Research Article

A Systematic Review and Meta-Analysis of Randomized Controlled Trials of Fecal Microbiota Transplantation for the Treatment of Inflammatory Bowel Disease

Xi-Yue Tan, Yu-Jia Xie, Xing-Long Liu, Xin-Yun Li, and Bo Jia

College of Basic Medicine, Chengdu University of Traditional Chinese Medicine, Chengdu 611137, China

Correspondence should be addressed to Bo Jia; jiabochengdutcm@163.com

Received 20 April 2022; Accepted 6 June 2022; Published 26 June 2022

Academic Editor: Duygu AĞAGÜNDÜZ

Copyright © 2022 Xi-Yue Tan et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Objectives. Inflammatory bowel disease (IBD) is a chronic recurrent inflammatory disease of the gastrointestinal tract, and its prevalence is increasing worldwide. Fecal microbiota transplantation (FMT) is an emerging therapy that modifies the patient’s gut microbiota by transplanting feces from a healthy donor to achieve disease remission. However, its efficacy and safety need to be further investigated.

Methods. PubMed, the Cochrane Library, Web of Science, Embase, and Google Scholar databases (up to 8th November 2021) were searched and literature was screened by title and abstract as well as full text. The primary outcome was clinical remission, with the clinical response as a secondary outcome. Risk ratios (RR) with 95% confidence intervals (CI) were reported.

Results. A total of 14 trials were included in this study. In terms of clinical remission, FMT had a significant effect compared to placebo (RR = 1.44, 95 CI%: 1.03 to 2.02, I² = 38%, P = 0.03), with no significant risk of study heterogeneity. Moreover, FMT led to significant results in clinical response compared to placebo with moderate between-study heterogeneity (RR = 1.34, 95 CI%: 0.92 to 1.94, I² = 51%, P = 0.12). Subgroup analysis showed a higher clinical remission for fresh fecal FMT (40.9%) than that for frozen fecal FMT (32.2%); the efficacy of gastrointestinal (GI) pretreatment, the severity of disease, route of administration, and the donor selection remain unclear and require more extensive study. Safety analysis concluded that most adverse events were mild and self-resolving. The microbiological analysis found that the patient’s gut microbiota varied in favor of the donor, with increased flora diversity and species richness. Conclusion. FMT is a safe, effective, and well-tolerated therapy. Studies have found that fresh fecal microbiota transplant can increase clinical remission rates. However, more randomized controlled trials and long-term follow-ups are needed to assess its long-term effectiveness and safety.

1. Introduction

Inflammatory bowel disease (IBD), which includes Crohn’s disease (CD) and ulcerative colitis (UC), is a chronic recurrent inflammatory disease of the gastrointestinal tract [1]. The prevalence of IBD is increasing worldwide [2, 3]. The annual incidence of UC and CD in North America is similar, 0–19.2 per 100,000 and 0–20.2 per 100,000, respectively [2]. UC is characterized by diffuse mucosal inflammation, with lesions limited to the colonic mucosa and typical symptoms of diarrhea, mucus, and bleeding; CD can cause transmural inflammation that can accumulate anywhere in the gastrointestinal tract, most commonly in the terminal ileum or rectum, with abdominal pain, watery diarrhea, and weight loss as the main clinical features [1, 4, 5]. The etiology of IBD is uncertain, but it is now believed that genetic susceptibility, environmental exposure, gut microbiota, and the immune system are involved. Current treatment is aimed at controlling the progression of inflammation and achieving disease remission. Commonly used drugs are corticosteroids, antitumor necrosis factor-alpha (TNF-α) drugs, immunomodulators (e.g., methotrexate), biologic therapies, and surgery. IBD not only imposes a heavy financial burden and reduces the quality of life but also has a significant impact on the resources of the healthcare system [6, 7].

Fecal microbiota transplantation (FMT) is a therapy in which fecal matter from a healthy donor is infused into the gastrointestinal tract of a recipient patient, changing the gut microbiota to promote disease remission. FMT has been used to treat severe and refractory cases of IBD, with documented improvements in clinical symptoms and drug-refractory conditions. This therapy has been shown to be safe and effective in reducing inflammation, improving symptoms, and achieving disease remission.
patient’s intestinal microbiota, leading to remission [8]. It has been suggested that the development of chronic inflammatory diseases such as IBD is associated with alterations in the composition of the intestinal microbiota (known as ecological dysbiosis) [9]. The differences in gut microbiota composition between IBD patients and healthy individuals are mainly in terms of microbial diversity and flora richness [10]. Extensive data confirm FMT as an effective treatment for recurrent Clostridium difficile infection (rCDI) [11, 12]. European Microbiology, the European Society for Infectious Diseases, and the American College of Gastroenterology all recommend FMT as a treatment for rCDI and have included it in their treatment guidelines [13, 14]. FMT is currently considered an emerging treatment for IBD [15, 16]. Studies have confirmed the efficacy of FMT in the treatment of IBD; Fang et al. [17] reported a clinical remission rate of 28.8% and a clinical response rate of 53% during follow-up in patients with IBD, concluding that fresh or frozen donor stool, route of administration, and antibiotic pretreatment or not had no effect on the efficacy of FMT for IBD. Caldeira et al. [18] showed an overall clinical remission rate of 37% and an overall clinical response rate of 54% in patients with IBD, suggesting that frozen fecal material was associated with a higher rate of clinical remission. However, several issues remain unresolved in the FMT protocol, such as type of stool (fresh or frozen), route of administration (lower or upper gastrointestinal tract), donor selection (healthy anonymous, or relative, or partners), and whether to pretreat the gastrointestinal (GI) tract (i.e., using vancomycin and/or polyethylene glycol, or antibiotic, or macrogol solution) prior to treatment with FMT.

Therefore, the purpose of this systematic review and meta-analysis is to synthesize data from published randomized controlled trials on FMT for the treatment of patients with IBD and to evaluate the efficacy and safety of FMT for the treatment of IBD.

2. Methods

This study was conducted and reported in accordance with the Preferred Reporting Items for Systematic Evaluation and Meta-Analysis (PRISMA (available (here))) guidelines.

