

# Systematic review of randomized controlled trials of probiotics, prebiotics, and synbiotics in inflammatory bowel disease

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**Background:** Probiotics are microorganisms that are ingested either in combination or as a single organism in an effort to normalize intestinal microbiota and potentially improve intestinal barrier function. Recent evidence has suggested that inflammatory bowel disease (IBD) may result from an inappropriate immunologic response to intestinal bacteria and a disruption in the balance of the gastrointestinal microbiota in genetically susceptible individuals. Prebiotics, synbiotics, and probiotics have all been studied with growing interest as adjuncts to standard therapies for IBD. In general, probiotics have been shown to be well-tolerated with few side effects, making them a potential attractive treatment option in the management of IBD.

**Aim:** To perform a systematic review of randomized controlled trials on the use of probiotics, prebiotics, and synbiotics in IBD.

**Results:** In our systematic review we found 14 studies in patients with Crohn's disease (CD), 21 studies in patients with ulcerative colitis (UC), and five studies in patients with pouchitis. These were randomized controlled trials using probiotics, prebiotics, and/or synbiotics. In patients with CD, multiple studies comparing probiotics and placebo showed no significant difference in clinical outcomes. Adding a probiotic to conventional treatment improved the overall induction of remission rates among patients with UC. There was also a similar benefit in maintaining remission in UC. Probiotics have also shown some efficacy in the treatment of pouchitis after antibiotic-induced remission.

**Conclusions:** To date, there is insufficient data to recommend probiotics for use in CD. There is evidence to support the use of probiotics for induction and maintenance of remission in UC and pouchitis. Future quality studies are needed to confirm whether probiotics, prebiotics, and synbiotics have a definite role in induction or maintenance of remission in CD, UC, and pouchitis. Similar to probiotics, fecal microbiota transplantation provides an alternate modality of therapy to treat IBD by influencing the intestinal flora.

**Keywords:** inflammatory bowel disease, Crohn's disease, ulcerative colitis, pouchitis, probiotics, prebiotics, synbiotics

## Introduction

Inflammatory bowel disease (IBD) is a chronic relapsing inflammatory disorder of the gastrointestinal tract, comprising ulcerative colitis (UC) and Crohn's disease (CD). IBD affects more than 1 million people in the United States and 2.5–3 million people in Europe.<sup>1,2</sup> The standard treatment approach for patients with CD and UC has traditionally been with the use of anti-inflammatory agents, like mesalamine, corticosteroids, immunomodulators such as azathioprine, and biologic agents such as infliximab. Although the exact pathophysiology of IBD is unknown, the current

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paradigm is multifactorial and involves an abnormal intestinal immune response to the gut microbiota in a genetically susceptible host.<sup>3</sup>

Interest in microbiota-based therapy has grown markedly, due in large part to an improved safety profile with fewer side effects when compared to traditional therapy. It is important to note, however, that there are only a limited number of large, well-designed prospective trials that could evaluate for potential rare side effects in IBD patients, many of whom are immunocompromised. Collectively, the gut microbiota contains a total of  $10^{13}$  to  $10^{14}$  microorganisms, with the majority ( $10^{12}$  microorganisms) hosted within the colon.<sup>4</sup> *Firmicutes* and *Bacteroidetes* are the main bacterial phyla in the gut, comprising about 90% of the microbiota, followed to a lesser proportion by *Actinobacteria*, *Proteobacteria*, *Verrucomicrobia*, *Fusobacteria*, and *Cyanobacteria*.<sup>5</sup> Furthermore, mucosa-associated bacterial communities in the colon are significantly different in composition from those in feces.<sup>6</sup>

Studies in human and mouse models over the past several years have investigated the relationship between dysbiosis of the gut microbiota and intestinal diseases. Mouse models have further advanced our understanding of the pathophysiology of IBD, showing that bacteria play a fundamental role in the initiation and development of chronic intestinal disease. A pivotal study showed that germfree interleukin (IL)-10-deficient mice did not develop colitis compared to IL-10 deficient mice with normal enteric bacteria.<sup>7</sup> Other studies have shown that germfree mice given dextran sodium sulfate, which induces colitis, developed more severe disease than did non-germfree mice, suggesting a protective effect of the microbiota.<sup>8</sup> In human model analysis using 16S ribosomal (r)RNA sequencing, the microbiota of IBD patients differed from that of healthy individuals. In IBD, the diversity and bacterial load of *Bacteroidetes* and *Firmicutes* are decreased, especially in regions of active inflammation. These groups of bacteria are particularly important as they produce short-chain fatty-acid metabolites. The short-chain fatty-acid substrates have potent anti-inflammatory properties, are important sources of energy for colonic epithelial cells, and may enhance epithelial barrier integrity. Finally, microbiota changes in IBD have a higher proportion of *Actinobacteria* and *Enterobacteria*, phyla which contain many of the known gut pathogens.<sup>9</sup>

Because of this association between the intestinal microflora and IBD, multiple studies have investigated the use of probiotics, prebiotics, and synbiotics to modify the gut microbiota in order to replace or augment conventional IBD therapies. Probiotics have been defined as live

microorganisms that confer a health benefit to the host when ingested. Prebiotics are nondigestible substances acting as a nutritive substrate to stimulate the growth and metabolism of protective endogenous enteric bacteria. Synbiotics are a combination of both probiotics and prebiotics.<sup>10</sup> Probiotics are believed to contribute to the normal functioning of the intestinal commensal microbiota. They also displace pathogenic microbes, improve host mucosal barrier function, and modulate the innate host immune function.<sup>11</sup> Finally, recent data suggest that a new approach to altering the gut microbiota by fecal microbiota transplantation has gained popularity in the treatment of *Clostridium difficile* infection and may be a promising option in treating IBD, as well.<sup>12</sup> For that reason it also will be discussed in this review.

Multiple studies have been done recently on the use of probiotic, prebiotic, and synbiotic agents as an adjuvant therapy in the treatment of CD and UC. Treatment approaches for these conditions can be divided into treatment during an acute flare (induction therapy) and treatment for long-term control of symptoms (maintenance therapy). Pouchitis is a clinical diagnosis for nonspecific, idiopathic inflammation of the ileal reservoir created by an ileal pouch–anal anastomosis, which is a commonly employed procedure following total proctocolectomy for patients with UC.<sup>13</sup> Previous studies have shown clinical benefit of some probiotics in pouchitis and UC, but less clearly in CD.<sup>14–17</sup> The aim of the present article is to provide an overview of randomized controlled trials (RCTs) in human studies of CD, UC, and pouchitis and to draw practical clinical conclusions about the efficacy of probiotics, prebiotics, and synbiotics in treating patients with these conditions.

