrier function in patients undergoing colectomy. *Br J Surg*. 2007; 94:546–554. [PubMed: 17443852]
73. Reynolds J, Jones J, Evans D, et al. Do preoperative oral antibiotics influence sepsis rates following elective colorectal surgery in patients receiving perioperative intravenous prophylaxis. *Surg Res Commun*. 1989; 7:71–77.
74. Rohwedder R, Bonadeo F, Benati M, et al. Single-dose oral ciprofloxacin plus parenteral metronidazole for perioperative antibiotic prophylaxis in colorectal surgery. *Chemotherapy*. 1993; 39:218–224. [PubMed: 8508692]
75. Sadahiro S, Suzuki T, Tanaka A, et al. Comparison between oral antibiotics and probiotics as bowel preparation for elective colon cancer surgery to prevent infection: prospective randomized trial. *Surgery*. 2014; 155:493–503. [PubMed: 24524389]
76. Stellato TA, Danziger LH, Gordon N, et al. Antibiotics in elective colon surgery. A randomized trial of oral, systemic, and oral/systemic antibiotics for prophylaxis. *Am Surg*. 1990; 56:251–254. [PubMed: 2194417]
77. Sun W-C, Hsu H-H, Liu H-C, et al. Can mechanical bowel preparation with oral antibiotics reduce surgical site infection and anastomotic leakage rates following elective colorectal resections? *Formos J Surg*. 2018; 51:21–25.
78. Takesue Y, Yokoyama T, Akagi S, et al. A brief course of colon preparation with oral antibiotics. *Surg Today*. 2000; 30:112–116. [PubMed: 10664331]
79. Taylor EW, Lindsay G. Selective decontamination of the colon before elective colorectal surgery. West of Scotland Surgical Infection Study Group. *World J Surg*. 1994; 18:926–931. [PubMed: 7846921]
80. Uchino M, Ikeuchi H, Bando T, et al. Efficacy of preoperative oral antibiotic prophylaxis for the prevention of surgical site infections in patients with Crohn disease: a randomized controlled trial. *Ann Surg*. 2017
81. Vo E, Massarweh NN, Chai CY, et al. Association of the addition of oral antibiotics to mechanical bowel preparation for left colon and rectal cancer resections with reduction of surgical site infections. *JAMA Surg*. 2018; 153:114–121. [PubMed: 29049477]
82. Wren SM, Ahmed N, Jamal A, et al. Preoperative oral antibiotics in colorectal surgery increase the rate of *Clostridium difficile* colitis. *Arch Surg*. 2005; 140:752–756. [PubMed: 16103284]
83. Zmora O, Mahajna A, Bar-Zakai B, et al. Colon and rectal surgery without mechanical bowel preparation: a randomized prospective trial. *Ann Surg*. 2003; 237:363–367. [PubMed: 12616120]
84. ERAS Compliance Group. The impact of enhanced recovery protocol compliance on elective colorectal cancer resection: results from an international registry. *Ann Surg*. 2015; 261:1153–1159. [PubMed: 25671587]
85. Hendry PO, Hausel J, Nygren J, et al. Determinants of outcome after colorectal resection within an enhanced recovery programme. *Br J Surg*. 2009; 96:197–205. [PubMed: 19160347]
86. Nelson RL, Gladman E, Barbateskovic M. Antimicrobial prophylaxis for colorectal surgery. *Cochrane Database Syst Rev*. 2014; 5

Mini Abstract

This meta-analysis examines the role of oral antibiotic preparation, with and without mechanical bowel preparation in elective colorectal surgery. Combined oral antibiotic and mechanical bowel preparation significantly reduces surgical site infection rates *versus* mechanical preparation alone, but this effect is similar to that of oral antibiotics alone.