2.1. Search Methodology. English articles of randomized controlled trials were searched in PubMed, the Cochrane Library, Web of Science, Embase, and Google Scholar databases (up to 8th November 2021). The databases were used (“Fecal Microbiota Transplantation” OR “Microbiota Transplantation, Fecal” OR “Transplantation, Fecal Microbiota” OR “Bacteriotherapy” OR “Intestinal Microbiota Transfer”) AND (“IBD” OR “inflammatory bowel disease” OR “Crohn’s disease” OR “CD” OR “ulcerative colitis” OR “UC” OR “colitis”) for the keywords combinations and wildcards were thoroughly searched to ensure the completeness of the search results.

2.2. Inclusion and Exclusion Criteria. Studies of randomized controlled trials of patients with IBD treated with FMT were screened by a set of inclusion and exclusion criteria. Inclusion criteria were as follows: (1) patients with IBD aged > 18 years with no gender restriction; (2) randomized controlled trials; (3) studies with clearly described clinical endpoints; (4) journal articles; and (5) FMT by different modalities (i.e., nasogastric tube, nasoduodenal tube, colon/colonoscopy, enema, or oral) was allowed. Exclusion criteria were as follows: (1) in vitro or animal testing; (2) languages other than English or Chinese; (3) patients with concomitant Clostridium difficile infection (CDI) infection or other pathogens; (4) population noncompliance, e.g., pediatric IBD; (5) duplicate literature; (6) other types of literature: reviews, editorials, case reports, abstracts, meta-analysis, observational studies, surveys, and questionnaires; (7) no clinical results reported, or studies with incomplete data.

We went through the PRISMA flowchart to select studies for inclusion. One reviewer performed the initial screening based on titles and abstracts. Titles and abstracts of potentially eligible records were independently reviewed by a second reviewer. Identified studies were then independently screened in full by two reviewers. Final inclusion was determined by both reviewers based on inclusion and exclusion criteria. The senior author discussed and resolved any discrepancies in the selection process.

2.3. Data Extraction and Quality Assessment. Data were extracted and evaluated independently by two reviewers, recorded on self-designed forms, and any disagreements were discussed and resolved by a third reviewer. Extracted study data included (1) first author, study year, and site; (2) participant characteristics: total number of participants, number of controls, number of experiments, and disease severity; (3) type of intervention: type of stool (fresh or frozen), route of administration, relationship to donor, gastrointestinal (GI) pretreatment, IBD type, and follow-up time; (4) clinical outcomes: primary outcome (clinical remission) and secondary outcome (clinical response).

Study quality was assessed using the Cochrane Bias Tool and risks of bias included: selection bias, performance bias, detection bias, attrition bias, reporting bias, and other sources of bias.

2.4. Statistical Analysis. The efficacy of FMT for IBD will be assessed based on clinical remission or clinical response as defined by the respective study authors. Safety will be assessed by study-reported adverse events and serious adverse events. The analysis will be performed using Review Manager, version 5.1. Results were expressed as risk ratios (RR) and 95% confidence interval (CI). Mantel–Haenszel statistics (M-H) were used to represent the results. The heterogeneity of the study results was evaluated graphically using the inconsistency relative index ($I^2$), and the $I^2$ statistic ($I^2 \geq 50\%$ using a random-effects model and $I^2 < 50\%$ using a fixed-effects model) was used to evaluate statistical heterogeneity [19], and $P$ value $< 0.05$ was considered statistically significant. In addition, subgroup analysis was performed according to disease severity, route of administration, type of stool storage (fresh/frozen), GI pretreatment, IBD type, gender, and donor-
3.1. Literature Search. A total of 5941 studies, including 5459 plots and Egger’s test. Sensitivity analysis was performed using STATA 16.0 software. To assess the stability of the results, a patient relationship. To assess the stability of the results, sensitivity analysis was performed using STATA 16.0 software. Potential publication bias was assessed using funnel plots and Egger’s test.

3. Results

3.1. Literature Search. A total of 5941 studies, including 5459 foreign language studies, and 482 Chinese language studies, were obtained by searching the database using the search strategy. The journal article title information of these studies was imported into the Endnote software. After removing duplicates, the search in the database yielded 3555 records. During the title and abstract screening process, 3391 records were considered irrelevant, and subsequently, 164 studies were evaluated after full-text assessment, and 150 studies were excluded using predetermined exclusion criteria. Ultimately, 14 studies were obtained, including 12 studies in English and 2 studies in Chinese. All included studies were published as journal articles. The study selection flowchart is shown in Figure 1. The data from these studies are presented in Table 1.

The 14 studies included 12 UC studies [20–23, 25, 26, 28–33] and 2 CD studies [24, 27] that included a total of 666 patients. FMT was the primary intervention in the RCTs trial group. In contrast, the control group took different measures, with 1 studying different routes of administration by gastroscopy or colonoscopy [27], 7 using placebo controls (including 2 autologous stools [20, 23], 2 sham procedures [24, 28], and 3 isotonic saline [21, 22, 25]), 1 using a special UC diet [30], 4 choosing conventional treatment as a control [20–22, 24, 25, 26, 28–30], and FMT led to significant results compared to placebo with moderate study heterogeneity. Eight studies reported clinical response [20–23, 26, 28, 30, 31], and FMT led to significant results in clinical response compared to placebo with moderate between-study heterogeneity (RR = 1.34, 95 CI%: 0.92 to 1.94, P = 0.12). Forest plots are shown in Figures 3 and 4.

3.2. Bias Analysis of Included Studies. Figures 2(a) and 2(b) summarizes the risk of bias for randomized controlled trials. The risk of bias assessment showed that 6 studies reported the method of randomization, of which 5 studies used computerized randomization as low risk [20–22, 25, 28], 1 study chose randomization according to the order of admission as high risk [33], and the remaining studies did not inform the specific grouping method. Seven studies took participant and researcher allocation concealment [20–22, 24, 25, 28, 30], 6 studies used participant and evaluator blinding [20–23, 25, 28], and the rest were not mentioned. Follow-up was not mentioned in only 1 study [33]. The remaining trials clearly reported the number of missed visits and dropouts and the reasons for them and adequately addressed the issue of selective reporting bias. However, given the poor methodological quality, we noted an unclear risk of bias in most of the included trials.