## Methods

We performed a literature search in Medline, PubMed, Cochrane database, Science Direct, Ovid, and Scopus. The results were limited to full-text articles published in English. The search terms used were “inflammatory bowel disease”, “IBD”, “inflammation”, “ulcerative colitis”, “Crohn’s disease”, and “pouchitis”. These terms were cross-referenced with the terms “probiotic”, “prebiotic”, and “synbiotic” in all possible combinations. Only RCTs performed among human subjects were selected for this review. Studies both in pediatric and adult populations were included. The initial search yielded 4,440 results: the search terms “inflammation” or “inflammatory bowel disease” and “prebiotic”, “probiotic”, or “synbiotic” yielded 3,188 results; the remaining 1,252 results were obtained by searching the individual types of inflammatory bowel disease with “probiotic”, “prebiotic”, or “synbiotic”. Several of these results provided redundant articles.

The authors reviewed the titles and/or abstracts to exclude studies that were not RCTs performed with IBD patients using probiotics, prebiotics, or synbiotics. We excluded any animal studies or human cell or tissue culture studies. Those trials that did not define clear randomization of the control and the study group were excluded. After applying the exclusion criteria, we selected 82 studies for further review. Full-text versions of these articles were reviewed in detail by the authors. Finally, 40 RCTs were selected for inclusion in our systematic review, including 14 CD studies, 21 UC studies, and five pouchitis studies. The excluded articles consisted of nine CD studies, 26 UC studies, and seven pouchitis studies. A description of the article selection process is shown in Figure 1.

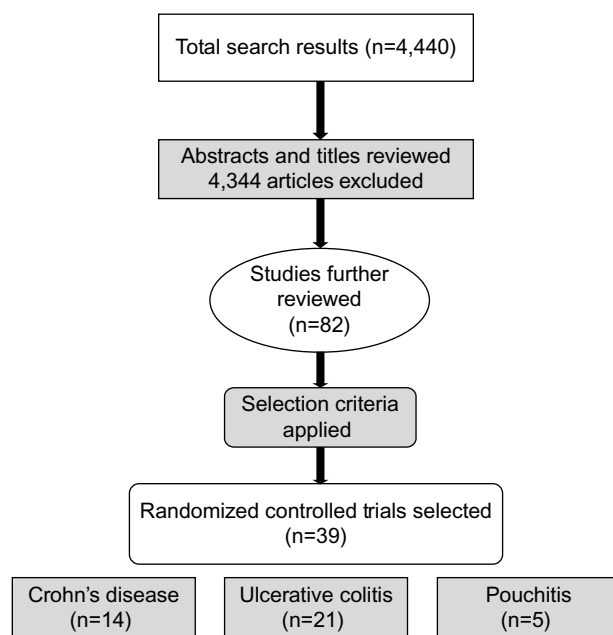
## Results

### Crohn's disease

In our literature search we found 14 RCTs that aimed to determine the outcome following probiotic, prebiotic, or synbiotic treatment for the management of CD<sup>18–31</sup> (described in Table 1). One study was performed among CD and UC patients.<sup>30</sup> Ten studies included only probiotics in the intervention group.<sup>19–25,27,31</sup> Of these, five used *Lactobacillus*.<sup>21–25</sup> The species *Lactobacillus GG* (LGG) was used in three trials and *Lactobacillus johnsonii* was used in two trials.<sup>21–23</sup> The primary end point in three studies was endoscopic recurrence rate and in two studies was clinical relapse rate. The Crohn's Disease Activity Index for adults

(CDAI, which measures the clinical severity of CD and helps in determining if there is active disease, relapse and severity of relapse)<sup>32</sup> was used in eight studies conducted in the adult population.<sup>20,22,24,25,27–29,31</sup> A study by Bousvaros et al<sup>23</sup> performed in a pediatric population used the Pediatric Crohn's Disease Activity Index score<sup>33</sup> to evaluate the severity of clinical symptoms; this study was stopped prematurely at 42 months because an interim data analysis showed no efficacy with use of the probiotics and because enrolling the pediatric population was becoming more challenging for the long duration of the study. In essence, none of the *Lactobacillus*-containing probiotics used in the CD study populations showed any meaningful effect on endoscopic remission or on the CDAI. Among the other five probiotic-only trials, *Saccharomyces boulardii* was used in four studies.<sup>18,20,27,31</sup> One study by Malchow<sup>19</sup> conducted in 1997 among CD patients used *Escherichia coli* strain Nissle 1917. This was a double-blinded placebo-controlled pilot study conducted over 1 year in 28 patients who were in remission from CD. The remission rates among the probiotic and placebo groups were not statistically different. The study also showed that *E. coli* strain Nissle 1917 produced no clinical harm among the study population.

Plein and Hotz<sup>18</sup> performed a pilot study in 1997 using *S. boulardii*. This was a double-blinded placebo-controlled trial conducted among 17 patients with CD. The study showed clinical benefit in decreasing the frequency of stools and improving CDAI scores.<sup>32</sup> In 2000, a study published by Guslandi et al<sup>20</sup> showed clinical benefit with use of *S. boulardii* as determined by a decrease in the rate of clinical relapse, which was measured by the CDAI; however, the study contained only 16 patients in each of the control and intervention groups. In 2013, Bourreille et al<sup>31</sup> published a study comparing the use of *S. boulardii* in 84 patients with 81 controls. This larger trial did not show any clinical benefit in relapse rate as determined by the CDAI score at the end of 40 weeks of follow-up. The study was well designed and included a total of 165 individuals. The final study from Garcia Vilela's group<sup>27</sup> in 2008 examined 14 and 17 CD patients in remission (defined by CDAI) who were treated with *S. boulardii* or placebo, respectively. They measured intestinal permeability with a lactulose:mannitol ratio, which has been shown to be associated with increased disease activity. While this study showed a reduction in the lactulose:mannitol ratio in the probiotic group, the data were difficult to interpret or apply given the wide variability of intestinal permeability in CD patients and lack of correlation with CDAI or endoscopic remission. These three studies described above showed benefit in using