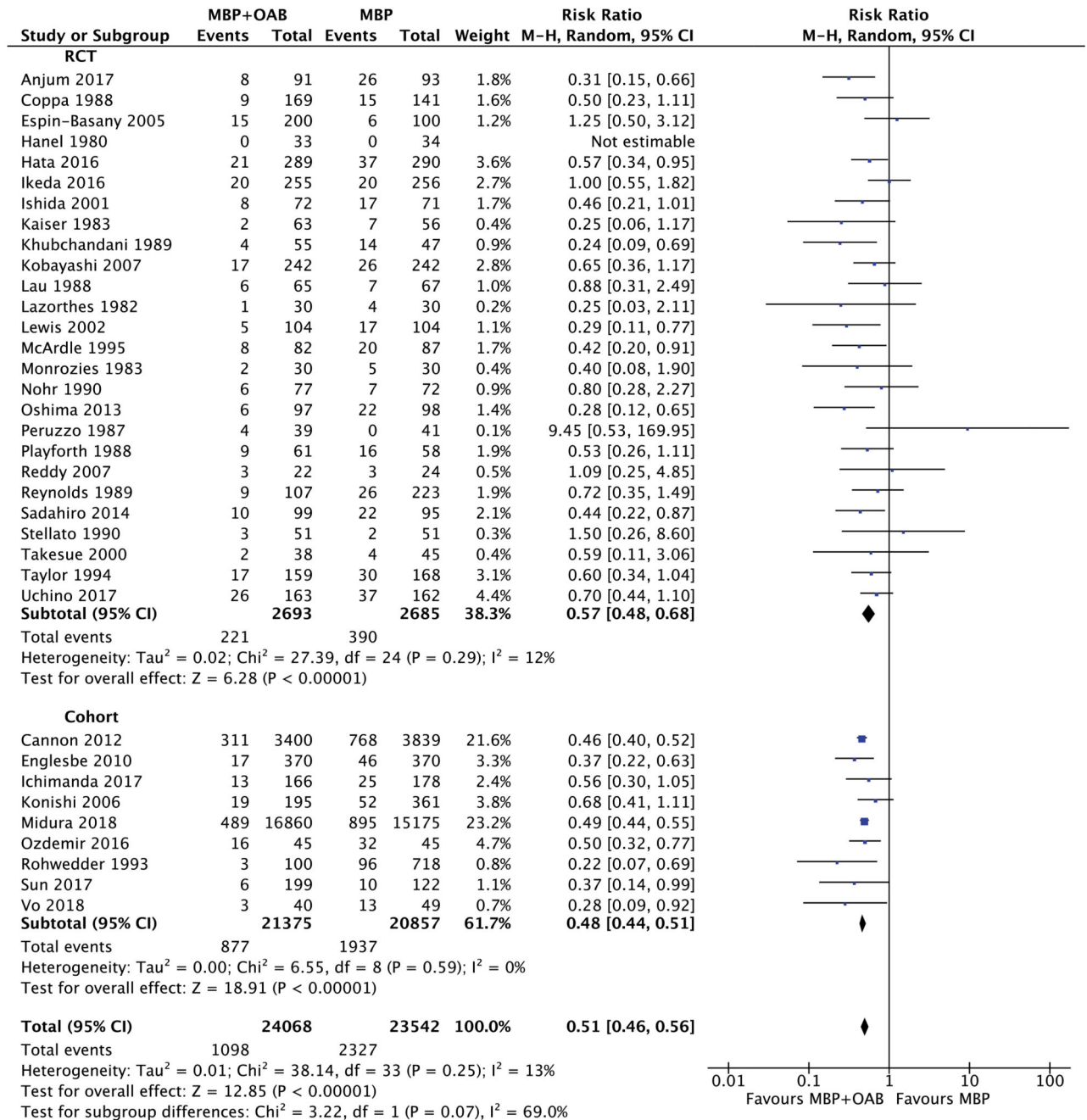


Figure 1. Forest plot comparing surgical site infection rate for patients receiving MBP+OAB versus MBP alone, divided by evidence from RCTs and cohort studies. A Mantel–Haenszel random effects model was used to perform the meta-analysis and risk ratios are quoted including 95% confidence intervals.

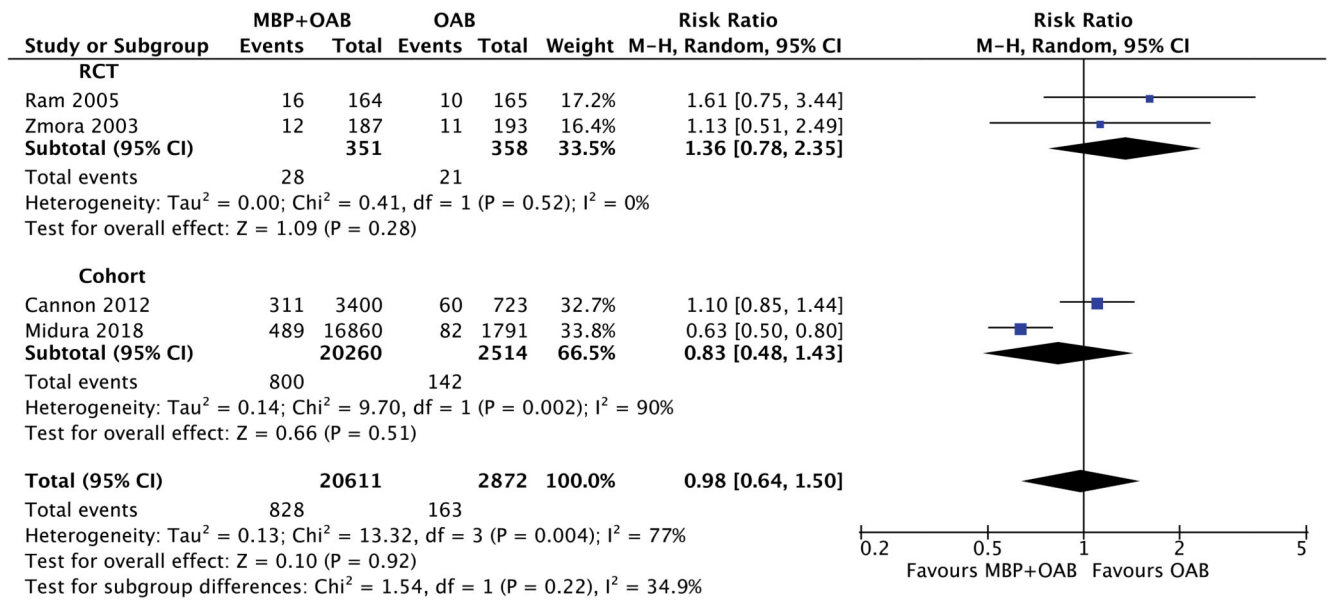


Figure 2. Forest plot comparing surgical site infection rate for patients receiving MBP+OAB versus OAB alone, divided by evidence from RCTs and cohort studies. A Mantel–Haenszel random effects model was used to perform the meta-analysis and risk ratios are quoted including 95% confidence intervals.