3.3. Clinical Remission/Response of FMT for IBD. Eleven studies reporting clinical remission in FMT for IBD (438 UC patients and 17 CD patients) were included in our analysis [20–26, 28–31]. In terms of clinical remission, FMT had a significant effect compared to placebo (RR = 1.44, 95 CI%: 1.03 to 2.02, I² = 38%, P = 0.03), with no significant risk of study heterogeneity. Eight studies reported clinical response [20–23, 26, 28, 30, 31], and FMT led to significant results in clinical response compared to placebo with moderate between-study heterogeneity (RR = 1.34, 95 CI%: 0.92 to 1.94, I² = 51%, P = 0.12). Forest plots are shown in Figures 3 and 4.

3.4. Subgroup Analysis

3.4.1. Disease Severity. Of 455 patients, 266 were defined as having mild-moderate disease, of which 82 achieved clinical remission, and 172 were described as having active disease (including refractory UC and recurrent UC), of which 79 achieved clinical remission. The clinical remission rate for mild-moderate IBD was 30.8% (RR = 1.31, 95 CI%: 0.68 to 2.54, I² = 56%), with moderate study heterogeneity. In comparison, the clinical remission rate for active IBD was 45.9% (RR = 1.47, 95 CI%: 0.94, 2.29, I² = 37%), with a low risk of study heterogeneity. No statistically significant differences were observed between these two studies (P = 0.14) (Figure 5).

3.4.2. Type of Stool. Of the 445 patients, 202 patients used frozen stools (65 patients achieved clinical remission), 105 patients used fresh stools (43 patients achieved clinical remission), 136 patients used frozen or fresh stools (58 patients achieved clinical remission), and 12 patients used frozen stools and oral capsules (2 patients achieved clinical remission). Clinical remission rates were 32.2% with frozen stool, 40.9% with fresh stool, 42.6% with frozen or fresh stool, and 16.7% with frozen stools and oral capsules (Figure 6). No

---

Figure 1: Selection flowchart.
<table>
<thead>
<tr>
<th>Clinical response</th>
<th>Clinical remission</th>
<th>Follow-up</th>
<th>Study</th>
<th>Study type</th>
<th>Country</th>
<th>N</th>
<th>Severity</th>
<th>Donor</th>
<th>Route</th>
<th>Fresh/frozen</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>dFMT:</strong> 18/38 (47%); aFMT: 6/35 (17%); at week 8</td>
<td><strong>dFMT:</strong> 21/38 (55%); aFMT: 8/35 (23%); at week 8</td>
<td>8 weeks</td>
<td>Costello et al. [20]</td>
<td>UC Australia</td>
<td>73</td>
<td>Mild to moderate</td>
<td>Unrelated</td>
<td>Colonoscopy and enema (lower)</td>
<td>Frozen</td>
<td></td>
</tr>
<tr>
<td>FMT: 9 (24%); placebo: 2 (5%) at week 7</td>
<td>FMT: 15 (39%); placebo: 9 (24%) at week 7</td>
<td>6 weeks</td>
<td>Moayyedi et al. [21]</td>
<td>UC Canada</td>
<td>75</td>
<td>Active</td>
<td>Unrelated</td>
<td>Retention enema (lower)</td>
<td>Fresh or frozen</td>
<td></td>
</tr>
<tr>
<td>FMT: 18 (44%); placebo: 8 (20%) at week 8</td>
<td>FMT: 22 (54%); placebo: 9 (23%) at week 8</td>
<td>8 weeks</td>
<td>Paramsothy et al. [22]</td>
<td>UC Australia</td>
<td>81</td>
<td>Mild to moderate</td>
<td>Unrelated</td>
<td>Colonoscopy and enema (lower)</td>
<td>Frozen</td>
<td></td>
</tr>
<tr>
<td>dFMT: 6/38 (26.1%); aFMT: 8/25 (32%), at week 12</td>
<td>dFMT: 10/23 (43.5%); aFMT: 25/23 (30.4%); at week 12</td>
<td>12 weeks</td>
<td>Rossen et al. [23]</td>
<td>UC Netherlands</td>
<td>48</td>
<td>Mild to moderate</td>
<td>Relatives, partners, or volunteers</td>
<td>Nasoduodenal tube (upper)</td>
<td>Fresh</td>
<td></td>
</tr>
<tr>
<td>FMT: 7/8 (87.5%); sham FMT: 4/9 (44.4%), at week 10; FMT: 4/8 (50.0%); sham FMT: 3/9 (33.3%), at week 24</td>
<td>—</td>
<td>12 weeks</td>
<td>Sokol et al. [24]</td>
<td>UC France</td>
<td>17</td>
<td>Active</td>
<td>Unrelated</td>
<td>Colonoscopy (lower)</td>
<td>Fresh</td>
<td></td>
</tr>
<tr>
<td>FMT: 20/30 (66.7%) at week 48</td>
<td>—</td>
<td>48 weeks</td>
<td>Sood et al. [25]</td>
<td>UC India</td>
<td>61</td>
<td>Active</td>
<td>Unrelated</td>
<td>Colonoscopy infusion and enema (lower)</td>
<td>Fresh or frozen</td>
<td></td>
</tr>
<tr>
<td>FMT: 3/10 (30%); FMT: 4/10 (40%) at week 12; Gastroscopy: 10/13 (76.9%); colonoscopy: 11/14 (78.6%); at week 2</td>
<td>FMT: 7/10 (70%); FMT: 6/10 (60%) at week 12; Gastroscopy: 9/13 (69.2%); colonoscopy: 9/14 (64.3%) at week 2</td>
<td>12 weeks</td>
<td>Wei et al. [26]</td>
<td>UC China</td>
<td>20</td>
<td>Mild to moderate</td>
<td>Unrelated</td>
<td>Colonoscopy (lower)</td>
<td>Fresh</td>
<td></td>
</tr>
<tr>
<td>FMT: 2/6 (33%); placebo: 0/6 (0%) at week 12</td>
<td>cFMT: 3/6 (50%); placebo: 1/6 (17%) at week 12</td>
<td>8 weeks</td>
<td>Yang et al. [27]</td>
<td>CD China</td>
<td>27</td>
<td>Active</td>
<td>Relatives, partners, or volunteers</td>
<td>Colonoscopy infusion and oral (lower or upper)</td>
<td>Fresh</td>
<td></td>
</tr>
<tr>
<td>FMT: 9/10 (90%); control: 5/10 (50%) at week 8</td>
<td>—</td>
<td>6–38 months</td>
<td>Crothers et al. [28]</td>
<td>UC USA</td>
<td>17</td>
<td>Mild to moderate</td>
<td>Unrelated</td>
<td>Colonoscopy (lower)</td>
<td>Frozen and capsules</td>
<td></td>
</tr>
<tr>
<td>FMT: 2/17 (11.8%); FMT + UCED: 4/19 (21.1%); UCED: 6/15 (40.0%) at week 8</td>
<td>FMT: 6/17 (35.3%); FMT + UCED: 8/19 (42.1%); UCED: 9/15 (60.0%) at week 8</td>
<td>12 weeks</td>
<td>Fang et al. [29]</td>
<td>UC China</td>
<td>20</td>
<td>Active</td>
<td>Unrelated</td>
<td>Colonoscopy (lower)</td>
<td>Frozen</td>
<td></td>
</tr>
<tr>
<td>FMT: 3/8 (37.5%); 5-ASA: 4/8 (50%) at week 12</td>
<td>FMT: 5/8 (62.5%); 5-ASA: 5/8 (62.5%) at week 12</td>
<td>12 weeks</td>
<td>Sarbagali Shabat et al. [30]</td>
<td>UC Israel</td>
<td>51</td>
<td>Mild to moderate</td>
<td>Unrelated</td>
<td>Colonoscopy and enema (lower)</td>
<td>Frozen</td>
<td></td>
</tr>
<tr>
<td>FMT: 20/30 (66.7%); control: 12/30 (40.0%) at week 4</td>
<td>FMT: 5/8 (62.5%); 5-ASA: 5/8 (62.5%) at week 12</td>
<td>12 weeks</td>
<td>Schierová et al. [31]</td>
<td>Czech republic</td>
<td>16</td>
<td>Active</td>
<td>Unrelated</td>
<td>Enema (lower)</td>
<td>Frozen</td>
<td></td>
</tr>
<tr>
<td>FMT: 48/50 (96%); 5-ASA: 35/50 (70%) at week 2</td>
<td>FMT: 5/8 (62.5%); 5-ASA: 5/8 (62.5%) at week 12</td>
<td>3 months</td>
<td>Chen et al. [32]</td>
<td>UC China</td>
<td>60</td>
<td>NR</td>
<td>Unrelated</td>
<td>Enema (lower)</td>
<td>Fresh or frozen</td>
<td></td>
</tr>
<tr>
<td>—</td>
<td>—</td>
<td>—</td>
<td>Zhang et al. [33]</td>
<td>UC China</td>
<td>100</td>
<td>NR</td>
<td>Relatives, partners</td>
<td>Enema (lower)</td>
<td>NR</td>
<td></td>
</tr>
</tbody>
</table>