**Figure 1** Flowchart showing randomized controlled study selection criteria for inclusion in the systematic review.

Table 1 Probiotics, prebiotics and synbiotics in Crohn's disease

Publication (year, country)	First author	Type of intervention	Agent used	N	Control group	Intervention group	Primary end point	Statistical benefit (P-value)	Trend toward clinical benefit or harm
1993, Germany	Plain <sup>18</sup>	Probiotic	<i>Saccharomyces boulardii</i>	17	7	10	Clinical remission	Yes (<0.01)	Benefit
1997, Germany	Malchow <sup>19</sup>	Probiotic	<i>Escherichia coli</i> strain Nissle 1917	28	12	16	Clinical relapse rate	No (n too small to calculate P-value)	Benefit
2000, Italy	Guslandi <sup>20</sup>	Probiotic	<i>Saccharomyces boulardii</i>	32	16	16	Clinical relapse rate	Yes (0.04)	Benefit
2002, Italy	Prantera <sup>21</sup>	Probiotic	<i>Lactobacillus GG</i>	45	22	23	Endoscopic recurrence rate	No (0.297)	Harm
2004, Germany	Schultz <sup>22</sup>	Probiotic	<i>Lactobacillus GG</i>	11	6	5	Mean time to clinical relapse	No (0.5)	Benefit
2005, USA	Bousvaros <sup>23</sup>	Probiotic	<i>Lactobacillus GG</i>	75	36	39	Clinical relapse rate	No (0.18)	Harm
2006, France	Marteau <sup>24</sup>	Probiotic	<i>Lactobacillus johnsonii</i>	98	50	48	Endoscopic recurrence rate	No (0.15)	Benefit
2007, Belgium	Van Gossum <sup>25</sup>	Probiotic	<i>Lactobacillus johnsonii</i>	70	36	34	Endoscopic recurrence rate	No (0.32)	Benefit
2007, Israel	Chermesh <sup>26</sup>	Synbiotic	Synbiotic 2000	30	10	20	Endoscopic, clinical and laboratory parameters	No (not significant)	None
2008, Brazil	Garcia Vilela <sup>27</sup>	Probiotic	<i>Saccharomyces boulardii</i>	31	17	14	Intestinal permeability (actulose:mannotol ratio)	Yes (0.0005)	Not applicable
2010, UK	Steed <sup>28</sup>	Synbiotic	<i>Bifidobacterium longum</i> and Synergy 1	24	11	13	Clinical response	Yes (0.02)	Benefit
2011, UK	Benjamin <sup>29</sup>	Prebiotic	Fructo-oligosaccharides	103	49	54	Clinical response	No (0.067)	Benefit
2007, Germany	Hafer <sup>30</sup>	Prebiotic	Lactulose	31*	9	8	Clinical response (CDAI and SAI)	No (>0.1 and 0.097)	Harm
2013, France	Bourreille <sup>31</sup>	Probiotic	<i>Saccharomyces boulardii</i>	165	81	84	Clinical relapse rate	No (0.5)	None

Notes: Randomized controlled trials that have studied the use of various probiotic/prebiotic/synbiotic agents in Crohn's disease with primary end points; number of patients involved in the study, and outcomes as shown. \*Study by Hafer et al<sup>30</sup> comprised a total of 31 study subjects, of whom 17 had Crohn's disease and 14 had ulcerative colitis.

Abbreviations: CDAI, Crohn's Disease Activity Index; N, total number of patients enrolled at the beginning of the study; n, number; SAI, Severity and Activity Index.

*S. boulardii*, but their major limitation was the small size of the study populations.

Studies using prebiotics and synbiotics in CD are further described. Two studies were found using only prebiotics for therapy in CD.<sup>29,30</sup> In 2011, Benjamin et al<sup>29</sup> performed a study with 103 subjects (49 in the control group and 54 in the intervention group) using fructo-oligosaccharides. Clinical response was measured using CDAI score; however, there was no clinical benefit at the end of 4 weeks of therapy. Another trial using lactulose as a prebiotic was performed in 2007 by Hafer's group among patients with UC and CD.<sup>30</sup> There was no clinical benefit in treatment among the CD patients but improvement in quality of life was seen in those with UC. Hafer et al<sup>30</sup> used the CDAI and Severity and Activity Index<sup>34</sup> scoring systems to determine clinical severity of the disease. Both studies showed no improvement in the scores in the prebiotic group; in fact, placebo groups showed a better response.

Two studies evaluated the use of a synbiotic preparation among patients with CD.<sup>26,28</sup> One study, published in 2010 by Steed et al,<sup>28</sup> consisted of eleven patients in the placebo group in comparison to 14 in the synbiotic group. The synbiotic preparation consisted of *Bifidobacterium longum* and Synergy 1, a preferential growth substrate consisting of an inulin/oligofructose mix. At the end of 3 months and 6 months of therapy, there was significant improvement in CDAI scores in the synbiotic group in comparison to the placebo group. A major limitation of the study was that the baseline CDAI scores were higher in the placebo group. A second synbiotic study conducted by Chermesh et al<sup>26</sup> used Synbiotic 2000, a combination of four probiotic and four prebiotic agents. It consisted of four lactic acid-producing bacteria: *Pediococcus pentoseceus*, *Lactobacillus affinolactis*, *Lactobacillus paracasei susp paracasei 19* and *Lactobacillus plantarum 2362*. It also consisted of four fermentable fibers:  $\beta$ -glucans, inulin, pectin, and resistant starch. The study included ten patients in the placebo group and 20 in the intervention group. Clinical, laboratory, and endoscopic parameters were used to compare the two groups for 24 months. At completion of the study, there was no statistically significant benefit among the intervention group in terms of improvement in clinical symptoms (CDAI score), endoscopic features (Rutgeerts' scoring system)<sup>35</sup> or laboratory measurements. Given the findings above, there are no clinical benefits found with prebiotics and synbiotics in patients with CD.

## Ulcerative colitis

Probiotics, prebiotics, and synbiotics have all been studied as therapeutic adjuncts in the management of ulcerative colitis.

A total of 21 RCT studies were identified that evaluated the use of various probiotics, prebiotics, and/or synbiotics used as induction therapy, maintenance of remission, and/or combination of both in patients with UC.

