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**Quality of bowel preparation in patients with inflammatory bowel disease undergoing colonoscopy: What factors to consider?**

Gravina AG *et al*. Bowel preparation in IBD patients

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**Abstract**

An adequate bowel preparation in patients with inflammatory bowel disease (IBD) is a prerequisite for successful colonoscopy for screening, diagnosis, and surveillance. Several bowel preparation formulations are available, both high- and low-volume based on polyethylene glycol. Generally, low-volume formulations are also based on several compounds such as magnesium citrate preparations with sodium picosulphate, oral sulphate solution, and oral sodium phosphate-based solutions. Targeted studies on the quality of bowel preparation prior to colonoscopy in the IBD population are still required, with current evidence from existing studies being inconclusive. New frontiers are also moving towards the use of alternatives to anterograde ones, using preparations based on retrograde colonic lavage.

**Key Words:** Bowel preparation; Colonoscopy; Inflammatory bowel disease; Crohn’s disease; Ulcerative colitis; Artificial intelligence

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**Core Tip:** Obtaining adequate bowel preparation is challenging when treating a patient with inflammatory bowel disease (IBD) undergoing colonoscopy. Colonoscopy has multidimensional value ranging from diagnosis to disease surveillance and cancer screening. Although numerous data are available on bowel preparations in the general population, it is still unclear which preparation is the best for both efficacy and safety in patients with IBD. In addition, the factors that increase the risk of suboptimal preparation in IBD patients remain unclear.

**INTRODUCTION**

Adequate bowel preparation is crucial for successful colonoscopy in diagnostic, therapeutic, and screening indications; however, it remains one of the main challenges in patients undergoing colonoscopy[1,2]. Thus, several requirements were identified for managing this issue, including improving the palatability and tolerability of bowel preparation products and adopting a tailored, patient focused approach by taking into account the patient's choice of product[1]. Management of inflammatory bowel disease (IBD), including Crohn’s disease (CD) and ulcerative colitis (UC), requires regular endoscopic surveillance. In patients with IBD, endoscopic procedures are indicated for initial diagnosis, monitoring of disease activity, evaluation of therapeutic response, and cancer screening[3]. The latter indication is significant as patients with IBD are at an increased risk of developing colorectal cancer[4]. In addition, a proportion of patients with IBD undergo surgical intervention and therefore require postoperative endoscopic re-evaluation to identify any postsurgical recurrence, especially in CD[5].

Emerging evidence has emphasised “treat-to-target” approaches in the management of IBD by defining specific treatment goals and punctuating the timing of evaluation. Assessment of these uses standardised endoscopic scores, the calculation of which requires adequate bowel preparation in all regions of the colon and last ileum that can be explored by colonoscopy and ileocolonoscopy[6,7]. However, especially in IBD patients, there is a need to weigh up the safety of intestinal preparations, especially under severe disease activity conditions[8]. Although rare, significant adverse events associated with bowel preparation, such as mucosal inflammation[9], intestinal perforations[10], and ischaemic colitis[11], have been reported.

The purpose of the present review is to provide an overview of the current available evidence on bowel preparation formulations, specifically evaluated in IBD, to determine the factors of successful and unsuccessful bowel preparation in patients with IBD, and to provide clues on the appropriate choice of formulation for IBD patients.

**TYPE OF BOWEL PREPARATIONS STUDIED IN IBD**

Several bowel preparations are available for patients undergoing colonoscopies. They can be categorized into high-volume (volume of at least 3 L), isosmotic, polyethylene glycol (PEG) formulations, and low-volume (volume less than 3 L, but with the addition of osmotically active adjuvants such as ascorbic-acid, citrate, and bisacodyl) PEG[12]. There are also osmotically active, non-PEG, low-volume solutions. Examples include magnesium citrate preparations or other preparations based on sodium picosulphate, oral sulphate, and oral sodium phosphate[12]. However, they have a high potential risk of adverse events due to their osmotic properties.

There are currently no studies that provide a definitive and detailed overview of the application of various bowel preparations available to patients with IBD. Table 1 summarises the bowel preparation regimens studied specifically for IBD patients. Case reports of patients with IBD and the relevant adverse events associated with bowel preparation are shown in Table 2.

***Bowel preparation studied in IBD: Origins***

An early trial in 1982 focused on evaluating bowel preparation in patients with UC who underwent colonoscopy for dysplasia screening[13]. The study compared two bowel preparation products, castor oil (30 mL) and senna tablets (75 mg sennosides). However, the authors highlighted post-colonoscopy complaints (particularly complaints that were more common with senna tablets) and many UC flare-ups in several cases requiring steroid treatment. In this study, they targeted patients with UC with inactive or presumably mild disease activity. The efficacy rate shown with both preparations exceeded 80%. In any case, to date, such preparations are insupportable and fall into disuse, especially given the abundant availability of new generation products with better safety margins.