UC, ulcerative colitis; CD, Crohn’s disease; N, number of patients; aFMT, autologous FMT; dFMT, donor FMT; cFMT, capsules FMT; FMTP, FMT combined with pectin; FMT, fecal microbiota transplantation; 5-ASA, 5-aminosalicylic acid; UCED, ulcerative colitis exclusion diet; NR, not reported.
A statistically significant difference was observed between the type of stool groups for this analysis ($P = 0.10$).

3.4.3. Route of Administration. Of the 443 patients, 395 patients were treated through the lower GI tract, 151 of whom achieved clinical remission, and 48 patients were treated through the upper GI tract, 15 of whom achieved clinical remission. The clinical remission rate was 38.2% for patients treated with the lower GI tract and 31.2% for those treated with the upper GI tract (Figure 7). No statistically significant differences were found between routes of administration in this analysis ($P = 0.09$).
3.4.4. Donor Selection. Of the 354 patients, 306 patients used stool from healthy anonymous donors, of whom 124 achieved clinical remission, and 48 patients used stool from relatives or close friends or volunteers, of whom 15 achieved clinical remission. The clinical remission rate was 37.6% with stools from healthy anonymous donors and 31.2% with stools from healthy relatives, partners, or volunteers (Figure 8). No statistically significant differences were found between these two studies (P = 0.10).

3.4.5. Gastrointestinal Pretreatment. Of the 455 patients, 214 patients underwent GI pretreatment, of which 95 patients achieved clinical remission, and 241 patients did not undergo GI pretreatment, of which 73 patients achieved clinical remission. The clinical remission rate was 44.4% with GI pretreatment (RR = 1.39, 95 CI%: 0.90 to 2.14, I² = 32%) and 30.3% without GI pretreatment (RR = 1.45, 95 CI%: 0.81 to 2.58, I² = 49%) (Figure 9). No statistically significant differences were found between these two studies (P = 0.10).

3.4.6. IBD Type. Of the 455 patients, 438 were UC patients, of which 161 patients achieved clinical remission, and 17 were CD patients, of which 7 patients achieved clinical remission. The clinical remission rate was 36.8% had UC

---

### Table 1: Forest plot of studies evaluating induction of clinical response post-FMT for IBD. FMT, fecal microbiota transplantation; IBD, inflammatory bowel disease.