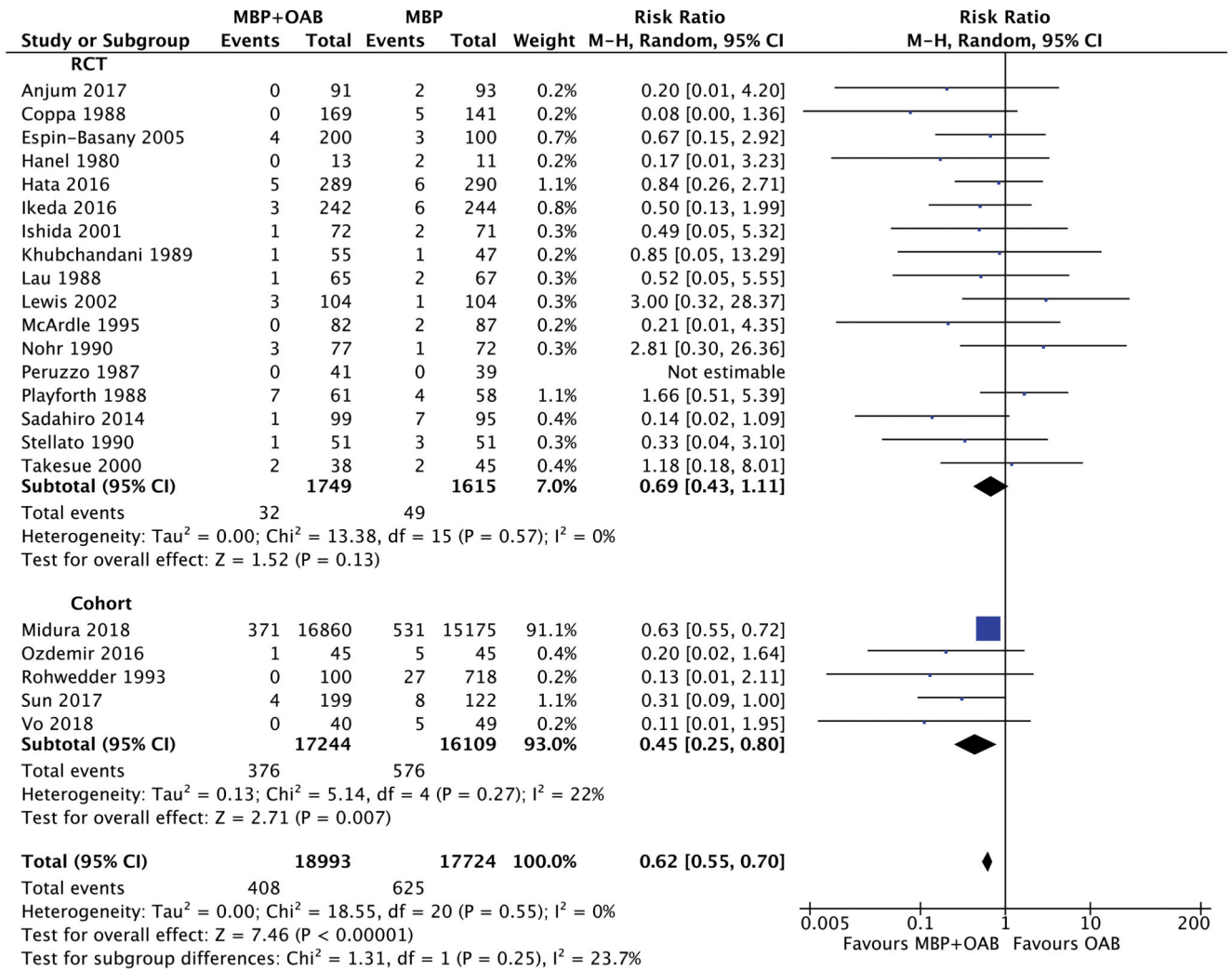


Figure 3. Forest plot comparing anastomotic leak rate for patients receiving MBP+OAB versus MBP alone, divided by evidence from RCTs and cohort studies. A Mantel-Haenszel random effects model was used to perform the meta-analysis and risk ratios are quoted including 95% confidence intervals.

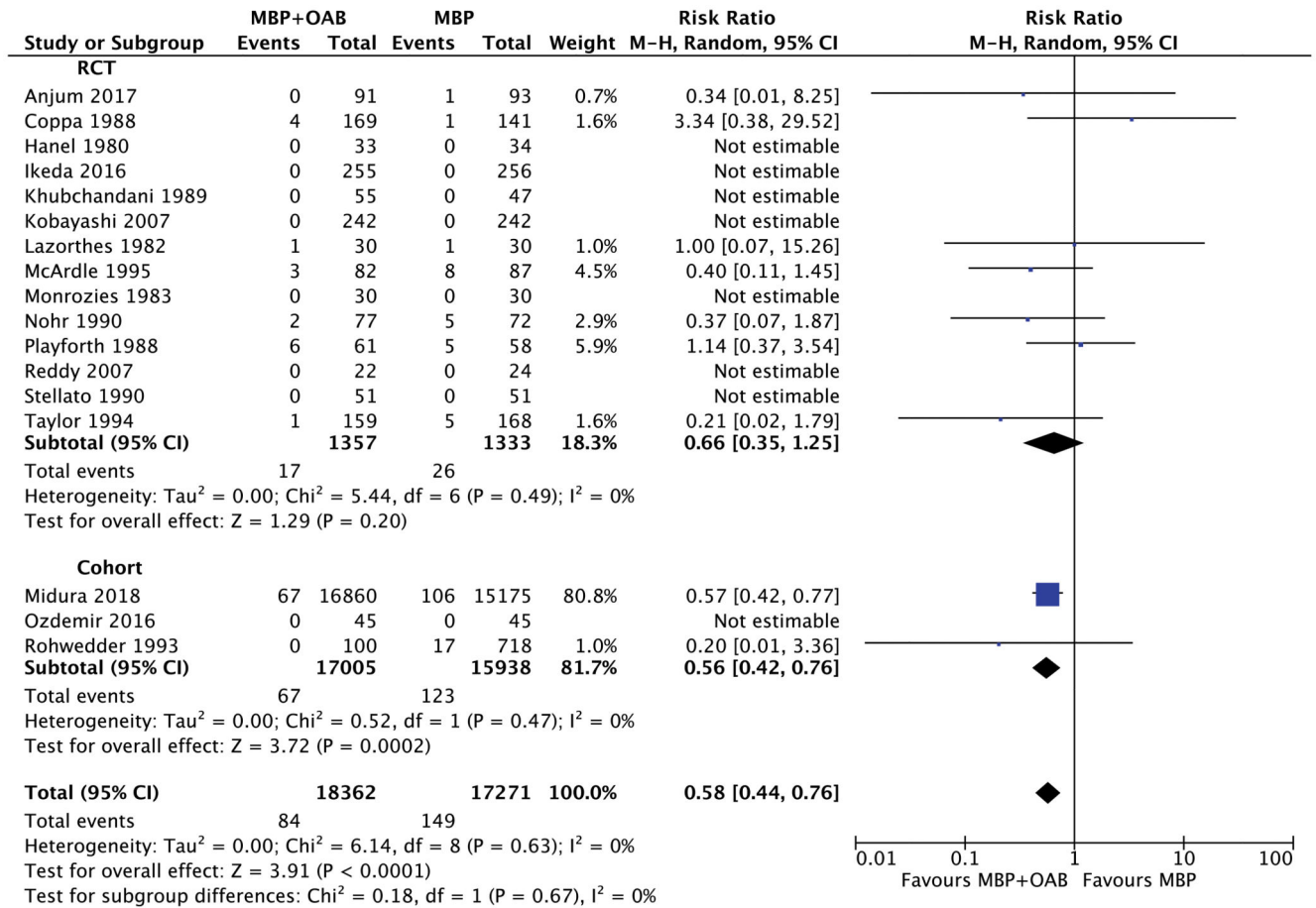


Figure 4. Forest plot comparing 30-day mortality rates for patients receiving MBP+OAB versus MBP alone, divided by evidence from RCTs and cohort studies. A Mantel-Haenszel random effects model was used to perform the meta-analysis and risk ratios are quoted including 95% confidence intervals.