***High- and low-volume plus adjuvant preparations of polyethylene glycol in patients with IBD***

PEG-based preparations were first introduced in 1980 by Davis *et al*[14] as an alternative to traditional laxatives. Several studies have examined high-volume PEG preparations in patients with IBD. In general, these formulations have been studied alone or in comparison to a control group of other solutions such as low-volume solutions.

An early Italian study[15] evaluated the efficacy of simethicone in addition to a 4 L PEG Electrolyte Lavage Solution (ELS). A high efficacy rate of 96% was shown in the group taking only 4 L PEG-ELS (considering at least acceptable preparation). In contrast, preparation was adequate and excellent in 50% and 27% of IBD participants, respectively. The authors defined the preparation as excellent in the absence of formed stools and the presence of low fluid content. The preparation was defined as adequate in the absence of formed stools and the moderate presence of clear fluid. Regarding the safety profile, the preparation showed poor tolerability in only 14% of participants with no serious adverse events and the main complaint was abdominal bloating.

However, two subsequent studies compared high-volume and low-volume PEG solutions with bisacodyl and ascorbic acid adjuvants.

Bisacodyl is a diphenylmethane compound. It has an extrinsic laxative action owing to its dual prokinetic and secretion properties after conversion to an intestinally active metabolite. It has shown comparable properties in some intestinal motility parameters to other drugs such as prucalopride, linaclotide, and tegaserod[16].

In a noninferiority study with 211 participants, Kastenberg *et al*[17] compared a high-volume 4 L PEG-ELS solution with a low-volume solution (2 L PEG) with bisacodyl adjuvant at 5 mg, and concluded a noninferiority of the latter over the former. They focused exclusively on patients with UC. Patients had a free choice of a split or non-split regimen, but nonetheless were instructed to eat a low fibre diet for the 3 d prior to preparation. Bisacodyl was administered by patients in the afternoon of the day before the procedure, and only after the procedure was initiated. Colon cleansing was assessed using the Ottawa scale[17]. The efficacy rate was higher in the low-volume group (83%), compared with 77.1% in the high-volume group, but the difference was not significant. In addition, the presence of bubbles was significantly worse in the low-volume group. Disease activity and the type of administration (split or not) did not influence patient compliance. In terms of safety, severe adverse reactions were not observed in both solutions. The low-volume PEG preparation with bisacodyl in this study, showed greater potential in avoiding gastrointestinal disorders associated with bowel preparation, including bloating, cramping, anal irritation, nausea, and vomiting.

In any case, the use of bisacodyl at high dosages > 5 mg should be avoided[18]. Cases of ischaemic colitis after bisacodyl use have been reported. Two cases were reported after using 15 mg bisacodyl in two women over 50 years of age with ischaemic colitis. In both cases, the patients were discharged without adverse outcomes; however, in one of the two cases, haemodynamic deterioration was experienced[19]. They were not IBD patients; however, in the latter patients, bisacodyl dosing must be weighed carefully. Certainly, such preparations should be evaluated with extreme caution in elderly patients with cardiovascular and ischaemic colitis risk factors[20] and even more so in patients with coexisting IBD.

Neurointestinal reactions are further rare but serious adverse reactions. In some studies, the use of bisacodyl as a laxative induced changes in colonic redundancy and colonic dilatation with loss of haustral markings. This is likely to have been caused by neuronal or colonic muscle damage[21]. In addition, *in vitro* evidence in rat bladder epithelium, suggests that bisacodyl has the potential to induce proliferative epithelial injury[21]. However, such evidence is still experimental and does not definitively impose a contraindication for this product. Moreover, the contraindication for bisacodyl in patients with advanced congestive heart failure should also be considered[18].

A subsequent study by Kim *et al*[22] investigated the low-volume solution adjuvant ascorbic acid, and compared this with classic high-volume 4 L PEG-ELS. As in the previous study, the authors exclusively targeted 109 patients with inactive UC. In addition, in this study, patients were given a free choice between a split and non-split regimen, and were advised to eat a fibre-free diet for 2 d before the endoscopic examination. Bowel preparation quality was assessed with the Boston Bowel Preparation Scale (BBSP)[23] and a score of equal to or greater than 6 defined a successful preparation. The authors also assessed possible disease recurrence 4 wk after the endoscopic procedure using the Simple Clinical Colitis Activity Index[24], with a cut-off greater than 4 defined as recurrence. In both groups, the split regimen was the most widely used, and more than 90% of patients in both groups achieved an effective preparation. The safety profile was slightly worse in the 4 L PEG group, but not significantly different from that of the low-volume preparation. However, patients treated with 4 L PEG experienced a significantly higher rate of nausea. There was no difference in disease recurrence at 1 mo, which was more, but not significantly greater than that in the low-volume group with ascorbate (25% *vs* 22.6%, respectively).