### Ulcerative colitis – induction

Our literature search found nine studies using various types of probiotics, prebiotics, and synbiotics for induction therapy in UC<sup>30,36–43</sup> (described in Table 2). The majority of these studies enrolled UC patients with mild to moderate disease activity, with the largest trials using VSL#3. VSL#3 is a multibacterial culture for oral use, consisting of the following species: *Bifidobacterium breve*, *B. longum*, *Bifidobacterium infantis*, *Lactobacillus acidophilus*, *L. plantarum*, *L. paracasei*, *Lactobacillus bulgaricus*, and *Streptococcus thermophilus*.<sup>44</sup> Tursi et al<sup>36</sup> in 2004 evaluated 30 patients receiving VSL#3 at a dose of 900 billion bacteria daily with low-dose balsalazide, 30 UC patients with medium-dose balsalazide alone, and another 30 UC patients with mesalamine alone. The rate of clinical remission was comparable between low-dose balsalazide with VSL#3 and medium-dose balsalazide, but statistical analysis showed that a daily dose of VSL#3 with balsalazide was significantly superior to mesalamine in achieving symptomatic remission. This dose of VSL#3 was lower than the dose used in the largest of the UC induction of remission studies done by Sood et al<sup>39</sup> in 2009 and Tursi et al<sup>40</sup> in 2010. These two studies used VSL#3 at a dose of 3600 billion bacteria daily. These two studies enrolled 147 and 144 patients, respectively, with mild to moderate UC and concomitant use of aminosalicylates or thiopurines. Both studies showed a decrease in the Ulcerative Colitis Disease Activity Index<sup>45</sup> by 50% or more in the treatment arm. At 12 weeks, Sood et al<sup>39</sup> observed that 42.9% of the VSL#3 treated patients had achieved remission compared with 15.7% of patients with placebo ( $P < 0.001$ ). Tursi et al<sup>40</sup> observed that 47.7% of patients treated with VSL#3 were in remission compared to 32.4% in the placebo group; however this did not reach statistical significance. Limitations of this study include a high placebo response rate and relatively short duration of study.

On a cellular level, VSL#3 affected dendritic cells, which shape bacterial recognition and T-cell responses, in patients with acute UC in a study by Ng et al.<sup>41</sup> There was an increase in anti-inflammatory IL-10 production and decrease in proinflammatory IL-12p40 production by colonic dendritic cells following VSL#3 treatment. These data suggest an association between a change toward beneficial dendritic cell cytokine profiles and clinical efficacy.

**Table 2** Probiotics and ulcerative colitis induction

Publication (year, country)	First author	Type of intervention	Agent used	N	Control group	Intervention group	Primary end point	Statistical benefit (P-value)	Trend toward clinical benefit or harm
2004, Italy	Tursi <sup>36</sup>	Probiotic	VSL#3	90	60	30	Rate of clinical remission	Yes (<0.02)	Benefit
2004, Japan	Kato <sup>37</sup>	Probiotic	Bifidobacteria-fermented milk	20	10	10	Clinical remission: improved CAI	Yes (0.05)	Benefit
2005, UK	Furrie <sup>38</sup>	Synbiotic	<i>Bifidobacterium longum</i> and Synergy 1	18	9	9	Sigmoidoscopy score and inflammatory markers	No (0.06) and yes (<0.05)	Benefit
2009, India	Sood <sup>39</sup>	Probiotic	VSL#3	147	70	77	Improvement in clinical activity	Yes (0.001)	Benefit
2010, Italy	Tursi <sup>40</sup>	Probiotic	VSL#3	144	73	71	Improvement in clinical activity	Yes (0.01)	Benefit
2010, UK	Ng <sup>41</sup>	Probiotic	VSL#3	28	14	14	Effect on dendritic cell cytokines	Yes (<0.05)	Benefit
2010, Germany	Matthes <sup>42</sup>	Probiotic	<i>Escherichia coli</i> Nissle 1917 enemas	90	20	24, 23, 23 (different doses)	Dose-dependent clinical remission	Yes (0.0446)	Benefit
2007, Germany	Hafer <sup>30</sup>	Prebiotic	Lactulose	31*	7	7	Clinical and endoscopic response	No (0.092)	Benefit
2012, People's Republic of China	Li <sup>43</sup>	Probiotic	Bifid Triple Viable	82	41	41	Clinical and mucosal inflammation	Yes (<0.01)	Benefit

**Notes:** Randomized controlled trials showing probiotics/prebiotics/synbiotics used for induction therapy in ulcerative colitis. \*Study by Hafer et al<sup>30</sup> comprised a total of 31 study subjects, of whom 17 had Crohn's disease and 14 had ulcerative colitis.

**Abbreviations:** CAI, Clinical Activity Index.

Similar increased IL-10 production was seen in patients treated with steroids.

Three other trials evaluated other probiotic regimens and one trial evaluated a synbiotic regimen in remission of UC. Bifidobacteria-fermented milk (BFM), which contains *Bifidobacterium* strains and *L. acidophilus*, was studied by Kato et al<sup>37</sup> in a small study of 20 patients. They achieved their primary end point of a decrease in the Clinical Activity Index (CAI) score<sup>46</sup> of at least 3 points by 12 weeks. The BFM-treated group's CAI score was significantly lower than the placebo score at 12 weeks ( $P<0.05$ ) along with improved Endoscopic Activity Index<sup>47</sup> and histological scores. Li et al<sup>43</sup> evaluated the use of Bifid Triple Viable, which is a combination of *Bacillus acidophilus*, *Bifidobacterium bifidum*, and fecal streptococci, combined with 5-aminosalicylic acid compared to patients taking 5-aminosalicylic acid alone. Clinical symptoms based on the Mayo Score<sup>48</sup> and colonic mucosal inflammation improved more in the treatment group. Matthes et al<sup>42</sup> evaluated the efficacy of *E. coli* strain Nissle 1917 in patients with proctitis and proctosigmoiditis. This was the only study that evaluated delivery of a probiotic via an enema for the treatment of acute UC. Their study examined several different doses of the probiotic, which resulted in small subgroups. Per protocol analysis indicated a statistically significant difference in dose-dependent efficacy ( $P=0.0446$ ); however, they failed to show statistical significance in the intent-to-treat analysis.

A small study done by Furrie et al<sup>38</sup> used a novel synbiotic consisting of *B. longum* and a combination of a prebiotic (Synergy 1) for the probiotic strain for 4 weeks. Much of their focus was on the effects of the treatment with respect to inflammatory markers and the presence of the bacteria in mucosal biopsies. There was a statistically significant reduction in the inflammatory markers among the synbiotic group when compared to the placebo group. Clinically they observed a trend toward improved sigmoidoscopy scores (without statistical significance) and no significant difference in CAI scores. However the data were limited by both a small study population (n=18) and a high dropout rate. As mentioned previously, Hafer et al,<sup>30</sup> published a study in 2007 which showed no clinical or endoscopic benefit in the use of lactulose as a prebiotic among 17 CD and 14 UC patients. However, UC patients showed an improvement in the quality of life. Use of the prebiotic in CD and UC showed no benefit in terms of clinical activity index, endoscopic score, or immunohistochemical parameters; however, UC patients showed an improvement in the quality of life. The study was limited by the size of the population and lack of placebo use in

the control group, meaning that we cannot rule out a placebo effect in the treatment group.

### Ulcerative colitis – maintenance

Nine studies have been published examining the use of probiotics in the maintenance of UC, and three studies evaluated both combined induction and maintenance of remission in UC. These 12 studies<sup>49–60</sup> are described in Table 3.