<table>
<thead>
<tr>
<th>Study or Subgroup</th>
<th>Experimental Events</th>
<th>Total</th>
<th>Control Events</th>
<th>Total</th>
<th>Weight (%)</th>
<th>Risk Ratio</th>
<th>Risk Ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>M-H, Fixed, 95% CI</td>
<td>M-H, Fixed, 95% CI</td>
</tr>
<tr>
<td>Crothers 2021 [28]</td>
<td>3</td>
<td>6</td>
<td>1</td>
<td>6</td>
<td>3.2</td>
<td>3.00</td>
<td>[0.42, 21.30]</td>
</tr>
<tr>
<td>Moayyedi 2015 [21]</td>
<td>15</td>
<td>38</td>
<td>9</td>
<td>37</td>
<td>13.5</td>
<td>1.62</td>
<td>[0.81, 3.24]</td>
</tr>
<tr>
<td>Paramsothy 2017 [22]</td>
<td>22</td>
<td>41</td>
<td>9</td>
<td>40</td>
<td>14.5</td>
<td>2.38</td>
<td>[1.26, 4.53]</td>
</tr>
<tr>
<td>Rossen 2015 [23]</td>
<td>11</td>
<td>23</td>
<td>13</td>
<td>25</td>
<td>15.9</td>
<td>0.92</td>
<td>[0.52, 1.63]</td>
</tr>
<tr>
<td>Sarbagili Shabat 2021 [30]</td>
<td>6</td>
<td>17</td>
<td>9</td>
<td>15</td>
<td>12.3</td>
<td>0.59</td>
<td>[0.27, 1.26]</td>
</tr>
<tr>
<td>Schierova 2020 [31]</td>
<td>5</td>
<td>8</td>
<td>5</td>
<td>8</td>
<td>12.4</td>
<td>1.00</td>
<td>[0.47, 2.14]</td>
</tr>
<tr>
<td>Wei 2016 [26]</td>
<td>7</td>
<td>10</td>
<td>6</td>
<td>10</td>
<td>14.3</td>
<td>1.17</td>
<td>[0.61, 2.23]</td>
</tr>
<tr>
<td>Total (95% CI)</td>
<td>181</td>
<td>176</td>
<td>100.0</td>
<td>1.34</td>
<td>[0.92, 1.94]</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Figure 4:** Forest plot of studies evaluating induction of clinical response post-FMT for IBD. FMT, fecal microbiota transplantation; IBD, inflammatory bowel disease.

---

### Table 2: Forest plot of disease severity.

<table>
<thead>
<tr>
<th>Study or Subgroup</th>
<th>Experimental Events</th>
<th>Total</th>
<th>Control Events</th>
<th>Total</th>
<th>Weight (%)</th>
<th>Risk Ratio</th>
<th>Risk Ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>M-H, Fixed, 95% CI</td>
<td>M-H, Fixed, 95% CI</td>
</tr>
<tr>
<td>4.5.1 Mild to moderately</td>
<td>Costello 2018 [20]</td>
<td>18</td>
<td>38</td>
<td>6</td>
<td>35</td>
<td>11.8</td>
<td>2.76</td>
</tr>
<tr>
<td></td>
<td>Crothers 2021 [28]</td>
<td>2</td>
<td>6</td>
<td>0</td>
<td>6</td>
<td>1.3</td>
<td>5.00</td>
</tr>
<tr>
<td></td>
<td>Paramsothy 2017 [22]</td>
<td>18</td>
<td>41</td>
<td>8</td>
<td>40</td>
<td>13.8</td>
<td>2.20</td>
</tr>
<tr>
<td></td>
<td>Rossen 2015 [23]</td>
<td>7</td>
<td>23</td>
<td>8</td>
<td>25</td>
<td>11.0</td>
<td>0.95</td>
</tr>
<tr>
<td></td>
<td>Sarbagili Shabat 2021 [30]</td>
<td>2</td>
<td>17</td>
<td>6</td>
<td>15</td>
<td>Not estimable</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Wei 2016 [26]</td>
<td>3</td>
<td>10</td>
<td>4</td>
<td>10</td>
<td>6.3</td>
<td>0.75</td>
</tr>
<tr>
<td>Subtotal (95% CI)</td>
<td>118</td>
<td>116</td>
<td>44.2</td>
<td>1.65</td>
<td>[0.96, 2.86]</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Figure 5:** Forest plot of disease severity.

<table>
<thead>
<tr>
<th>Study or Subgroup</th>
<th>Experimental Events</th>
<th>Total</th>
<th>Control Events</th>
<th>Total</th>
<th>Weight (%)</th>
<th>Risk Ratio</th>
<th>Risk Ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>4.5.2 Active</td>
<td>Fang 2021 [29]</td>
<td>9</td>
<td>10</td>
<td>5</td>
<td>10</td>
<td>15.3</td>
<td>1.80</td>
</tr>
<tr>
<td></td>
<td>Moayyedi 2015 [21]</td>
<td>9</td>
<td>38</td>
<td>2</td>
<td>37</td>
<td>4.6</td>
<td>4.38</td>
</tr>
<tr>
<td></td>
<td>Schierova 2020 [31]</td>
<td>3</td>
<td>8</td>
<td>4</td>
<td>8</td>
<td>7.0</td>
<td>0.75</td>
</tr>
<tr>
<td></td>
<td>Sood 2019 [25]</td>
<td>727</td>
<td>31</td>
<td>20</td>
<td>30</td>
<td>29.0</td>
<td>1.31</td>
</tr>
<tr>
<td>Subtotal (95% CI)</td>
<td>87</td>
<td>35</td>
<td>55.8</td>
<td>1.47</td>
<td>[0.94, 2.29]</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Figure 5:** Forest plot of disease severity.
3.4.1 Frozen
Costello 2018 [20] 18 38 6 35 10.8 2.76 [1.24, 6.16]
Paramsothy 2017 [22] 18 41 8 40 12.5 2.20 [1.08, 4.46]
Sarbagili Shabat 2021 [30] 2 17 6 15 4.5 0.29 [0.07, 1.24]
Schierowo 2020 [31] 3 8 4 8 6.7 0.75 [0.24, 2.33]
Subtotal (95% CI) 104 98 34.5 1.26 [0.53, 3.01]
Total events 41 24
Heterogeneity: $Tau^2 = 0.52$; $Chi^2 = 9.57$, df = 3 ($P = 0.02$); $I^2 = 69$
Test for overall effect: $Z = 0.53$ ($P = 0.60$)

4.1.2 Fresh
Fang 2021 [29] 9 10 5 10 13.6 1.80 [0.94, 3.46]
Rossen 2015 [23] 7 23 8 25 10.2 0.95 [0.41, 2.21]
Sokol 2020 [24] 2 8 3 9 6.5 1.50 [0.47, 4.76]
Wei 2016 [26] 3 10 4 10 6.0 0.75 [0.22, 2.52]
Subtotal (95% CI) 51 54 36.3 1.31 [0.85, 2.04]
Total events 23 20
Heterogeneity: $Tau^2 = 0.00$; $Chi^2 = 2.47$, df = 3 ($P = 0.48$); $I^2 = 0$
Test for overall effect: $Z = 1.22$ ($P = 0.22$)