Moreover, in the context of ascorbate-based preparations, Maida *et al*[25] evaluated a preparation with 1 L of PEG in 185 patients with IBD, compared with 226 non-IBD controls. The study concluded a higher clearance rate (assessed by the BBSP) than in non-IBD controls (92.9% *vs* 85.4%), and a similar safety profile between IBD and non-IBD. No correlation was found between disease activity and incidence of adverse events. The rate of non-severe adverse events was 22.2% in IBD patients compared with 21.2% in controls, and there were no severe adverse events in the study.

Patients with phenylketonuria or glucose-6-phosphate dehydrogenase deficiency should not undergo this type of preparation due to the presence of ascorbate[26]. Some case reports of serious adverse events have also been described with ascorbic acid. For example, an > 70-year-old hypertensive diabetic woman experienced ischaemic colitis with 200 g of 3350 PEG in 1 L and 21 g of ascorbic acid solution[27]. An 82-year-old woman with atrial fibrillation and mild chronic renal failure received a 1 L PEG preparation with ascorbate and experienced non-occlusive mesenteric ischaemic colitis[28]. Both were elderly, non-IBD patients, with cardiovascular risk factors. Therefore, similar to bisacodyl, such preparations should be weighed carefully in elderly patients with IBD and cardiovascular risk factors. In addition, attention should be paid to patients with advanced heart failure, unstable angina, and creatinine clearance < 30 mL/min. In these patients, adequate hydration should be considered when such a formulation is used[18].

Finally, an attempt with very low-volume PEG-based preparations (1 L PEG) was made by Neri *et al*[29]. In a cohort of 103 IBD patients, with good distribution between CD and UC, Neri *et al*[29] showed an adequate preparation rate of 85.4% (assessed by BBSP), with an impact of disease activity on preparation rate. This evidence, coupled with a good safety profile, makes this preparation a good choice for patients with a very low tolerance for high-volume preparations.

Moreover, a final study by Mohsen *et al*[30] evaluated a comparative single-blinded randomised trial for both preparation type and population (IBD *vs* non-IBD patients). The authors compared a preparation based on sodium picosulphate/magnesium citrate in combination with PEG and a low-volume preparation of PEG with ascorbic acid. The two preparations did not differ in terms of efficacy (assessed by the Ottawa scale) or safety. Although, in the group of IBD patients on the low-volume preparation of PEG with ascorbic acid, the incidence of abdominal pain increased. Overall, only 10.5% of the patients had inadequate preparation.

***What are the alternatives to polyethylene glycol-based preparations in focused studies in patients with IBD?***

Alternatives to PEG-based preparations include those based on magnesium citrate plus picosulphate and oral sulphate solutions, a compound of magnesium sulphate, sodium sulphate, and potassium sulphate. However, few studies have evaluated these preparations for IBD treatment.

A 2019 French multicentre study examined several low-volume solutions based on sodium picosulphate, sodium phosphate, and trisulphate (sodium, magnesium, and potassium sulphate)[31]. All three of these non-PEG regimens had been administered in more than 50% of the sample in a split regimen, with a previous low fibre diet lasting between one and more than 3 d, depending on the regimen. The efficacy rate in the preparation with sodium picosulphate reached 78.4%, comparable with 76.7% of the 2 L PEG examined in the same study. Both formulations were significantly more effective than the control 4 L PEG group. In addition, patient tolerance was better in the picosulphate group than in the 2 L and 4 L PEG groups. The safety profile of sodium picosulphate was not extremely harmful because there was one case of fever and another of vomiting in CD patients (however, the sample receiving this formulation consisted of only approximately 80 patients). Oral sulphate tablets were evaluated in the study by Kim *et al*[32], however, only in 110 patients with inactive IBD in non-inferiority comparison with split regimens with a 2 L PEG preparation with ascorbic acid. The study showed a greater tolerance for the oral sulphate-based preparation, so much so that more than 90% of the patients taking it stated that they would reuse it for subsequent colonoscopy.

Regarding safety, although there were no severe events, a small percentage of patients (only two) on oral sulphate showed disease flare-ups after colonoscopy. Furthermore, caecal intubation was achieved more quickly in the PEG-based preparation (100%) than in the sulphate preparation (92.8%). Finally, the preparation with sulphate achieved a lower and, therefore, better score for bubble presence. There is still little evidence to justify the use of such alternative preparations to PEG-based preparations in patients with IBD, so much so that the recommendations of endoscopic reference societies tend to recommend PEG-based solutions in patients with IBD[18].