Table 1
Risk of bias within randomized controlled trials included within the meta-analysis

Reference	Random sequence generation	Allocation concealment	Blinding of participants and personnel	Blinding of outcome assessment	Incomplete outcome data	Selective reporting	Other bias
Anjum <i>et al.</i> 201749	+	+	?	+	+	+	
Coppa <i>et al.</i> 199850	?	?	-	+	-	-	
Espin-Basany <i>et al.</i> 200551	?	?	?	+	+	+	
Hanel <i>et al.</i> 198052	-	?	?	+	-	+	
Hata <i>et al.</i> 201653	+	+	-	-	+	+	36 patients in the MBP+OAB group received reduced doses of kanamycin due to prescription error
Ikeda <i>et al.</i> 201655	+	+	-	+	+	+	
Ishida <i>et al.</i> 200156	+	-	-	-	+	?	
Kaiser <i>et al.</i> 198357	?	+	+	+	+	-	Different IV antibiotic regimens given to the two groups
Khubchandani <i>et al.</i> 198958	?	?	+	+	-	-	Different IV antibiotic regimens given to the two groups
Kobayashi <i>et al.</i> 200759	+	?	-	-	-	-	
Lau <i>et al.</i> 198861	+	?	?	?	+	+	
Lazorthes <i>et al.</i> 198262	?	?	?	?	?	-	Different IV antibiotic regimens given to the two groups
Lewis 200263	-	-	+	+	+	+	
McArdle <i>et al.</i> 199564	?	?	?	?	+	?	Different IV antibiotic regimens given to the two groups
Monrozies <i>et al.</i> 198565	?	?	?	?	+	+	Different IV antibiotic regimens

Reference	Random sequence generation	Allocation concealment	Blinding of participants and personnel	Blinding of outcome assessment	Incomplete outcome data	Selective reporting	Other bias
Nohr <i>et al.</i> 199066	?	?	+	+	-	+	given to the two groups Different IV antibiotic regimens given to the two groups
Oshima <i>et al.</i> 201367	?	?	-	-	+	+	
Peruzzo <i>et al.</i> 198769	?	?	?	?	+	+	
Playforth <i>et al.</i> 198870	?	?	?	?	+	+	
Ram <i>et al.</i> 200571	-	?	?	?	+	+	
Reddy <i>et al.</i> 200772	+	+	-	-	+	+	Group also randomized to probiotics – not included within meta-analysis
Reynolds <i>et al.</i> 198973	+	-	?	?	-	-	Two different IV antibiotic regimens in the MBP group
Sadahiro <i>et al.</i> 201475	+	-	+	+	?	?	Group also randomized to probiotics – not included within meta-analysis
Stellato <i>et al.</i> 199076	+	?	+	+	-	+	
Takesue <i>et al.</i> 200078	?	?	?	?	-	?	
Taylor <i>et al.</i> 199479	?	?	-	-	-	+	
Uchino <i>et al.</i> 201780	+	+	-	+	-	?	<i>C. difficile</i> toxin and faecal cultures only pre-op
Zmora <i>et al.</i> 200383	+	+	?	?	-	+	

+ Low risk of bias; - High risk of bias; ? Unclear risk of bias

Table 2

Patient demographics in studies included

Reference	Study methodology	Number of patients	Indication for surgery	Location of resection	Laparoscopic or open	OAB agent	MBP agent	Parenteral agent	Comparison included
Ajum <i>et al.</i> 201749	RCT	190	Gastrointestinal tract fistula IBD Trauma Malignancy	Partial small bowel resection – 39 Right colectomy – 67 Left colectomy – 50 LAR – 34	Laparoscopic – 40 Open – 150	Metronidazole 400 mg and levofloxacin 200 mg TDS on the day before surgery.	Sodium phosphate 133 ml twice a day on the day before surgery.	Second generation cephalosporin + metronidazole 30-60 min pre-incision, every 3 h intra-op then 24 h post-op.	MBP+OAB vs. MBP
Cannon <i>et al.</i> 201245	Retrospective database study – Veterans Affairs Surgical Quality Improvement Program	9940	Neoplasm – 7871 IBD – 176 Diverticulitis – 644 Not stated – 1248	Ileocolic resection – 984 Partial colectomy – 6847 Rectal resection – 1771 Total colectomy – 338	Not stated	Erythromycin, neomycin or metronidazole.	Polyethylene glycol, phospho-soda or magnesium citrate.	Not stated	MBP+OAB vs. MBP MBP+OAB vs. OAB MBP+OAB vs. no prep OAB vs. no prep OAB vs. MBP
Coppa <i>et al.</i> 198850	RCT	350	Cancer – 255 Inflammatory – 46 Other – 9	Not stated	All open	Neomycin 8 g/day and erythromycin 4 g/day in divided doses for 24 h pre-op.	Fleet phospho-soda between 1 and 3 days pre-op, and saline enemas for the last two days.	Cefoxitin 1-2 g according to patient body weight given preoperatively, intraoperatively and every 6 h for the first post-op day.	MBP+OAB vs. MBP
Englesbe <i>et al.</i> 201043	Retrospective propensity-matched database study – Michigan Surgical Quality Collaborative – Colectomy Best Practices Project	740	Not stated	Segmental colectomy Ileocolic resection	Open and laparoscopic	Neomycin and erythromycin 76.3% Neomycin alone 7.9% Erythromycin alone 2.6% Metronidazole alone 2.6% Clindamycin alone 2.6%	Polyethylene glycol 20.9% Phospho-soda 5.9% Fleet enema 38.5% Magnesium citrate 5% Other 29.7%	Not stated	MBP+OAB vs. MBP
Espin-Basany <i>et al.</i> 200551	RCT	300	Cancer – 269 IBD – 4 Diverticular disease – 21 Not stated – 6	Segmental resection – 120 Sigmoidectomy – 69 Anterior resection – 27 TME-colectomy – 66 APR – 18	Not stated	Neomycin 1 g and metronidazole 1 g EITHER TDS the day before surgery OR OD the day before surgery.	Sodium phosphate 45 ml diluted in 90 ml water BD the day before surgery	Cefoxitin 1 g pre-incision and two doses at 8 and 16 h post-op.	MBP+OAB vs. MBP
Hanel <i>et al.</i> 198052	RCT	77	Adenoma – 2 Carcinoma – 48 IBD – 4 Diverticular disease – 7 Hodgkin's disease – 1 Villus papilloma – 1 Cecal volvulus – 2 Sigmoid volvulus – 2	Right colectomy – 15 Left colectomy – 6 Transverse colectomy – 2 Sigmoid colectomy – 10 Colonic bypass – 1 Cecostomy – 1 Colostomy – 1 Colostomy closure – 5 Colotomy and polypectomy – 2 Anterior resection – 14	All open	Metronidazole 1 g QDS for four days and neomycin 1 g TDS for two days prior to surgery.	Four day standard mechanical preparation including a low residue diet, and alternating enemas or washouts.	Clindamycin 7 mg/kg and cephalosolin sodium 1 g given at the start of the anesthetic.	MBP+OAB vs. MBP