4.1.3 Frozen or fresh
Moayyedi 2015 [21] 9 38 2 37 4.4 4.38 [1.01, 18.94]
Sood 2019 [25] 27 31 20 30 23.6 1.31 [0.98, 1.74]
Subtotal (95% CI) 69 67 28.0 2.05 [0.52, 8.03]
Total events 36 22
Heterogeneity: $Tau^2 = 0.75$; $Chi^2 = 3.60$, df = 1 ($P = 0.06$); $I^2 = 72$
Test for overall effect: $Z = 1.03$ ($P = 0.30$)

4.1.4 Frozen and capsules
Crothers 2021 [28] 2 6 0 6 1.3 5.00 [0.29, 86.43]
Subtotal (95% CI) 6 6 1.3 5.00 [0.29, 86.43]
Total events 2 0
Heterogeneity: Not applicable
Test for overall effect: $Z = 1.11$ ($P = 0.27$)

Total (95% CI) 230 225 100.0 1.44 [1.03, 2.02]
Total events 102 66
Heterogeneity: $Tau^2 = 0.10$; $Chi^2 = 16.08$, df = 10 ($P = 0.10$); $I^2 = 38$
Test for overall effect: $Z = 2.15$ ($P = 0.03$)
Test for subgroup differences: $Chi^2 = 1.19$, df = 3 ($P = 0.76$); $I^2 = 0$

Figure 6: Forest plot of type of stool.

(RR = 1.44, 95 CI%: 1.00 to 2.07, $I^2 = 44$%) and 41.2% had CD (RR = 1.50, 95 CI%: 0.47 to 4.76) (Figure 10). No statistically significant differences were found between these two studies ($P = 0.10$).

3.4.7. Gender. Of the 407 patients, 221 were male and 186 were female. The percentage of males was 54.1% (RR = 0.84, 95 CI%: 0.56 to 1.25, $I^2 = 0$%) and 45.7% (RR = 1.36, 95 CI%: 0.91 to 2.03) for females (Figure 11). No statistically significant differences were found between these two studies ($P = 0.76$).

3.5. Safety and Adverse Events. Adverse events during and after treatment with FMT were reported in the vast majority of studies, with only one study not reporting follow-up. Self-limiting symptoms of gastrointestinal distress were the most common adverse events, including abdominal pain, nausea, diarrhea, abdominal distention, increased stool frequency, loss of appetite, and transient fever. These adverse events were mild and resolved on their own. Paramsothy et al. [22] found worsening or discomfort in two patients undergoing FMT, but the study found no significant association between individual donors or donor lots and adverse events. Therefore, consideration may not be related to FMT. Rosen et al. [23] reported four serious adverse events in their study (including two in the donor FMT group and two in the autologous FMT group), of which one patient was found to have small bowel CD, one required cervical cancer surgery, one was admitted with abdominal pain after 11 weeks of treatment that resolved on its own, and one became severely ill from a primo Cytomegalovirus infection 7 weeks after the first infusion and was found to be in the autologous FMT group, so no serious adverse events occurred related to donor FMT. No adverse events were considered to be associated with FMT. The duration of follow-up ranged from 4 weeks to 12 months.

3.6. Microbiological Analysis. Currently, for microbiological analysis, patient stool samples are examined by extracting bacterial DNA or high-throughput 16s rRNA gene
Heterogeneity: Not applicable

Test for overall effect: $Z = 0.12 \ (P = 0.91)$

Rossen 2015 [23]  
Subtotal (95% CI)  
Total events 7  
Heterogeneity: Not applicable
Test for overall effect: $Z = 0.12 \ (P = 0.91)$

4.4.2 Lower
Costello 2018 [20]  
Fang 2021 [29]  
Moayyedi 2015 [21]  
Paramsothy 2017 [22]  
Sarbagili Shabat 2021 [30]  
Schierova 2020 [31]  
Sokol 2020 [24]  
Sood 2019 [25]  
Wei 2016 [26]  
Subtotal (95% CI)  
Total events 93  
Heterogeneity: $\tau^2 = 0.13; \ Chi^2 = 14.52, df = 8 \ (P = 0.07); I^2 = 45\%$
Test for overall effect: $Z = 2.06 \ (P = 0.04)$

Total (95% CI)  
Total events 100  
Heterogeneity: $\tau^2 = 0.11; \ Chi^2 = 15.21, df = 9 \ (P = 0.09); I^2 = 41\%$
Test for overall effect: $Z = 2.02 \ (P = 0.04)$
Test for subgroup differences: $Chi^2 = 0.89, df = 1 \ (P = 0.35); I^2 = 0\%$

Favours control    Favours FMT

Figure 7: Forest plot of the route of administration.

4.3.1 Anonymous
Costello 2018 [20]  
Crothers 2021 [28]  
Fang 2021 [29]  
Moayyedi 2015 [21]  
Paramsothy 2017 [22]  
Sarbagili Shabat 2021 [30]  
Schierova 2020 [31]  
Sokol 2020 [24]  
Sood 2019 [25]  
Wei 2016 [26]  
Subtotal (95% CI)  
Total events 95  
Heterogeneity: $\tau^2 = 0.12; \ Chi^2 = 15.40, df = 9 \ (P = 0.08); I^2 = 42\%$
Test for overall effect: $Z = 2.20 \ (P = 0.03)$

4.3.3 relatives, partners, or volunteers
Rossen 2015 [23]  
Subtotal (95% CI)  
Total events 7  
Heterogeneity: Not applicable
Test for overall effect: $Z = 0.12 \ (P = 0.91)$

Total (95% CI)  
Total events 102  
Heterogeneity: $\tau^2 = 0.10; \ Chi^2 = 16.08, df = 10 \ (P = 0.10); I^2 = 38\%$
Test for overall effect: $Z = 2.15 \ (P = 0.03)$
Test for subgroup differences: $Chi^2 = 0.98, df = 1 \ (P = 0.32); I^2 = 0\%$

