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Effect of preoperative oral antibiotics in combination with mechanical bowel preparation on inflammatory response and short-term outcomes following left-sided colonic and rectal resections

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Golder et al’s propensity-matched retrospective observational study found the addition of preoperative oral antibiotics and mechanical bowel preparation to prophylactic intravenous antibiotics was associated with reduced postoperative complications and severity of systemic inflammatory response. We believe several statistical issues may have led to the treatment effect being overstated.

First, more patients in the control group had malignant disease; this can lead to a heightened inflammatory state. The P values that would have made this obvious were removed between Table 1 and Table 2. Second, the historical nature of the control group, with unmeasured aspects of treatments likely to improve over time, will lead to improved outcomes in the treatment group. Taken together, these two effects could lead to false rejection of the null hypothesis.

Third, the postoperative Glasgow Prognostic Score (poGPS) is determined solely by albumin and C-reactive protein (CRP) concentrations. Determining statistical differences in poGPS at the same time as differences in CRP and albumin, at exactly the same cut-off values, artificially inflates the number of hypotheses being tested. In addition, no Bonferroni correction for multiple hypothesis testing was undertaken for the 15 P values given in Table 3; again, this could lead to a type I error and is an example of P-hacking.

We feel these points have led to Golder and colleagues overstating the likelihood of any treatment effect. Gut decontamination in colorectal surgery is an important issue that deserves a large high-quality RCT.

Disclosure

The authors declare no conflict of interest.

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1 Golder AM, Steele CW, Conn D, MacKay GJ, McMillan DC, Horgan P et al. Effect of preoperative oral antibiotics in combination with mechanical bowel preparation on inflammatory response and short-term outcomes following left-sided colonic and rectal resections. BJS 2019; 3: 830–839.

2 Watt DG, McSorley ST, Park JH, Horgan PG, McMillan DC. A postoperative systemic inflammation score predicts short- and long-term outcomes in patients undergoing surgery for colorectal cancer. Ann Surg Oncol 2017; 24: 1100–1109.

Authors’ reply: Effect of preoperative oral antibiotics in combination with mechanical bowel preparation on inflammatory response and short-term outcomes following left-sided colonic and rectal resections

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We acknowledge Hartrick and colleagues for their interest in and comments regarding our study. They correctly note that our study involved two longitudinal groups (control group before 2015 and test group after 2015). This was stated clearly in the Methods section and acknowledged in the limitations paragraph of the Discussion. For several reasons, we do not believe this introduced a significant bias to the results. The proportion of these patients was small (less than 15 per cent) and, although the authors have raised concerns regarding the possible heightened inflammatory state in malignant compared with benign disease, we believe the propensity score-matched for the preoperative systemic inflammatory response (modified Glasgow Prognostic Score) with good balance between test and control groups (Cramer’s V = 0.018). By convention, P values for significance are not usually presented after matching, as the very fact that these variables have been used to generate the propensity scores leads to inherent bias and renders inference illogical. For this reason, Cramer’s V was calculated before and after matching (Tables 1 and 2 respectively), along with a ‘butterfly plot’ of propensity score distribution. Both methods showed improvement in balance after matching.

The authors correctly state that the C-reactive protein and albumin cut-offs for days 3 and 4 were the same as those used to calculate the postoperative Glasgow Prognostic Score on days 3 and 4. The outcomes included in Table 3 are largely related – both the postoperative inflammatory state on postoperative days 3 and 4 and the development of postoperative complications. As a result, the likelihood of a type I error is substantially less than it would have
been had the study reported 15 unrelated outcomes. Regardless of statistical values, absolute numbers/percentages are shown in Table 3 and are of clear clinical significance (a reduction in the overall complication rate from 55 to 28 per cent, a reduction in the infective complication rate from 37 to 20 per cent, and a reduction in the surgical-site infection rate from 23 to 10 per cent). This was, however, a relatively small study with approximately 100 patients in each group, and clearly not powered to detect significant differences in less frequently observed complications including deep surgical-site infections and anastomotic leaks.

As the above outcomes of interest are likely to be related, a Bonferroni correction is perhaps an overly conservative way of correcting for multiple testing. Given the interrelationship of our outcomes, a different analysis to correct for multiple testing such as the Benjamini–Hochberg procedure may be more appropriate 1. Indeed, we have now carried out such a post hoc analysis. Using this correction, all of the outcomes reported as statistically significant in Table 3 remained so when the false discovery rate was set at 5 per cent, and the majority remained statistically significant when the false discovery rate was set at 10 per cent. Therefore, the suggestion of ‘P-hacking’ is unlikely to be the case and is supportive of the peer review process.

Hartrick and colleagues state in their letter that the use of oral antibiotics and mechanical bowel preparation in resectional colorectal surgery is an important issue requiring further prospective research in the form of large prospective RCTs. As acknowledged in the final paragraph of the Discussion section of our article (‘This strategy is worthy of further investigation’), we are in clear agreement. Indeed, we look forward to the reporting of those trials currently underway, in particular the COLON-PREP trial (EudraCT no. 2017-002542-72). This is of particular interest given the recent negative findings of the MOBILE trial, contrary to most of the published meta-analyses 2–7 in the field.

Disclosure

The authors declare no conflict of interest.

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[Correction added on 17 April 2020, after first online publication: The article title was previously missing and has been inserted in this current version.]

Cluster-randomized crossover trial of chlorhexidine–alcohol versus iodine–alcohol for prevention of surgical-site infection (SKINFECT trial)

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We read with interest the work of Charehbili and colleagues 1, which ‘aimed to investigate whether there is a superiority of chlorhexidine–alcohol over iodine–alcohol for preventing SSI’.

This cluster-randomized crossover trial was conducted in five hospitals and 3665 patients were included. The authors found that the incidence of surgical-site infection (SSI) was not different between the groups: 3.8 per cent among patients in the chlorhexidine–alcohol group versus 4.0 per cent in those in the iodine–alcohol group (odds ratio 0.96, 95 per cent c.i. 0.69 to 1.35).

We commend the authors for performing this interesting study, as these results are useful for the choice of the most appropriate preoperative antiseptic. However, we have several statistical suggestions and queries that we would like to communicate to the authors.

The authors concluded that ‘Preoperative skin disinfection with chlorhexidine–alcohol is similar to that for iodine–alcohol with respect to reducing the risk of developing an SSI’. This may be due to an underpowered study.

In fact, sample size was estimated by simulation. Although this approach is