In fact, previous evidence has shown that in settings other than IBD, solutions based on sodium phosphate or sodium picosulphate for intestinal preparation had a 10-fold increased risk of developing intestinal mucositis compared with preparations with PEG[33].

Cases of post-preparation shock with picosulphate-based solutions have also been described[34] as well as of acute renal failure[35]. Unlike PEG-based solutions, these solutions are hyperosmolar and challenging to handle and contraindicated in patients with heart failure, rhabdomyolysis, hypermagnesemia, gastrointestinal ulcerative lesions, and renal failure[18].

***Role of simethicone as an additional component in bowel preparation***

Simethicone is a compound of dimethicone and 4%-7% silicon dioxide. This surfactant can reduce the surface tension of bubbles in the intestinal lumen by removing them. This improves clarity of endoscopic examination and reduces abdominal tension (Figure 1)[36].

Few studies have investigated the role of simethicone in intestinal preparations, specifically for patients with IBD. Metanalytic evidence has suggested its potential to improve mucosal cleanliness and visibility, providing evidence to increase the detection rate of adenomas and polyps[37,38]. Therefore, Lazzaroni *et al*[15] began experimenting in a randomised controlled trial with high-volume 4 L PEG-ELS in IBD, comparing with a double-arm design by adding 120 mg of simethicone.

In the study, the mean age of the participants was under 40 years for both arms. The cohort mainly consisted of UC patients. Regarding efficacy rates, bowel preparation was at least acceptable in 96% of cases in both arms. Although efficacy was not found to be a significant distinguishing element between the arms with or without simethicone, it was interesting to note a significant ameliorative effect on the presence of bubbles. Bubbles were either not detected or minimally impacted the examination in 98% of the sample with simethicone *vs* 85% of patients taking 4 L PEG alone. The tolerability of patient preparation increased in favour of the simethicone-based preparation. Generally, in this study, the addition or non-addition of simethicone did not dramatically affect the differences in safety. These were mainly comparable in terms of nausea, cramping pain, and abdominal bloating, except for sleep disturbances and general malaise, which were drastically lower in patients in the simethicone group (19% *vs* 44%). Bezzio *et al*[39] also showed good efficacy and tolerability of a 2 L PEG-ELS solution with added simethicone in 126 patients with IBD.

Further evidence on simethicone in patients with IBD, was provided by studies in a different setting, namely, studies of patients undergoing small-bowel capsule endoscopy[40-42]. One study observed that adding 80 mg of simethicone to a 2 L PEG preparation improved visualisation of the intestinal mucosa more in the proximal rather than in the distal tract, but only in non-CD patients. This effect was likely due to altered motility in CD patients[42]. In addition, in the paediatric population, adding simethicone to the PEG preparation appears to give good results in small-bowel capsule endoscopy; however, good visualisation of the terminal ileum remains challenging[41].

In any case, it is noteworthy that the most recent European guidelines recommend adding simethicone to intestinal preparations, cautiously leaning towards improved cleanliness and tolerability, despite a strong need for evidence to reaffirm this recommendation[18].

A study which was not solely conducted in the IBD population, analysed the optimal timing of simethicone administration in bowel preparation. The study suggested that optimal simethicone administration in the PEG-based preparation was in the evening of the day before colonoscopy. Moreover, the ameliorative effect is primarily at the expense of caecal intubation and bubble improvement[43]. This evaluation was further conducted in another study, combining a PEG preparation with ascorbic acid. Again, patients who took simethicone in the evening of the day before colonoscopy, showed fewer bubbles and improved detection of diminutive adenomas less than or equal to 5 mm[44].

Although not directly analysed in patients with IBD, improvement in the ease of caecal intubation[43] should be considered. Patients with ileal CD, both at the first diagnosis and at follow-up, should be carefully studied in the small intestinal tract. One of the most widely used scores is the Simple Endoscopic Score for Crohn's Disease[45] uses a thorough assessment of the ileum to identify the presence of ulcerative lesions and/or stenosis. This need is also reaffirmed by another endoscopic score, the Crohn's Disease Index of Severity, most commonly used in patients with CD[46].

The use of intraprocedural simethicone requires several technical considerations. A spectroscopic study of residual fluid samples detected with borescopes in the colonoscope channel found residual simethicone[47]. This study indicated that as simethicone is an inert and hydrophobic compound, it can reduce the effectiveness of endoscope reprocessing. In addition, simethicone is often included in solutions containing sugars which can potentially increase intra-endoscopic microbial growth[48]. Moreover, several dimethicone crystals have been detected in the waterjet channel of a damaged endoscope[49]. Therefore, European guidelines have warned about using simethicone at the lowest effective dose[50], exclusively in the biopsy channel and not in the auxiliary water channel[18].