Reference	Study methodology	Number of patients	Indication for surgery	Location of resection	Laparoscopic or open	OAB agent	MBP agent	Parenteral agent	Comparison included
Hata <i>et al.</i> 201653	RCT	579	Colorectal malignancy Adenoma	APR - 7 Proctocolectomy - 2 Colectomy - 376 Anterior resection - 183 APR - 20	All laparoscopic	Kanamycin 1 g and metronidazole 750 mg BD at 13 h and 9 h pre-op.	Sodium picosulphate 75 mg and magnesium citrate 34 g with 180 ml water the day before surgery.	Cefmetazole 1 g 30 min pre-incision then every 3 h intra-op.	MBP+OAB vs. MBP
Ichimanda <i>et al.</i> 201754	Retrospective case controlled series	344	All colorectal cancer	Not stated. Primary site: Colon - 181 Rectum - 163	Laparoscopic - 293 Open - 51	Kanamycin 1 g TDS and metronidazole 1 g TDS for 24 h prior to surgery.	Polyethylene glycol 2 L and nasoside (Pulsemid) 24 mg.	Second generation cephem on the day of surgery until the second post-op day.	MBP+OAB vs. MBP
Ikeda <i>et al.</i> 201655	RCT	511	Colorectal malignancy	Colonic surgery - 309 Anterior resection - 177 APR - 25	All laparoscopic	Kanamycin 1 g and metronidazole 750 mg BD the day before surgery.	Magnesium citrate and sodium picosulphate the day before surgery.	Cefmetazole 1 g at least 30 min pre-incision, every 3 h intra-op and for 24 h post-op.	MBP+OAB vs. MBP
Ishida <i>et al.</i> 200156	RCT	143	Cancer - 135 IBD - 4 Diverticular disease - 1 Not stated - 3	Colectomy - 76 Anterior resection - 47 APR - 9 Total proctectomy with J pouch - 3 Total pelvic exenteration - 4 Other - 4	Not stated	Kanamycin 2 g/day and erythromycin 1.6 g/day in 4 divided doses from 2 days prior to surgery.	Polyethylene glycol 2 L given the day before surgery.	Cefotiam 1 g after induction, 1 g at one hour after completion of surgery and 4 additional doses given BD for 2 consecutive days.	MBP+OAB vs. MBP
Kaiser <i>et al.</i> 198357	RCT	119	Local malignancy - 50 Metastatic malignancy - 30 Diverticulitis - 17 Polyps - 9 IBD - 9 Not stated - 4	Right colectomy - 34 Left colectomy - 25 Sigmoid resection - 25 APR - 11 Anterior resection - 7 Subtotal colectomy - 6 Operative colotomy - 6 Total colectomy - 3 Colostomy closure - 2	All open	Neomycin 1 g TDS and erythromycin 1 g TDS the day prior to surgery.	Magnesium citrate and cleansing enemas for 2 days prior to surgery.	Cefoxitin 2 g with the 'on call' medications, 1 g intra-operatively and 1 g every 6 h following surgery for four doses in the MBP alone group. Cefazolin 1 g with the 'on call' medications, 500 mg intra-operatively and 1 g every 6 h following surgery for four doses in the MBP+OAB group.	MBP+OAB vs. MBP
Khubchandani <i>et al.</i> 198958	RCT	155	'Colonic surgery'	Not stated	All open	Neomycin 1 g and erythromycin 1 g at 1 pm, 2 pm and 10 pm the day before surgery.	Castor oil 60 ml the afternoon of admission and saline enemas the night of	Metronidazole 1 g given 1 h before surgery, then 500 mg at 6 and 12 h post-op in MBP alone group.	MBP+OAB vs. MBP

Reference	Study methodology	Number of patients	Indication for surgery	Location of resection	Laparoscopic or open	OAB agent	MBP agent	Parenteral agent	Comparison included
Kim <i>et al.</i> 201441	Retrospective propensity-matched database study – Michigan Surgical Quality Collaborative – Colectomy Best Practices Project	1914	Not stated	Ileocolic resection with anastomosis Segmental colectomy with anastomosis	Open – 1049 Laparoscopic – 865	Not stated	Not stated	Not stated	MBP+OAB vs. no prep
Kobayashi <i>et al.</i> 200759	RCT	484	Colorectal malignancy	Surgical procedure: Colon – 241 Rectum – 243	Not stated	Kanamycin 1 g and erythromycin 400 mg TDS the day before surgery.	Polyethylene glycol 2 L the morning of the day before surgery.	Cefmetazole 1 g at induction, an additional dose if operation exceeded 3 h, then BD for 3 days post-op.	MBP+OAB vs. MBP
Konishi <i>et al.</i> 200660	Retrospective case controlled series – National Nosocomial Infection Surveillance program	556	Not stated	Right colectomy – 94 Left colectomy – 155 Other colectomy – 90 LAR – 126 APR – 51 Total colectomy or panproctocolectomy – 34 Hartmann's procedure – 6 Additional concomitant procedures: Ostomy closure – 47 Ostomy formation – 106 Multiple organ resection – 93	Open – 515 Laparoscopic – 41	Kanamycin and metronidazole.	Oral laxative and glycerine enema.	Second generation cephalosporin given 30 min prior to incision, repeated every 3 h intra-op and stopped within 24 h after the operation.	MBP+OAB vs. MBP
Lau <i>et al.</i> 198861	RCT	194	All cancer	Right colectomy – 39 Left colectomy – 7 Transverse colectomy – 9 Sigmoid colectomy – 22 Subtotal colectomy – 10 Pelvic exenteration – 2 Palliative bypass – 4 Anterior resection – 39 LAR – 17 APR – 45	All open	Neomycin 1 g and erythromycin 1 g at 1 pm, 2 pm and 11 pm the day prior to surgery.	3 days of oral bisacodyl, magnesium sulphate and saline enemas prior to surgery.	Metronidazole 500 mg and gentamycin 2mg/kg body weight given 30 min prior to surgery, then repeated at 8 h intervals for two further doses.	MBP+OAB vs. MBP
Lazorthes <i>et al.</i> 198262	RCT	90	Cancer – 51 Colostomy closure – 23 Benign disease – 16	Colectomy – 30 APR – 9 Sphincter-saving resection – 23 Miscellaneous – 28	All open	Kanamycin 1 g QDS and metronidazole 250 mg QDS for 3 days prior to surgery.	Three days of low residue diet, enemas and magnesium sulphate purges.	Cephadrine 2 g at induction with metronidazole 500 mg infusion over 4 h in MBP alone group. Cephadrine 2 g and gentamycin 2 mg/kg as IM injection at time of premedication	MBP+OAB vs. MBP