A study by Kruis et al<sup>49</sup> in 1997 using *E. coli* Nissle 1917 in comparison with mesalamine in a population of 103 patients showed no benefit in the probiotic group in comparison with the mesalamine group. Another study by Kruis et al<sup>52</sup> in 2004 randomized 327 patients to receive either *E. coli* Nissle 1917 (n=162) or mesalamine (n=165). The *E. coli* group had a relapse rate of 56.4% compared to 33.9% in the placebo group over a 12 month period ( $P=0.003$ ). The CAI trended towards a greater increase in the *E. coli* group; however, there were no statistically significant differences in quality of life score, endoscopy scores, and histology scores. In 2011, Wildt et al<sup>57</sup> enrolled 32 patients with left-sided UC in remission to take either placebo (n=12) or a probiotic consisting of *L. acidophilus* La-5 and *Bifidobacterium animalis* subspecies *lactis* BB-12. Although no statistically significant benefit was seen in the primary end point of maintaining remission, there was a trend toward increased remission in the probiotic group. In 2004, Cui et al<sup>53</sup> studied 30 patients with UC initially treated with sulfasalazine and glucocorticoids and then randomly administered either Bifid Triple Viable capsule or an identical placebo (starch). Fecal analysis and histological analysis were performed after 8 weeks or in cases of relapse. The study showed that high concentration of luminal gram-positive bacteria were seen in the probiotics group (n=15), which could prolong the time to relapse of ulcerative colitis. Also, proinflammatory cytokines were inhibited by probiotics, and messengerRNA expression of anti-inflammatory cytokine IL-10 was elevated by the effect of probiotics. In 2010, Hegazy and El-Bedewy<sup>58</sup> used a probiotic preparation containing *Lactobacillus delbrueckii* and *Lactobacillus fermentum* for 8 weeks in newly diagnosed UC patients (n=15) who were being treated with sulfasalazine compared to those taking sulfasalazine and placebo. Histological improvement of inflammation was significantly improved in the probiotic group compared to the sulfasalazine group and was associated with decreased fecal calprotectin and mucosal inflammatory cytokines.

There were two open-label prospective RCTs, one evaluating the use of probiotics and the other evaluating the use of synbiotics for maintenance therapy in UC. Ishikawa et al<sup>51</sup> in

2003 enlisted 21 patients in an open-label RCT using BFM (*B. breve*, *B. bifidum*, and *L. acidophilus*) combined with standard therapy in eleven patients. There were significantly fewer episodes of symptomatic exacerbations in the probiotic group observed over a period of one year when compared to standard therapy (27% versus 90%,  $P=0.0075$ ). In 2006, Zocco et al<sup>54</sup> studied a sample size of 187 patients in an open-label RCT using LGG as the probiotic with three treatment groups: LGG alone, LGG combined with mesalamine, and mesalamine alone. The primary end point was the number of relapses over 12 months. There was no significant difference in the relapse rates among the three groups, and the authors concluded that they established equivalent efficacy. Fujimori et al,<sup>55</sup> in 2009, also conducted a large (n=120) prospective open-label RCT using a synbiotic in patients with UC who were either in remission or had mildly active disease. The patients were randomly assigned into one of the three groups: probiotic (*B. Longum*) group, prebiotic (psyllium) group and synbiotic (a combination of the aforementioned prebiotic and probiotic) group. We arbitrarily have labeled these three groups as group a, b, and c in Table 3. The primary end point was scores on the Inflammatory Bowel Disease Questionnaire,<sup>61</sup> which assesses health-related quality of life in IBD, at 4 weeks. Results showed a statistically significant improvement in scores for the synbiotic group at the end of the study. This improvement was not seen in the probiotic or prebiotic groups. The limitations of this study included a high dropout rate, no endoscopic or histological examinations, and short duration. In 2012, Oliva et al<sup>60</sup> performed a double-blinded RCT among a pediatric population with UC in remission. Fifteen controls received placebo and mesalamine and 16 patients received a probiotic, *Lactobacillus reuteri* ATCC 55730 with mesalamine. The results of the study showed improvement in clinical and endoscopic findings (Mayo score) and decrease in severity of inflammation/inflammatory markers in the probiotic group.

### Ulcerative colitis – induction and maintenance

The persistent effect of both induction and maintenance of remission in UC was evaluated in three double-blind RCTs. In 1999, Rembacken et al<sup>50</sup> studied 116 patients with active UC over a period of 12 months. Patients received pretreatment with oral gentamicin followed by a nonpathogenic strain of *E. coli* Nissle 1917 serotype O6:K5:H1 and were compared to patients receiving pretreatment with gentamicin and standard treatment with mesalamine with respect to induction of remission and prevention of relapse. Thirty-nine of 57 patients (68%) in the *E. coli* group achieved remission compared to

**Table 3** Probiotics in ulcerative colitis maintenance and induction + maintenance

Publication (year, country)	First author	Type of intervention	Agent used	N	Control group	Intervention group	Primary end point	Statistical benefit (P-value)	Trend toward clinical benefit or harm
1997, Germany	Kruis <sup>49</sup>	Probiotic	<i>Escherichia coli</i> Nissle 1917	103	53	50	Remission and relapse rate	No (>0.05)	None
1999, UK	Rembacken <sup>50</sup>	Probiotic	Nonpathogenic strain of <i>Escherichia coli</i> (serotype O6:K5:H1) Nissle 1917	116	59	57	Remission and relapse rate	No (>0.05) and yes (0.0059)	Benefit
2003, Japan	Ishikawa <sup>51</sup>	Probiotic	Bifidobacteria-fermented milk	21	10	11	Clinical relapse and duration of remission	Yes (0.0059 and 0.0174)	Benefit
2004, Germany	Kruis <sup>52</sup>	Probiotic	<i>Escherichia coli</i> Nissle 1917	327	165	162	Rate of relapse	Yes (0.003)	Benefit
2004, People's Republic of China	Cui <sup>53</sup>	Probiotic	Bifid Triple Viable capsule	30	15	15	Rate of relapse and fecal/mucosal response	Yes (<0.01 and <0.05)	Benefit
2006, Italy	Zocco <sup>54</sup>	Probiotic	LGG LGG + mesalazine	187	62	65,* 60**	Maintenance of clinical remission	No (>0.05)	Benefit
2009, Japan	Fujimori <sup>55</sup>	Probiotic, prebiotic, synbiotic	<i>Bifidobacterium longum</i> , psyllium, combined	120	0	40, <sup>a</sup> 40, <sup>b</sup> 40 <sup>c</sup>	Improvement in quality of life: IBDQ score	Yes (0.03)	Benefit
2009, Italy	Miele <sup>56</sup>	Probiotic	VSL#3	29	15	14	Maintenance of remission and relapse rate	Yes (<0.01 and 0.014)	Benefit
2009, Denmark	Wildt <sup>57</sup>	Probiotic	<i>Lactobacillus acidophilus</i> La-5 and <i>Bifidobacterium animalis</i> subsp. <i>Lactis</i> BB-12	32	12	20	Maintenance of clinical remission	No (0.37)	Benefit
2010, Egypt	Hegazy <sup>58</sup>	Probiotic	<i>Lactobacillus delbruekii</i> and <i>Lactobacillus fermentum</i>	30	15	15	Decrease in colonic inflammation	Yes (<0.05)	Not applicable
2011, Japan	Ishikawa <sup>59</sup>	Synbiotic	<i>Bifidobacterium breve</i> Yakult and galacto-oligosaccharide	41	20	21	Endoscopic score: Matt's Classification	Yes (<0.05)	Benefit
2012, Italy	Oliva <sup>60</sup>	Probiotic	<i>Lactobacillus reuteri</i> ATCC 55730	31	15	16	Clinical and endoscopic remission: Mayo Score and inflammatory markers	Yes (<0.01)	Benefit