***Role of high-volume colonic lavage as a bowel preparation strategy: A retrograde strategy***

In addition to conventional oral bowel preparations, a promising method of retrograde bowel preparation (already evaluated in non-IBD patients), high-volume water irrigation with colonic lavage, has recently been explored for IBD. This is undoubtedly a preparation modality that overcomes the obstacles and predictors of poor bowel preparation observed for oral bowel preparation. Moreover, it is likely to be more “palatable” to patients who experience problems with oral bowel preparation. However, it can be practiced only in a hospital setting, and after prescription by experienced practitioners. In a recent study, Gajera *et al*[51] examined this modality (Figure 2) in a retrospective study of more than 300 patients with IBD. The efficacy of bowel preparation was above 90% (except in patients with severe hemorrhoidal disease, in whom it was just under 90%).

Interestingly, a high efficacy rate of 94% was recorded, even for patients with previous gastrointestinal surgery. No severe reactions were observed; the most frequent mild reaction was abdominal pain in approximately 15% of patients. This study was not restricted to a specific IBD and had a good distribution of UC and CD. In this study, IBD patients were required to take bisacodyl (10 mg) the day before colonic lavage. In other non-IBD cases, this was magnesium hydroxide, 1-5 d before.

***Comparisons between preparations: What evidence in IBD?***

There is still a need for numerous studies on the preparations already available in singles for patients with IBD, especially in the different IBD subgroups with increased severity, such as patients with perianal disease and CD with stenosing phenotype. However, there are even fewer available data of head-to-head comparisons among different regimens.

A meta-analysis by Restellini *et al*[52] was designed specifically for patients with IBD with adequate bowel preparation as the primary outcome. The study included four previously described studies by Gould *et al*[13], Lazzaroni *et al*, Manes *et al*[53], and Kim *et al*[22].

The study by Gould *et al*[13] was excluded owing to a lack of clinical applicability of castor oil or senna. Furthermore, the study by Lazzaroni *et al*[15] was excluded owing to a lack of robust data on the efficacy of preparation with the addition of simethicone. The remaining two studies compared high- and low-volume PEG-based solutions. Examining these two analytical methods, the authors concluded that there were no relevant differences in preparation quality. Shifting the perspective onto the patient’s tolerability and willingness on the choice of regimen for subsequent colonoscopy, the winning regimens were the addition of simethicone to the 4 L PEG preparation, and the 2 L PEG preparation with bisacodyl or ascorbate compared with high volumes. Meta-analyses comparing PEG-based and non-PEG preparations are still needed. Even fewer comparative data exist between oral and retrograde rectal anterograde IBD preparations. An additional study, particularly relevant to colorectal cancer prevention in IBD, advised IBD patients undergoing chromoendoscopy to follow a clear fluid diet the day before colonoscopy[54].

***Predictors of IBD patients' poor bowel preparation?***

The extreme phenotypic variability of patients with IBD and the different regimens available for bowel preparation pose a problem for stratifying bowel preparation, identifying which patients are most at risk of poor bowel preparation, and with which preparations.

Several predictors of poor bowel preparation have been identified in PEG-based and UC-based studies, including male sex, non-split regimen, poor patient compliance (less than 100% intake), and moderate to severe discomfort during preparation[53]. The split regimen was a predictor of good bowel cleansing success in the study by Maida *et al*[25] with a 1 L PEG regimen with ascorbate. In another study that included both PEG-based and non-PEG regimens, PEG 2 L or 4 L regimens were associated with greater efficacy. Having a CD or colonoscopy in private *vs* public centres has also contributed to this[31]. Other evidence indicates that patients with active CD, experience more abdominal pain during bowel preparation, and patients with worse anxiety experience more symptoms during bowel preparation[55].

In contrast, a recent study by Kumar *et al*[56] highlighted an interesting finding relating bowel preparation to disease activity and biological therapy. Moderate-to-severe disease activity and biological therapy were predictors of suboptimal bowel preparation. Additional predictors of poor preparation identified in the study were, a non-split regimen, and patient age of over 65 years.

One Digestive Disease Week 2022 abstract presented a retrospective study of factors of inadequate bowel preparation for colonoscopy related to underlying IBD in 309 patients. The authors described how the presence of diabetes mellitus and antidepressant use were independent general risk factors in this setting. Indeed, it is well known that diabetes mellitus is a non-negligible comorbidity in IBD patients, increasing the risk of hospitalisations and infections, while not increasing the general risk of IBD complications or mortality[57]. In patients with UC, a history of inadequate bowel preparation was an independent risk factor[58].