Reference	Study methodology	Number of patients	Indication for surgery	Location of resection	Laparoscopic or open	OAB agent	MBP agent	Parenteral agent in the MBP+OAB group.	Comparison included
Lewis 200263	RCT	208	Cancer – 150 IBD – 51 Rectal prolapse – 10 Not stated – 2 (5 patients withdrawn)	Anterior resection – 119 APR – 19 Right colectomy – 55 Left colectomy – 13 Transverse colectomy – 4	Not stated	Neomycin 2 g and metronidazole 2 g BD the day before surgery.	Sodium phosphate the day before surgery, with saline enemas if this did not result in a clear effluent.	Amikacin 1 g and metronidazole 1 g on the day of surgery. MBP+OAB vs. MBP	
McArdle <i>et al.</i> 199564	RCT	169	Cancer/cancer related – 151 IBD – 13 Diverticular disease – 5	Right colectomy – 35 Left colectomy – 26 Anterior resection – 24 APR – 17 Total colectomy – 5 Hartmann's procedure/reversal – 15 Bypass – 7 Small bowel resection – 14 Formation or revision of stoma – 18 Others – 8	All open	Ciprofloxacin 1 g 1 h prior to surgery – one group received no further doses and one group received ciprofloxacin 750 mg BD for 3 days.	MBP alone: Gentamycin 120 mg + metronidazole 500 mg at induction then one group received gentamycin 80 mg + metronidazole 500 mg at 8 and 16 h post-op and one group received gentamycin 80 mg + metronidazole 500 mg TDS for 3 days. MBP+OAB: metronidazole 500 mg at induction then in one group at 8 and 16 h post-op and in the other metronidazole 500 mg TDS for 3 days.	MBP+OAB vs. MBP	
Midura <i>et al.</i> 201831	Database study – ACS NSQIP	45,724	IBD Cancer Diverticulitis Others	Left colectomy Right colectomy Segmental colectomy	Open Laparoscopic Robotic	Not stated	Not stated	Not stated MBP+OAB vs. MBP MBP+OAB vs. OAB prep MBP vs. OAB OAB vs. no prep	
Mik <i>et al.</i> 201646	Retrospective cohort study	2240	Colorectal malignancy	Right colectomy – 413 Left colectomy – 171 Sigmoidectomy – 282 Hartmann's – 171 Anterior resection – 309 LAR – 381 APR – 163 Not stated – 350	All open	Erythromycin 500 mg and neomycin 500 mg TDS the day before surgery.	Oral macrogol the day before surgery.	Cefazolin 1 g and metronidazole 500 mg directly before incision, and broadened to 3 doses if surgery lasted longer than 3 h.	MBP+OAB vs. no prep
Monrozieres <i>et al.</i> 198365	RCT	60	Cancer – 34 Closure of colostomy – 8 Benign – 18	Colectomy – 35 Rectal surgery – 15 Others – 10	All open	Kanamycin 1 g QDS and metronidazole 1 g QDS for 3 days pre-op.	Magnesium sulphate and enemas.	MBP+OAB: Cephadrine 2 g at induction and IM gentamycin 2 mg/kg at premedication	

Reference	Study methodology	Number of patients	Indication for surgery	Location of resection	Laparoscopic or open	OAB agent	MBP agent	Parenteral agent	Comparison included
Nohr <i>et al.</i> 199066	RCT	149	Cancer – 116 Complicated diverticulitis – 9 Crohn's disease – 8 UC – 1 Not stated – 15	Right colectomy – 29 Rectal resection – 44 Sigmoid resection – 30 APR – 19 Others – 27	All open	Bacitracin 250 mg and neomycin 250 mg TDS for 2 days pre-op. Metronidazole 500 mg TDS the day before surgery.	Frangula bark 2 tablets 2 days pre-op and magnesium sulphate (7.5 g) daily for 2 days pre-op.	Ampicillin 1 g within 1 h pre-op in MBP +OAB group. Fosfomycin 8 g and metronidazole 1 g within 1 h pre-op in MBP alone group.	MBP+OAB vs MBP
Oshima <i>et al.</i> 201367	RCT	200	Ulcerative colitis	Restorative proctocolectomy with ileal pouch-anal anastomosis (IPAA)	All open	Kanamycin 500 mg and metronidazole 500 mg TDS the day before surgery.	Magnesium citrate 1.8 L the day before surgery.	Flomoxef 30 min before surgery, repeated every 3 h intra-op and then 24 h post-op.	MBP+OAB vs. MBP
Ozdemir <i>et al.</i> 201668	Retrospective cohort study	90	Colonic malignancy Ulcerative colitis	Right colectomy – 17 Left colectomy – 10 Transverse colectomy – 9 LAR – 45 Total colectomy – 8 Other – 1	All open	Gentamycin 240 ml and metronidazole 2 g at 11 and 9 h pre-op.	Sodium dibasic phosphate 45 ml BD at 12 and 10 h pre-op, fleet enema 8 and 3-4 h pre-op.	Cefazolin 1 g and metronidazole 500 mg during anesthetic induction, continued BD for 5 days post-op.	MBP+OAB vs. MBP
Peruzzo <i>et al.</i> 198769	RCT	80	Cancer – 61 Diverticular disease – 6 Colostomy – 12 Not stated – 1	Right colectomy – 17 Left colectomy – 27 Sigmoid colectomy – 9 Anterior resection – 13 APR – 2 Colostomy closure – 12	All open	Neomycin 1 g at 19, 18, and 9 h pre-op and 2 g oral tinidazole.	'According to standard practice'.	Cefoxitin 30 min pre-op then at 6 and 12 h post-op.	MBP+OAB vs. MBP
Playforth <i>et al.</i> 198870	RCT	119 + 83 non randomized cohort (not included)	Cancer (curative) - 66 Cancer (palliative) - 22 Inflammatory - 31	Right colon - 38 Left colon and rectum - 81	All open	Neomycin 1 g every 6 h and metronidazole 200 mg every 8 h for 24 h prior to surgery.	Mannitol 100 g in 1 L water the day before surgery.	Metronidazole 500 mg at the time of premedication.	MBP+OAB vs. MBP
Ram <i>et al.</i> 200571	RCT	329	Cancer – 268 Benign - 61	Right colectomy – 42 Left colectomy – 74 Sigmoidectomy – 86 Subtotal colectomy – 11 APR – 34 Transverse colectomy – 3 Anterior resection – 50 LAR – 29	All open	Not stated	Monobasic sodium phosphate 2.4 g and dibasic sodium phosphate 0.9 g given the day before surgery.	Metronidazole 500 mg and ceftriaxone 1 g given 1 h pre-induction and continued for 48 h post-op.	MBP+OAB vs. OAB
Reddy <i>et al.</i> 200772	RCT	92 (46 pertinent to this meta-analysis)	Cancer and benign	Right colectomy - 16 Left colectomy - 6	All open	3 g neomycin in three divided doses the day before surgery.	Sodium picosulphate	Not stated	MBP+OAB vs. MBP