**Notes:** Randomized controlled trials that have studied the use of various probiotic/prebiotic/synbiotic agents in the treatment of either induction, maintenance, or induction + maintenance of ulcerative colitis with primary end points, number of patients involved in the study, and outcomes as shown. \*LGG agent used. \*\*LGG + mesalazine agent used. <sup>a</sup>Probiotic type of intervention. <sup>b</sup>Prebiotic type of intervention. <sup>c</sup>Synbiotic type of intervention.  
**Abbreviations:** ATCC, American Type Culture Collection; IBDQ, Inflammatory Bowel Disease Questionnaire; LGG, *Lactobacillus GG*; N, total number of patients enrolled in the study at the beginning of the study.

44 (75%) in the mesalamine group, which was not statistically significant ( $P>0.05$ ). Thirty-two patients (73%) in the mesalamine group relapsed within 1 year compared with 26 (67%) in the *E. coli* group, which was borderline statistically significant ( $P=0.059$ ). The study showed that *E. coli* Nissle 1917 is equivalent to mesalamine in maintaining remission. A smaller study of 29 patients in 2009 by Miele et al<sup>56</sup> evaluated the use of VSL#3 in children with UC for both the induction and maintenance of remission over a 1-year period in conjunction with standard therapy. They observed higher remission rates in the group treated with VSL#3 compared to placebo (92.8% versus 36.4%,  $P<0.001$ ), although the mean duration for achieving induction with standard steroid therapy was not significantly different between the two groups. In 2011, Ishikawa et al<sup>59</sup> studied 41 patients, of whom 21 took a synbiotic preparation containing a *B. breve* strain (Yakult) as its probiotic component and galacto-oligosaccharide as the prebiotic component. The patients were followed for 1 year and evaluated for endoscopic improvement. The study reported improvement of endoscopic grading (Matt's Classification)<sup>62</sup> in the synbiotic group when compared to the standard therapy group.

## Pouchitis

Data regarding the effectiveness of probiotics for induction and maintenance therapy in pouchitis is very limited. There were no RCTs evaluating the use of prebiotics and synbiotics for management of pouchitis. A total of five RCTs using probiotics in pouchitis were found<sup>63-67</sup> (listed in Table 4). In all the studies, disease activity was measured by the Pouchitis Disease Activity Index (PDAI), which is an 18-point scale consisting of three categories: symptoms, endoscopic findings, and histological findings.<sup>68</sup> Four of the trials used VSL#3 as the probiotic agent<sup>63,65-67</sup> and one trial used only LGG.<sup>64</sup> Four of the five RCTs found probiotics to be effective in the maintenance of remission in pouchitis.<sup>63,65-67</sup> One trial found that there was no such effect.<sup>64</sup> The doses of culture selected for use were not uniform between these studies.

In 2000, Gionchetti et al<sup>63</sup> published a double-blinded, placebo-controlled trial that used probiotic VSL#3 as a maintenance treatment in patients with chronic relapsing pouchitis, defined as  $>3$  relapses per year. This study involved 40 patients who were in clinical and endoscopic remission after 1 month of antibiotic treatment with ciprofloxacin and rifaximin. Patients were randomized to receive 6 g daily of VSL#3 (1,800 billion bacteria/daily dose) or an identical placebo for 9 months. The patients were assessed clinically every month and endoscopically and histologically every

**Table 4** Probiotics in pouchitis

SL no	Publication (year, country)	First author	Type of intervention	Agent used	N	Control group	Study group	Primary end point	Benefit (P-value)	Harm
1	2000, Italy	Gionchetti <sup>63</sup>	Probiotic	VSL#3	40	20	20	Clinical relapse rate	Yes (<0.01)	Benefit
2	2003, Finland	Kuisma <sup>64</sup>	Probiotic	Lactobacillus GG	20	10	10	Clinical response, change in microbial flora	No (0.97)	None
3	2003, Italy	Gionchetti <sup>65</sup>	Probiotic	VSL#3	40	20	20	Clinical relapse rate	Yes (0.05)	Benefit
4	2004, UK	Mimura <sup>66</sup>	Probiotic	VSL#3	36	16	20	Remission maintenance rate	Yes (<0.0001)	Benefit
5	2008, Italy	Pronio <sup>67</sup>	Probiotic	VSL#3	31	13	18	Remission maintenance rate and inflammatory markers	Yes (0.006)	Benefit

**Note:** Randomized controlled trials that have studied the use of probiotic agents in pouchitis, with primary end points, number of patients involved in the study, and outcomes as shown.  
**Abbreviations:** N, total number of patients enrolled in the study at the beginning of the study; SL no, serial number.

2 months or in the event of a relapse. Relapse was defined as an increase of  $>2$  points in the clinical category of PDAI and was confirmed by endoscopic and histological findings. At the end of 9 months, 15% of the treatment group experienced a relapse while 100% in the placebo group experienced a relapse ( $P<0.01$ ). A subsequent double-blinded RCT in 2003 by Gionchetti et al<sup>65</sup> evaluated 40 patients for 1 year who were randomized to VSL#3 or placebo for disease maintenance in pouchitis. These patients underwent clinical, endoscopic, and histological assessment at 1 month, 3 months, 6 months, 9 months, and 12 months. Development of pouchitis occurred in 10% of those in the treatment group compared to 40% of those treated with placebo ( $P<0.05$ ). Additionally, the treatment group demonstrated significant improvement in quality of life, lower stool frequency, and increased concentrations of “protective bacteria”.