One study aimed to investigate the experiences of IBD patients who resorted to repeat colonoscopy through telephone interviews. Despite the small sample size of approximately 33 patients, it emerged that patients felt that repeated colonoscopy was a guarantee of their health, and an ongoing reminder of the chronic and incurable nature of IBD[59]. This underscores how beyond looking for predictors and patient compliance, healthcare providers should strive to assist patients who require continuity of care. This is also in view of the fact that, as demonstrated by the COVID-19 pandemic, patients with IBD are exposed to a higher rate of anxiety and depression than the general population[60], even in disease remission[61]. Moreover, IBD patients are exposed to a non-negligible rate of treatment nonadherence[62,63].

***Patients with active IBD: What factors to consider?***

Bowel preparations can result in mucosal damage, with some studies having associated cases of toxic megacolon[64-67]. Therefore, caution becomes of utmost importance in patients with active IBD. PEG-based or sulphate-based solutions can induce colic mucosal damage by inducing metabolic or chemical damage[22]. Such histological damage results are also seen with sodium picosulphate[68]. As stated earlier, the risk of mucosal inflammation with PEG solutions is around ten times lower than that with those based on picosulphate or sulphate. Furthermore, picosulphate solutions can cause ulcerative lesions at the esophagogastric level[69,70]. In light of these facts, it is inevitable that for patients with active IBD, the only advisable solutions based on the available evidence, are PEG-based solutions[18,71]. Clearly, within active UC, a distinction must be made between moderate to severe and severe acute UC. Severe acute UC generally includes several diagnostic criteria (the presence of more than six bloody evacuations per day and at least one of the following: Body temperature greater than 37.8 °C, heart rate greater than 90 bpm, haemoglobin less than 105 g/L, and C-reactive protein greater than 30 mg/L. Under these conditions, the risk of impending megacolon, toxic megacolon, and colectomy is very high. Therefore, in addition to the initiation of treatment, it is appropriate to screen for confounding factors of disease activity. First, a faecal culture screening for *Clostridium difficile* should be performed. However, an evaluation of rectal biopsy for cytomegalovirus should also be performed. For cytomegalovirus evaluation, a complete colonoscopy is not recommended due to the high risk of bowel perforation and a simple sigmoidoscopy is appropriate[72]. Therefore, the problem of choosing the appropriate bowel preparation arises. In such cases, a simple phosphate enema generally suffices, as regular oral bowel preparation may increase the risk of colic dilatation[71].

***Prospects: Role of artificial intelligence***

More recently, artificial intelligence (AI) applications are emerging for use in digestive endoscopy, with indications from diagnosis to treatment[73-75]. In addition, promising AI results have also emerged for adenoma and polyp detection rates[76]. However, to date, the assessment of bowel preparation has been performed by the general application of scales by an endoscopist. Thus, assessment of bowel preparation is strongly dependent on the endoscopist[1].

However, even in bowel preparation, AI systems have been used in various experiments. For example, the ENDOANGEL system, which is based on deep convolutional neural network (CNN) technology, showed a higher BBPS calculation accuracy than operators with less than one year of experience, and operators with more than three years of experience. Furthermore, ENDOANGEL has an overall accuracy in classifying colonoscopy images of 91.89% and was associated with a 30-second reminder system for the endoscopist on bowel preparation assessment. Such a reminder system has the potential to overcome the limitations of BBPS. The reminder system is based on the selection of representative segments of different colic localisations. In contrast, this system is based on continuous video images, which are more representative of the bowel preparation of the whole colon[77]. The experience of Lee *et al*[78] also supports the use of AI systems, using two CNN algorithms and set on a BBPS. This study also provided encouraging results with an accuracy for inadequate bowel preparation of 85.3%, and an area under the curve of more than 0.8 (0.918).

Furthermore, the κ index of agreement between raters without and with AI was similar. Su *et al*[79] developed an automated quality control system for lower endoscopy, based on CNN. However, the system was not explicitly designed for exclusive evaluation of bowel preparation. Instead, it evaluated the withdrawal phase and stability, as well as the detection rate of colorectal polyps. In addition, the CNN system showed significant superiority in evaluating bowel preparation.

Ultimately, despite the paucity of available studies, the applications of AI and CNN in the real-time assessment of bowel preparation have non-negligible potential. Such an application may have a positive impact, improving several parameters including accuracy in the assessment of bowel preparation, prediction of the difficulty of caecal intubation, and estimation of the sensitivity of the examination in cancer screening.

**CONCLUSION**

Bowel preparation remains one of the main difficulties encountered by IBD patients undergoing colonoscopy. In general, PEG-based preparations appear to have the best safety profile and are recommended by endoscopic reference scientific societies for patients with IBD. Indeed, the timing of bowel preparation plays a relevant role, with split regimens being preferred. Caution must be exercised in patients with active intestinal inflammation due to the risk of mucosal damage associated with bowel preparation. New forms of preparation are emerging in both modalities, such as retrograde technology, with the integration of AI into quality assessment. However, new evidence is needed to enable tailoring of preparations to individual IBD patients. This will improve patient compliance and procedure efficacy.