Reference	Study methodology	Number of patients	Indication for surgery	Location of resection	Laparoscopic or open	OAB agent	MBP agent	Parenteral agent	Comparison included
Reynolds <i>et al.</i> 198973	RCT	330	Cancer – 247 Benign – 5 Inflammatory lesion – 19 Others – 59	Anterior resection - 18 APR - 3 Subtotal colectomy - 2 Panproctocolectomy - 1 Right colectomy – 65 Left colectomy – 9 Sigmoid colectomy – 48 APR – 50 Anterior resection – 97 Panproctocolectomy – 2 Subtotal colectomy – 9 Hartmann's procedure – 10 Colostomy surgery – 35 Other – 5	All open	Metronidazole 400 mg eight hourly and neomycin 1 g six hourly for 48 h prior to surgery. Last dose of antibiotics given 8 and 12 h prior to surgery, respectively.	Magnesium sulphate up to 8x4 g doses for 48 h starting 72 h pre-op. Followed by two doses of sodium picosulphate the day before surgery.	Either piperacillin 2 g IV at induction and 3 further doses 8 hourly or metronidazole 500 mg and cefuroxime 1.5 g at induction followed by 3 further doses of metronidazole and 2 further doses of cefuroxime.	MBP+OAB vs. MBP
Rohwedder <i>et al.</i> 199374	Retrospective historical case controlled series	818 (100 MBP+OAB, 718 MBP)	Of those with MBP+OAB: Colorectal cancer – 89 Anal cancer – 1 Pelvic recurrence – 1 Villous tumour – 1 Diverticular disease – 6 UC – 1 Crohn's colitis – 1	Of those with MBP+OAB: Right colectomy – 14 Left colectomy – 25 LAR – 37 Miles APR – 12 Total colectomy – 6 Subtotal colectomy – 1 Double colectomy – 2 Other – 3	All open	Ciprofloxacin 750 mg taken between 1 and 3 h pre-op.	Polyethylene glycol the day before surgery.	Gentamycin 80 mg and metronidazole 500 mg at the beginning of induction, then gentamycin 80 mg every 8 h for 3 days.	MBP+OAB vs. MBP
Sadahiho <i>et al.</i> 201475	RCT	294	Colorectal malignancy	Not stated – tumour location: Right colon – 99 Transverse colon – 38 Left colon – 157	Open – 214 Laparoscopic – 80	Kanamycin sulphate 500 mg + metronidazole 500 mg TDS the day before surgery.	Sodium picosulphate 10 ml 2 days pre-op and 2 L polyethylene glycol the day before surgery.	Flomoxef 1 g 1 h pre-incision and further dose given if operative duration exceeded 3 h.	MBP+OAB vs. MBP
Stelato <i>et al.</i> 199076	RCT	146	Cancer – 123 Polyp – 11 Diverticular disease – 6 IBD – 6	Right colectomy - 44 Left colectomy - 17 Transverse colectomy - 4 Sigmoid colectomy - 30 LAR - 31 APR - 15 Subtotal colectomy - 5	All open	Neomycin 1g and erythromycin 1g TDS on the day before surgery.	Magnesium citrate 1.745 g in 296 ml in the morning and an enema (19 g sodium biphosphate and 7 g sodium phosphate in 118 ml) in the evening 2 days prior to surgery. Magnesium citrate 1.745 g in 296 ml in the morning and saline enemas until clear in the evening of	Cefoxitin 2 g at induction then at 6 and 12 h following the first dose.	MBP+OAB vs. MBP

Reference	Study methodology	Number of patients	Indication for surgery	Location of resection	Laparoscopic or open	OAB agent	MBP agent	Parenteral agent	Comparison included
Sun <i>et al.</i> 201877	Retrospective case controlled series	321	Malignancy – 306 Benign – 12 IBD – 3	Right colectomy – 86 Left colectomy – 24 Sigmoid colectomy – 65 LAR – 90 APR – 16 Laparoscopic anterior resection – 12 Laparoscopic sigmoidectomy – 15 Subtotal colectomy – 4 Laparoscopic right hemicolectomy – 8	Laparoscopic – 35 Open – 269	Neomycin 1 g and erythromycin 1 g at 20, 19 and 10 h prior to surgery.	Fleet phospho-soda 45 ml at 24 and 15 h before surgery then tap water enema at 2 h pre-op.	Cefazolin 1 g at induction.	MBP+OAB vs. MBP
Takesue <i>et al.</i> 200078	RCT	83	Dukes A – 16 Dukes B – 43 Dukes C – 24	Ileocecal resection – 5 Right colectomy – 14 Left colectomy – 3 Transverse colectomy – 6 Sigmoidectomy – 24 LAR – 24 Miles' APR – 7	All open	Kanamycin 500 mg and metronidazole 500 mg at 2 pm, 3 pm and 11 pm the day before surgery.	Polyethylene glycol commence at 10 am the day before surgery.	Cefmetazole 1 g given at induction, then TDS for 3 days following surgery.	MBP+OAB vs. MBP
Taylor <i>et al.</i> 199479	RCT	327	Benign – 53 Cancer – 259 IBD – 15	Anastomosis right colon – 93 Anastomosis left colon/rectum – 168 Hartmann's resection – 6 APR – 43 Not stated – 17	Not stated	Ciprofloxacin 500 mg BD the day before surgery.	Sodium picosulphate one sachet BD the day before surgery.	Piperacillin 4 g at induction of anaesthesia.	MBP+OAB vs. MBP
Uchino <i>et al.</i> 201780	RCT	325	Crohn's disease	Small bowel resection Colonic resection Rectal resection	All open	Kanamycin 500 mg and metronidazole 500 mg TDS the day before surgery.	Sodium picosulphate hydrate (20 ml of 0.75%) pre-operatively.	Flomoxef sodium 30 min before surgery, every 3 h intra-op then 24 h post-op.	MBP+OAB vs. MBP
Vo <i>et al.</i> 201881	Retrospective case control series	89	Colorectal cancer	Left colectomy - 14 Sigmoid colectomy – 16 LAR – 35 APR – 14 Subtotal colectomy or other – 10	Open – 21 Minimally invasive – 68	Neomycin sulphate 1 g and metronidazole hydrochloride 1 g TDS. Commenced one day prior to surgery.	Magnesium citrate 296 ml twice daily. Commenced one day prior to surgery.	Ertapenem – 82 Non-ertapenem - 7	MBP+OAB vs. MBP
Wren <i>et al.</i> 200582	Retrospective case controlled study	304	Not stated	Colon and/or rectal resection – 258 Colostomy creation or take down – 46	Open and laparoscopic	Neomycin 1 g and erythromycin 1 g	GoLYTELY, magnesium citrate or Fleet phospho-soda	Cephalosporin and metronidazole 59.2% Second generation cephalosporin 21.0% Fluoroquinolone and metronidazole or clindamycin 9.5% First-generation cephalosporin alone 3.9%	MBP+OAB vs. MBP