Pronio et al<sup>67</sup> conducted an open-labeled RCT involving 31 UC patients who had undergone colectomy with ileal pouch–anal anastomosis who had no signs or symptoms of pouchitis. These patients were randomized to receive either VSL#3 daily (450 billion bacteria/packet) or no treatment. They were evaluated clinically and endoscopically at baseline and after 3 months, 6 months, and 12 months. Patients with a total PDAI score  $>7$  were defined as having acute pouchitis. Patients treated with VSL#3 showed a small but statistically significant reduction in PDAI score at 3 months ( $P=0.014$ ) and maintained these scores at 6 months and 12 months, compared to no change in PDAI score in the placebo group. In addition, the study found that VSL#3 administration led to an expansion of mucosal regulatory T-cells that was speculated to have a role in controlling mucosal inflammation.

Mimura et al<sup>66</sup> conducted a double-blinded RCT involving 36 patients who had experienced pouchitis at least twice in 1 year or who required continuous antibiotics. Initially, remission was induced with oral antibiotic therapy consisting of metronidazole and ciprofloxacin for 4 weeks. The patients were randomized to receive either VSL#3 daily (300 billion bacteria/g) or placebo for 1 year or until relapse. Their PDAI score was evaluated before randomization, at 2 months, and at 12 months or until experiencing relapse. Remission was maintained in 85% of the treatment group compared to only 1% in the control group ( $P<0.0001$ ). Kuisma et al<sup>64</sup> conducted a double-blinded RCT that used LGG. In this study, 20 patients with a history of pouchitis were randomized to receive either two capsules of LGG twice daily ( $n=10$ ;  $[0.5-1.0] \times 10^{10}$  colony-forming units/capsule) or a placebo ( $n=10$ ) for 3 months. The effect was measured by fecal samples and mucosal biopsies taken for bacteriological analysis

and fecal bile acid composition before and after treatment. LGG was associated with altered pouch flora; however, no significant difference in PDAI scores was seen between the treatment and control group. There was no difference in clinical, endoscopic, or histological findings in the patients who were given the probiotic.

## Discussion

### Crohn's disease

There have been several studies evaluating the effect of probiotics in CD. Different probiotics have shown variable results with respect to clinical response when compared to a placebo or standard treatment. In a meta-analysis by Rahimi et al,<sup>69</sup> eight placebo-controlled RCTs were selected, of which six using *Lactobacillus* showed no superior benefit in maintaining remission and preventing clinical or endoscopic relapses. Interestingly, two studies that showed some clinical benefit did not use *Lactobacillus*; one used *Saccharomyces* and the other used *E. coli*. This clearly suggests that use of *Lactobacillus* as a single probiotic agent is of no benefit in CD. In our systematic review of the literature we have found a similar effect. After surgical resection, probiotics have been used to maintain remission among CD patients. A meta-analysis of seven studies by Doherty et al<sup>70</sup> in 2010 showed similar results as described above. Recently, Shen et al<sup>71</sup> published a systematic review with meta-analysis of RCTs that evaluated the effect of probiotics in IBD patients. They selected a total of 23 trials and performed a meta-analysis that showed therapeutic benefit with the use of probiotics among UC and pouchitis patients, but no such effect was noted in CD.

In our review there were a total of 14 RCTs that studied the use of probiotics, prebiotics, and/or synbiotics in CD.<sup>18-31</sup> Of these, four showed a benefit in outcome with use of intervention.<sup>18,20,27,28</sup> Three of the four used a probiotic<sup>18,20,27</sup> and one used a synbiotic in the study arm.<sup>28</sup> Three studies using *S. boulardii* showed a benefit with treatment; however these were limited by small study populations ranging from 17 to 31 patients per study (average 26.6).<sup>18,20,27</sup> A 2013 study using *S. boulardii* was performed in 165 patients by Bourreille et al.<sup>31</sup> This was a well-designed RCT that showed no statistically significant difference between use of placebo and the probiotic. Of the 14 RCTs, two additional studies showed no trend toward clinical benefit or harm.<sup>26,31</sup> There is a need for subsequent large, well-designed RCTs to evaluate the efficacy of other probiotic agents in CD.

The majority of the listed studies had significant limitations, including small study populations, high dropout rates, and lack of dose-response analyses. There was also a lack of

consistency with the probiotic agent, dose, and concomitant use of other medications that may affect outcomes, including antidiarrheals, aminosalicylates, immunomodulators, and corticosteroids. These studies also did not correlate CDAI scores with endoscopic remission. Therefore, most of these studies provide inconclusive evidence of a beneficial role of probiotics in the induction or maintenance of remission in CD. Well-designed RCTs are needed to determine if there is any benefit with the use of probiotics, prebiotics, or synbiotics in the treatment of CD.

## Ulcerative colitis

We reviewed 21 RCTs evaluating the use of various probiotics, prebiotics, and synbiotics in the management of UC. Overall, a myriad of different agents with varying doses makes drawing a conclusion regarding efficacy in achieving and/or maintaining remission in UC difficult. The limited size of the populations studied, differences in methodology, and duration of therapy limit the quality of some of the studies. The majority of studies (19/21) evaluating different agents for either induction and/or remission in mild to moderate UC showed at least a trend toward clinical benefit,<sup>30,36–43,50–57,59,60</sup> with 16 of the 19 studies showing statistically significant benefit.<sup>36–43,50–53,55,56,59,60</sup> A Cochrane review in 2007 evaluated the use of probiotics for the induction of remission in UC.<sup>72</sup> Four studies met criteria and the review concluded that probiotics combined with conventional therapy do not improve overall remission rates; however they may provide modest benefits in terms of reduction of disease activity in mild to moderate UC. The two largest studies in our review conducted after 2008 by Sood et al<sup>39</sup> and Tursi et al<sup>40</sup> support the use of VSL#3 in the induction of remission. The caveat is that VSL#3 was used as an adjunct to standard therapy, with greater than 90% of patients in the Tursi et al<sup>40</sup> trial on mesalamine and none of the patients receiving VSL#3 alone. In the Sood et al<sup>39</sup> trial, 96% of patients were on mesalamine, whether alone or in combination with other UC medications, with only 2.6% of patients receiving VSL#3 monotherapy. Currently there seems to be insufficient data to recommend for or against the use of other probiotics for the induction of remission in UC.