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**Footnotes**

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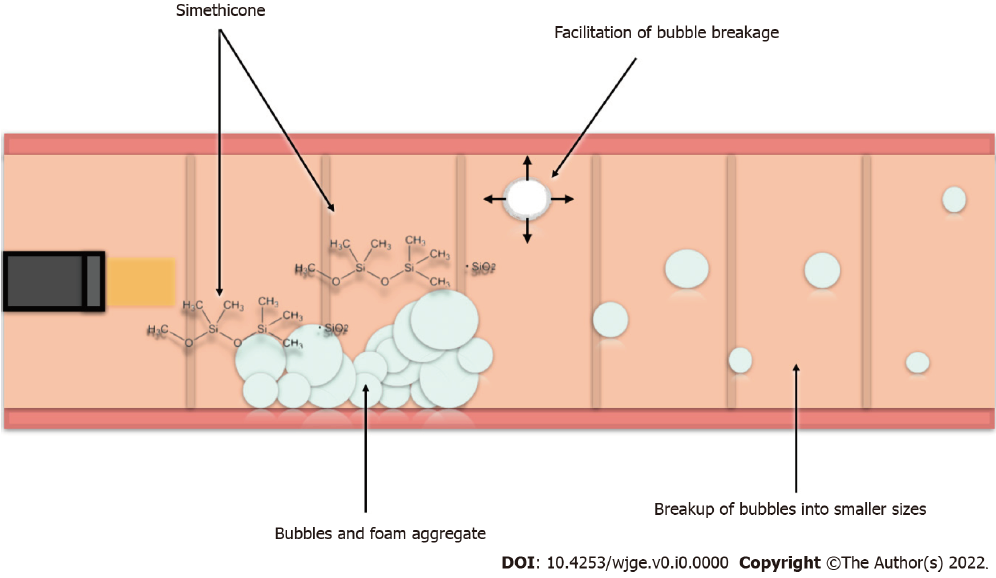
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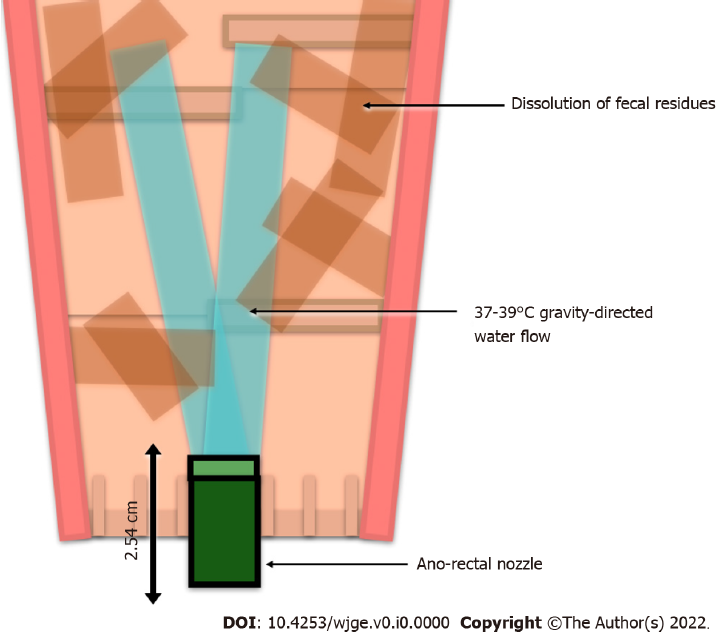
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**Figure Legends**



**Figure 1 Functioning of simethicone at the level of the intestinal mucosa.** Simethicone is a surfactant causing a reduction in the surface tension of intestinal bubbles. This reduction results in the aggregate of bubbles adhering to the colic mucosa being weaker with the facilitation of bubble reduction. As a result, larger bubbles are divided into smaller bubbles that have a greater ease of intestinal transit. The silicon dioxide component of dimethicone has an additional role, with an extensive molecular surface area that can promote bubble rupture. The breakdown of foam and bubbles and formed gas can be either absorbed by the intestinal wall or eliminated by intestinal transit. This likely explains the ameliorative effect on patients' symptoms.



**Figure 2 Colonic lavage process.** A nozzle is inserted approximately 2.54 cm into the rectum of the patient on a disinfected stand. According to the gravity gradient, a direct flow of water is dispensed from the nozzle at the rigidly controlled temperature of 37-39 °C, with immediate termination of the procedure if this temperature is exceeded. The water flow makes the stool soft, facilitating its dissolution and elimination. The procedure is generally complete within an hour, and the patient can subsequently undergo an endoscopic examination.