Favours control    Favours FMT

Figure 8: Forest plot of donor selection.
### Study or Subgroup
<table>
<thead>
<tr>
<th>Experimental Events</th>
<th>Control Events</th>
<th>Weight (%)</th>
<th>Risk Ratio M-H, Fixed, 95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>4.6.2 with gastrointestinal pretreatment</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Costello 2018 [20]</td>
<td>18 6 35 10.8 2.76 [1.24, 6.16]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Crothers 2021 [28]</td>
<td>2 6 1.3 5.00 [0.29, 86.43]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Rossen 2015 [23]</td>
<td>7 23 8 10.2 0.95 [0.41, 2.21]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sood 2019 [25]</td>
<td>27 31 20 30 23.6 1.31 [0.98, 1.74]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Wei 2016 [26]</td>
<td>10 4 10 6.0 0.75 [0.22, 2.52]</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Subtotal (95% CI)</strong></td>
<td>108 106 51.8 1.39 [0.90, 2.14]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total events</td>
<td>57 38</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Heterogeneity:** 
- Tau² = 0.08; Chi² = 5.89, df = 4 (P = 0.21); I² = 32%
- Test for overall effect: Z = 1.49 (P = 0.14)

### 4.6.3 without gastrointestinal pretreatment
<table>
<thead>
<tr>
<th>Experimental Events</th>
<th>Control Events</th>
<th>Weight (%)</th>
<th>Risk Ratio M-H, Fixed, 95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fang 2021 [29]</td>
<td>9 10 13.6 1.80 [0.94, 3.46]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Moayyedi 2015 [21]</td>
<td>9 38 2 37 4.4 4.38 [1.01, 18.94]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Paramosothy 2017 [22]</td>
<td>18 41 8 40 12.5 2.20 [1.08, 4.46]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sarbagili Shabat 2021 [30]</td>
<td>2 17 6 15 4.5 0.29 [0.07, 1.24]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Schierova 2020 [31]</td>
<td>3 8 4 8 6.7 0.75 [0.24, 2.33]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sokol 2020 [24]</td>
<td>4 8 3 9 6.5 1.50 [0.47, 4.76]</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Subtotal (95% CI)</strong></td>
<td>108 119 48.2 1.45 [0.81, 2.58]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total events</td>
<td>45 28</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Heterogeneity:** 
- Tau² = 0.24; Chi² = 9.74, df = 5 (P = 0.08); I² = 49%
- Test for overall effect: Z = 1.26 (P = 0.21)

**Figure 9:** Forest plot of gastrointestinal pretreatment.

### Study or Subgroup
<table>
<thead>
<tr>
<th>Experimental Events</th>
<th>Control Events</th>
<th>Weight (%)</th>
<th>Risk Ratio M-H, Fixed, 95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>4.2.1 Ulcerative colitis</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Costello 2018 [20]</td>
<td>18 6 35 10.8 2.76 [1.24, 6.16]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Crothers 2021 [28]</td>
<td>2 6 1.3 5.00 [0.29, 86.43]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fang 2021 [29]</td>
<td>9 10 13.6 1.80 [0.94, 3.46]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Moayyedi 2015 [21]</td>
<td>9 38 2 37 4.4 4.38 [1.01, 18.94]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Paramosothy 2017 [22]</td>
<td>18 41 8 40 12.5 2.20 [1.08, 4.46]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sarbagili Shabat 2021 [30]</td>
<td>2 17 6 15 4.5 0.29 [0.07, 1.24]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Schierova 2020 [31]</td>
<td>3 8 4 8 6.7 0.75 [0.24, 2.33]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sokol 2020 [24]</td>
<td>4 8 3 9 6.5 1.50 [0.47, 4.76]</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Subtotal (95% CI)</strong></td>
<td>222 216 93.5 1.44 [1.00, 2.07]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total events</td>
<td>98 98</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Heterogeneity:** 
- Tau² = 0.13; Chi² = 16.08, df = 9 (P = 0.07); I² = 44%
- Test for overall effect: Z = 1.95 (P = 0.05)

### 4.2.2 Crohn’s disease
<table>
<thead>
<tr>
<th>Experimental Events</th>
<th>Control Events</th>
<th>Weight (%)</th>
<th>Risk Ratio M-H, Fixed, 95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sokol 2020 [24]</td>
<td>4 8 3 9 6.5 1.50 [0.47, 4.76]</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Subtotal (95% CI)</strong></td>
<td>8 9 6.5 1.50 [0.47, 4.76]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total events</td>
<td>4 3</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Heterogeneity:** Not applicable
- Test for overall effect: Z = 0.69 (P = 0.49)

**Figure 10:** Forest plot of IBD type. IBD: inflammatory bowel disease.
sequencing to determine changes in patient stool flora before and after transplantation. FMT significantly changes the gut microbiota of IBD patients, increasing microbial diversity or abundance. Patients treated with donor's stool were found to have a more diverse gut microbiota in favor of the donor. A systematic review was conducted due to the lack of articles on microbiota analysis. Moayyedi et al. [21] found a statistically significant change in microbial composition with more diversity in the FMT group compared to the placebo group. The microbiota of the responders were more similar to those of the donor compared to the nonresponders but did not reach statistical significance. Sokol et al. [24] found that after excluding the two patients who failed FMT at week 6 (the microbiota of the donor was poorly colonized for these two patients), a significant difference was found between the FMT and placebo groups, and the similarity of the microbiota of the FMT group to the donor persisted until the end of follow-up. Crothers et al. [28] found that FMT did not increase Shannon diversity in the subjects but caused changes in gut microbiota levels, which were highly correlated with the donor after 2 weeks of FMT treatment and persisted until 20 weeks. This may be related to the antibiotic pretreatment of subjects prior to FMT treatment in this study. Fang et al. [29] reported that receiving single-donor FMT significantly reconstituted the intestinal microbiota and preserved documentation, with an increase in the proportion of Bacteroidetes and a decrease in the proportion of Aspergillus. The relative abundance of Escherichia coli in the gut of treated subjects decreased significantly; patients treated with FMT who achieved remission also tended to have higher Prevotella abundance.

3.7. Sensitivity Analysis And Publication Bias. Sensitivity analysis of clinical remission (primary outcome) showed that the effect sizes of the outcome indicators did not change significantly after the exclusion of any of the studies, suggesting that the results of the meta-analysis were more stable and credible. Funnel plots based on publication bias and Egger’s test indicated that the conclusions were not affected by publication bias (P = 0.714) (as shown in Figure 12).
overall study quality was low. Randomization, allocation concealment, and blinding of investigators or subjects or evaluators were the main factors that may have influenced the evaluation of bias, which explains the failure of randomized controlled trial studies to provide high-quality evidence. Most of the studies only mentioned "randomization" or "randomized grouping/assignment" in the text but did not specify the specific methods used and whether allocation concealed or blinding was implemented. In addition, these studies remained heterogeneous in design, with different treatment regimen designs (e.g., route of administration and choice of stool type) and different or poorly defined efficacy endpoints.