Reference	Study methodology	Number of patients	Indication for surgery	Location of resection	Laparoscopic or open	OAB agent	MBP agent	Parenteral agent	Comparison included
Zmora <i>et al.</i> 200383	RCT	380	Cancer – 296 Diverticular disease – 16 Hartmann's procedure (for closure) – 29 Benign polyp – 14 IBD – 13 Not stated – 12	Right colectomy – 113 Left colectomy – 33 Sigmoidectomy – 89 Anterior resection – 83 Closure of Hartmann's – 29 Subtotal/total abdominal colectomy – 24 Total proctectomy and ileal pouch – 9	Not stated	Neomycin and erythromycin	Polyethylene glycol 1 gallon 12 to 16 h pre-op. Rectal surgery – given Fleet enema.	Extended-spectrum penicillin 3.6% 'Broad spectrum antibiotics' continued for 24 h post-op.	MBP+OAB vs. OAB

APR – abdominoperineal resection; IBD – inflammatory bowel disease; LAR – low anterior resection; MBP – mechanical bowel preparation; OAB – oral antibiotics; RCT – randomized controlled trial

Table 3

Overall summary of results

Preparation considered	Outcome measure	All studies	RCTs only	Cohort studies only	
MBP+OAB vs. MBP	Surgical site infection	Significant ↓ with MBP+OAB (RR 0.51, 95% CI 0.46 to 0.56, p<0.00001, I ² =13%)	Significant ↓ with MBP+OAB (RR 0.57, 95% CI 0.48 to 0.68, p<0.00001, I ² =12%)	Significant ↓ with MBP+OAB (RR 0.48, 95% CI 0.44 to 0.51, p<0.00001, I ² =0%)	
	Anastomotic Leak	Significant ↓ with MBP+OAB (RR 0.62, 95% CI 0.55 to 0.70, p<0.00001, I ² =0%)	No difference (RR 0.69, 95% CI 0.43 to 1.11, p=0.13, I ² =0%)	Significant ↓ with MBP+OAB (RR 0.45, 95% CI 0.25 to 0.80, p=0.0007, I ² =22%)	
	30-day mortality	Significant ↓ with MBP+OAB (RR 0.58, 95% CI 0.44 to 0.76, p<0.00001, I ² =0%)	No difference (RR 0.66, 95% CI 0.35 to 1.25, p=0.20, I ² =0%)	Significant ↓ with MBP+OAB (RR 0.56, 95% CI 0.42 to 0.76, p=0.0002, I ² =0%)	
	Overall morbidity	Significant ↓ with MBP+OAB (RR 0.67, 95% CI 0.63 to 0.71, p<0.00001, I ² =0%)	No difference (RR 0.71, 95% CI 0.41 to 1.24, p=0.23, I ² =9%)	Significant ↓ with MBP+OAB (RR 0.67, 95% CI 0.63 to 0.71, p<0.00001, I ² =0%)	
	Development of ileus	Significant ↓ with MBP+OAB (RR 0.72, 95% CI 0.52 to 0.98, p=0.04, I ² =36%)	No difference (RR 0.62, 95% CI 0.14 to 2.67, p=0.52, I ² =50%)	No difference (RR 0.68, 95% CI 0.45 to 1.03, p=0.07, I ² =53%)	
	<i>C. difficile</i> infection	No difference (RR 0.94, 95% CI 0.55 to 1.61, p=0.81, I ² =37%)	No difference (RR 0.79, 95% CI 0.21 to 2.96, p=0.72, I ² =10%)	No difference (RR 0.97, 95% CI 0.54 to 1.75, p=0.92, I ² =64%)	
	MBP+OAB vs. OAB	Surgical site infection	No difference (RR 0.98, 95% CI 0.64 to 1.50, p=0.92, I ² =77%)	No difference (RR 1.36, 95% CI 0.78 to 2.35, p=0.28, I ² =0%)	No difference (RR 0.83, 95% CI 0.48 to 1.43, p=0.51, I ² =90%)
		Anastomotic Leak	No difference (RR 0.79, 95% CI 0.59 to 1.05, p=0.11, I ² =0%)	No difference (RR 1.39, 95% CI 0.47 to 4.10, p=0.55, I ² =0%)	---
		30-day mortality	Significant ↓ with MBP+OAB (RR 0.58, 95% CI 0.34 to 0.97, p=0.04, I ² =0%)	No difference (RR 1.02, 95% CI 0.30 to 3.50, p=0.97, I ² =0%)	---
		Overall morbidity	---	---	---
Development of ileus		Significant ↓ with MBP+OAB (RR 0.83, 95% CI 0.73 to 0.95, p=0.008, I ² =0%)	No difference (RR 1.25, 95% CI 0.68 to 2.33, p=0.47, I ² =0%)	---	
MBP+OAB vs. no preparation	<i>C. difficile</i> infection	---	---	---	
	Surgical site infection	---	---	Significant ↓ with MBP+OAB (RR 0.54, 95% CI 0.43 to 0.68, p<0.00001, I ² =82%)	
	Anastomotic Leak	---	---	Significant ↓ with MBP+OAB (RR 0.52, 95% CI 0.45 to 0.59, p<0.00001, I ² =0%)	

Preparation considered	Outcome measure	All studies	RCTs only	Cohort studies only
	30-day mortality	---	---	Significant ↓ with MBP+OAB (RR 0.36, 95% CI 0.17 to 0.76, p=0.008, I ² =46%)
	Overall morbidity	---	---	---
	Development of ileus	---	---	Significant ↓ with MBP+OAB (RR 0.72, 95% CI 0.68 to 0.77, p<0.00001, I ² =0%)
	<i>C. difficile</i> infection	---	---	---

MBP – mechanical bowel preparation; OAB – oral antibiotics; --- Insufficient data for conduct of meta-analysis

OAB vs. no preparation – only outcome was surgical site infection in cohort studies alone which demonstrated a significant ↓ with OAB. OAB vs. MBP – only outcome was surgical site infection in cohort studies alone which demonstrated no difference.