A Cochrane review by Naidoo et al<sup>73</sup> in 2011 evaluated the efficacy and safety of probiotics for the maintenance of remission in UC. Four studies met inclusion criteria and tested the effect of probiotics among 587 patients with UC in remission. The studies were either small or had methodological flaws. There was no clear benefit for probiotic treatment compared to either placebo or mesalamine. While

Kruis et al<sup>52</sup> and Zocco et al<sup>54</sup> were among the four studies included in the Cochrane review, they do provide the best evidence that there is a role for probiotics in maintenance of remission in UC. *E. Coli* Nissle 1917 and LGG, which were the two probiotics used in these studies, have been studied in other types of IBD, as noted elsewhere in this review, and according to these studies they appear safe. The findings of these studies need to be replicated in larger double-blinded RCTs. At this stage it is difficult to argue superiority of a specific probiotic. Both of these authors acknowledge the efficacy of various probiotic regimens. Zocco et al<sup>54</sup> state that the same probiotic should not be assumed to be “equally suitable for all individuals and the heterogeneity of clinical disorders such as UC suggests that strain-specific properties may be required for different patient categories”.

There are limited studies evaluating the use of prebiotics in UC. Our review included one study by Hafer's group<sup>30</sup> that showed no statistical benefit with use of prebiotic lactulose but did show a trend toward clinical benefit. Fujimori's group<sup>55</sup> conducted a head-to-head trial comparing the efficacy of probiotic, prebiotic, and synbiotic in UC. This study showed a superior benefit with the use of synbiotic compared to probiotic or prebiotic alone. In conclusion, larger well-designed RCTs are needed to further determine whether probiotics, type of probiotics, and/or synbiotics are of clear benefit for both the induction and/or the maintenance of remission in UC.

## Pouchitis

Since pouchitis has a similar pathogenic mechanism as UC, we expected a similar clinical benefit as seen in UC. Our review included five RCTs that evaluated the effect of probiotics in pouchitis (Table 4). In 2008, Elahi et al<sup>74</sup> published a systematic review and a meta-analysis of five placebo-controlled RCTs that showed clinical benefit, determined by PDAI, with the use of probiotics VSL#3 and LGG. Of the five trials selected in our review, four showed benefit in clinical outcome<sup>63,65–67</sup> and one showed no outcome benefit in terms of clinical and endoscopic findings.<sup>64</sup> This study used LGG culture as the probiotic agent. Among the four trials that showed benefit, the observed treatment effects were statistically significant, with some trials demonstrating an 85% remission rate in the VSL#3-treated groups.<sup>63,65–67</sup> The most significant limitation in all five studies was small sample sizes, with the largest being composed of 40 patients and the smallest consisting of 15 patients. Additional limitations include the short duration of the trials, with one being 3 months, another being 9 months, and the rest being 12 months. This warrants

subsequent larger double-blinded RCTs with a much longer duration of follow-up.

## Future directions: fecal microbiota transplantation

Fecal microbiota transplantation (FMT) is the infusion of a fecal suspension from a healthy individual into the gastrointestinal tract of another person in an attempt to treat an illness. The process has been described for decades, with the first reported case in humans performed by Eiseman and colleagues<sup>75</sup> in 1958 for pseudomembranous colitis. FMT has primarily been used to treat *C. difficile* infection, with cure rates of up to 95% in published studies.<sup>76</sup> This novel technique began to be employed because of a continual rise in *C. difficile* infections since the year 2000, especially in patients with recurrent infections that failed standard antibiotics therapy, in theory because of bowel flora deficiencies. Given the reduced diversity of luminal microbiota in IBD and specific deficiencies in *Bacteroidetes* and *Firmicutes* in IBD patients, FMT offers an attractive solution to replenish those species.<sup>9</sup> While probiotics may alter the metabolic or immunologic activity of residing native gut microbiota with a cultured single or a few strains, FMT seems to result in durable, long-term implantation of donor flora.<sup>77</sup> Borody et al<sup>78</sup> found marked success with FMT in six UC patients, causing reversal of inflammation in those cases. These cases required multiple infusions rather than the single episode of FMT that is typically done for *C. difficile* infection. A systematic review by Anderson et al<sup>79</sup> in 2012 found no RCTs, but reviewed 17 case reports or case series for a total of 26 patients receiving FMT for management of their IBD. Of those, 27 patients had UC, 12 had CD, and two were unclassified. Unfortunately, there was variability and incomplete data for the preparation and administration of FMT. No article reported complete data enabling protocol replication.<sup>79</sup> Variability was seen with the number of follow-up enemas – from 2 days to 59 weeks. There was also no standardization for reporting disease activity in most series. Twenty five patients had follow-up information and 19 of these reported complete resolution of their symptoms. Disease activity was reported only in 24 patients and 15 patients were in remission (15/24). Due to lack of standardized studies, clearly more rigorous studies are needed. More recently in 2013 Kunde et al<sup>80</sup> performed a single-center uncontrolled study of FMT in 10 children with UC, showing safety, tolerability, and clinical response. They used a standardized Pediatric Ulcerative Colitis Activity Index<sup>81</sup> to document clinical response and remission; the

first study of its kind. Adverse events were similar to other studies and were reported as bloating, flatulence, abdominal pain, cramping, diarrhea, fatigue, fever, and bloody stools. One subject had disabling hematochezia in the third week after FMT, but it was thought to be due to a UC flare rather than temporally relating to the FMT.

Given the findings above, the evidence for use of FMT is limited and weak overall, with most data based on case reports. Although limited, this evidence shows promise that FMT may be an effective and safe alternative treatment for IBD patients. Certainly, FMT holds enormous potential for a wide range of diseases. Future directions should include fecal and mucosal microbial community profiling, as this would help determine whether FMT's mechanism of action in IBD is similar to that in *C. difficile* infection. Standardization of FMT preparation, donor selection, route of administration, and duration of therapy with FMT are all needed and should be established through double-blinded RCTs.

## Conclusions

All of the Crohn's disease studies had significant limitations. The size of the study population in the majority of the studies was small. These studies were plagued by high dropout rates and lack of dose-response analyses. There was also lack of consistency with respect to concomitant use of other medications, including antidiarrheals, aminosalicylates, immunomodulators, and corticosteroids, which could have affected outcomes. None of these studies provide conclusive evidence of a beneficial role of probiotics in induction or maintenance of remission in CD. Larger RCTs are needed to show any benefit for the use of probiotics, prebiotics, or synbiotics for treatment of CD.

In the ulcerative colitis studies, various agents showed a trend toward improved rates in both induction of remission and maintenance in UC. The effect was noted both clinically and objectively on a macroscopic and molecular level, in conjunction with standard treatment. For pouchitis, the evidence appears to support a role for VSL#3 as a maintenance therapeutic option when pouchitis is in remission. However, the limited data available do not support the use of probiotics as primary therapy for pouchitis.

## Disclosure

The authors report no conflicts of interest in this work.

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