After FMT treatment, patients with IBD had a clinical remission rate of 36.9% (168/455) and a clinical response rate of 42.0% (150/357). Positive changes in clinical, endoscopic, or fecal microbiota were present. The clinical remission rate was 30.8% (82/266) for mild-moderate IBD and 45.9% (79/172) for active IBD. Our results suggest no significant difference in the degree of disease was found.

In our study, there was no significant difference found between the fresh stool (clinical remission rate: 40.9%) and frozen stool (clinical remission rate: 32.2%). Moreover, some studies have found that fresh FMT has superior clinical efficacy to frozen FMT [34]. Hamilton et al. [35] found that frozen stool microbiota from a healthy donor was more effective in treating recurrent CDI, thereby restoring the structure of the gut microbiota and clearing *C. difficile*. A randomized controlled trial [36] investigated the effectiveness of frozen FMT and fresh FMT in the treatment of recurrent CDI diarrhea, with clinical remission rates of 75.0% for frozen FMT and 70.3% for fresh FMT in a modified intention-to-treat analysis, showing similar efficacy and no major differences between the two stool types. The results of this study provide evidence for the use of fresh stools.

FMT can be administered via the upper GI tract (nasogastric tube, gastrostoduodenal tube, oral) or the lower GI tract (colonoscopy, cecum, enema), two different routes. A systematic review has shown that the lower GI tract is a safe and effective route for FMT in the treatment of CDI [37, 38]. In our study, the clinical remission rate was 38.2% (151/395) in the lower GI tract and 31.2% (15/48) in the upper GI tract. Our study suggests that no significant difference in the route of administration was found. Ramai et al. [39] concluded that colonoscopy has a better cure rate than the nasogastric tube for CDI. In addition to efficacy, patient compliance, cost-effectiveness, comfort of administration, level of invasiveness, risk of aspiration and infection, multiplicity of drugs to be administered, and relapse rate appear to be the main factors influencing physicians’ choice of route of administration [40]. Some studies suggest that the use of upper GI administration requires attention to adverse events such as patient vomiting [41] and aspiration pneumonia [42], while there are also risks associated with lower GI administration in patients with severe colitis. Therefore, more extensive studies are needed to find low-risk, convenient, and cost-effective routes of administration.

At present, there is no consensus on the optimal donor for FMT. Our study found that fecal donors mainly included healthy adults, relatives or friends, or anonymous volunteers. Clinical remission was 37.6% (153/407) for feces from healthy anonymous donors and 31.2% (15/48) for feces from relatives, close friends, or volunteers. In our study, there was no significant difference found between the efficacy of FMT derived from healthy anonymous donors and the FMT using relatives, close friends, or volunteers. Some studies agree with our results that the ideal donor is a healthy donor [43]. Others have concluded that there is no significant difference between healthy donors and relative or close friend FMT [44]. The advantage of healthy anonymous donors is that they are easy to extract and screen from stool banks, saving time and money for screening stool donors [45]. Considering the possibility that the sample size using relatives or close friends was insufficient, more randomized controlled trials are needed to support our conclusions.

Clinical remission rates were 44.4% (95/214) and 30.3% (73/241) in the group with and without gastrointestinal pretreatment, respectively. In our study, there was no significant difference found between pretreatment of the gastrointestinal tract and treatment with FMT. One study found that bowel cleansing prior to colonoscopy and/or stool disposal led to changes in symptoms and gut microbiota composition in irritable bowel syndrome (IBS) [46]. Bowel cleansing brings immediate changes to the gut microbiota [47, 48], which plays an integral role in IBD. However, the amount of relevant trials is insufficient, so a more extensive study in this area is warranted. The clinical response rate was 36.8% (161/438) in UC patients and 41.2% (7/17) in CD patients. In our study, there was no significant difference found between CD patients and UC patients and the number of RCTs in CD is relatively insufficient, so more extensive research in this area is needed to support this conclusion. In
our study, no significant differences were found in the gender of the patients.

Currently, the most common adverse events are self-limiting gastrointestinal discomfort, including abdominal pain, bloating, diarrhea, nausea, and other adverse events such as transient fever. These adverse events usually resolve on their own within 24 hours. The follow-up period is 6 weeks to 36 months.

The current systematic review still has some limitations. First, different infusion frequencies (single or multiple FMTs), number of donors, and preparation and storage of donor stools are factors that can affect the results, increasing heterogeneity between studies. Secondly, subgroup analysis (e.g., route of administration, type of stool, gastrointestinal pretreatment, and IBD type) has the limitation of a small number of studies and insufficient sample size. Finally, there is uncertainty regarding the selection of FMT donors, although some studies have shown no significant differences between healthy donors and relatives or close friends. Future larger and more extensive trials are needed to optimize the use of FMT, including donor selection, stool preparation and storage, and route of administration, to obtain better outcomes for patients.

5. Conclusions

In conclusion, our study found that FMT is an effective, feasible, and safe treatment for IBD. It has a positive effect in terms of clinical remission/clinical response. It can also be done by increasing the diversity and species richness of the gut microbiota. Still, further randomized controlled trials are needed to fully evaluate the long-term efficacy and safety of FMT in patients with IBD.

Data Availability

The data supporting this meta-analysis are from previously reported studies and datasets, which have been cited. The processed data are available from the corresponding author upon request.

Conflicts of Interest

The authors declare that there are no conflicts of interest regarding the publication of this article.

Authors’ Contributions

BJ initially conceived the study. XYT designed the study and produced the first draft of the study. YJX and XYL performed the preliminary search. XYT and XLL will read the full texts of the studies and extract the data. XYT and YJX will perform the data synthesis. BJ will arbitrate in cases of any disagreement. BJ and XLL revised the manuscript. All authors read and approved the final manuscript.

Acknowledgments

This work was supported by the National Natural Science Foundation of China (No. 325020015).

Supplementary Materials

See PRISMA_2020_checklist in the Supplementary Materials for detailed Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA). (Supplementary Materials)

References