**Table 1 Anterograde and retrograde bowel preparation regimens studied in patients with inflammatory bowel disease**

|  |  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
| **Ref.** | **Year** | **Design** | **N** | **IBD** | **Bowel preparation** | **Split dosing considered** | **Low fiber diet considered** | **Main result (Success), %** | **Post-colonoscopy IBD flare-up rate, % (days after colonoscopy)** | **Severe AEs** |
| Gould *et al*[13] | 1982 | CT | 23 | UC: 23 | Castor oil 30 mL | - | YES | 82.6 | 26 (14-21 d) | NO |
| Gould *et al*[13] | 1982 | CT | 23 | UC: 23 | Sennosides 75 mg | - | YES | 86.9 | 48 (14-21 d) | NO |
| Lazzaroni *et al*[15] | 1993 | CT | 48 | UC: 26; CD: 23 | 4 L PEG-ELS plus placebo | YES | - | 96 | - | NO |
| Lazzaroni *et al*[15] | 1993 | CT | 57 | UC: 35; CD: 21 | 4 L PEG-ELS plus simethicone | YES | - | 96 | - | NO |
| Manes *et al*[53] | 2015 | CT | 106 | UC: 106 | 2 L PEG plus bisacodyl | YES | YES | 83 | - | NO |
| Manes *et al*[53] | 2015 | CT | 105 | UC: 105 | 4 L PEG-ELS | YES | YES | 77.1 | - | NO |
| Kim *et al*[22] | 2017 | CT | 53 | UC: 53 | 4 L PEG-ELS | YES | YES | 96.2 | 5.7 (7 d); 1.9 (28 d) | NO |
| Kim *et al*[22] | 2017 | CT | 56 | UC: 56 | 2 L PEG plus ascorbate | YES | YES | 92.9 | 3.6 (7 d); 1.8 (28 d) | NO |
| Briot *et al*[31] | 2019 | nCT Prospective |  | UC: 21; CD: 57; Unspecified IBD: 2 | Picosulphate-based regimen | YES | YES | 78.4 | 0 | NO |
| Bezzio *et al*[39] | 2020 | nCT  Prospective | 189 | UC: 63; CD: 63 | 2 L PEG-ELS plus simethicone | YES | - | UC: 89.8; CD: 86.2 | - | NO |
| Maida *et al*[25] | 2021 | nCT Retrospective | 185 | UC: 95; CD: 90 | 1 L PEG plus ascorbate solution | YES | YES | 92.9 | - | NO |
| Mohsen *et al*[30] | 2021 | CT | 61 | - | 2 L Sodium; picosulphate, magnesium citrate PEG | YES | YES | 89.5 | - | NO |
| Mohsen *et al*[30] | 2021 | CT | 64 | - | 2 L PEG plus ascorbate solution | YES | YES | - | NO |
| Neri *et al*[29] | 2021 | nCT Prospective | 103 | UC: 47; CD: 56 | 1 L PEG-ELS | YES | YES | 85.4 | - | NO |
| Kim *et al*[32] | 2022 | nCT Prospective | 52 | UC: 35; CD: 17 | 2 L PEG plus ascorbate | YES | YES | 98.1 | 0 (7 and 28 d) | NO |
| Kim *et al*[32] | 2022 | nCT Prospective | 55 | UC: 37; CD: 18 | Novel oral sulphate tablets | YES | YES | 98.1 | 3,63 (7 and 28 d) | NO |
| Gajera *et al*[51] | 2022 | nCT Retrospective | 318 | UC: 182; CD: 104; Unspecified IBD: 28 | Colonic lavage | Not applicable | YES | 97 | - | NO |

IBD: Inflammatory bowel disease; PEG: Polyethylene glycol; ELS: Electrolyte lavage solution; UC: Ulcerative colitis; CD: Crohn’s disease; CT: Clinical trial; nCT: Non-clinical trial.

**Table 2 Case reports on patients with known inflammatory bowel disease and relevant adverse events related to bowel preparation**

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
| **Ref.** | **Bowel preparation** | **Age** | **IBD** | **Gender** | **AE** | **Comorbidity** | **IBD therapy** | **Outcome** |
| Loraine *et al*[34] | Sodium picosulphate/magnesium oxide/citric acid | 73 | CD | F | Shock | Severe COPD, hypertension, dyslipidemia, cardiomyopathy, diverticular disease | Azathioprine | Dead |
| Gonlusen *et al*[35] | Sodium picosulphate | 56 | CD | F | Acute renal failure | GERD, healed gastric ulcer, endometriosis, mitral valve prolapse, migraine | None | Favourable |

IBD: Inflammatory bowel disease; CD: Crohn’s disease; F: Female; AE: Adverse event; COPD: Chronic obstructive pulmonary disease; GERD: Gastroesophageal reflux disease.