

# British Society of Gastroenterology guidelines on the management of irritable bowel syndrome

Dipesh H Vasant<sup>1,2</sup> Peter A Paine,<sup>3</sup> Christopher J Black<sup>4</sup>,  
 Lesley A Houghton<sup>5,6</sup> Hazel A Everitt,<sup>7</sup> Maura Corsetti,<sup>8,9</sup>  
 Imran Aziz<sup>10</sup> Adam D Farmer,<sup>11,12</sup> Maria P Eugenicos,<sup>13</sup> Rona Moss-Morris,<sup>14</sup>  
 Yan Yiannakou,<sup>15</sup> Alexander C Ford<sup>16</sup>

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For numbered affiliations see end of article.

## Correspondence to

Professor Alexander C Ford,  
 Leeds Gastroenterology  
 Institute, St James's University  
 Hospital, Leeds, UK;  
[alex12399@yahoo.com](mailto:alex12399@yahoo.com)

DHV and PAP are joint first authors.

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## ABSTRACT

Irritable bowel syndrome (IBS) remains one of the most common gastrointestinal disorders seen by clinicians in both primary and secondary care. Since publication of the last British Society of Gastroenterology (BSG) guideline in 2007, substantial advances have been made in understanding its complex pathophysiology, resulting in its re-classification as a disorder of gut-brain interaction, rather than a functional gastrointestinal disorder. Moreover, there has been a considerable amount of new evidence published concerning the diagnosis, investigation and management of IBS. The primary aim of this guideline, commissioned by the BSG, is to review and summarise the current evidence to inform and guide clinical practice, by providing a practical framework for evidence-based management of patients. One

of the strengths of this guideline is that the recommendations for treatment are based on evidence derived from a comprehensive search of the medical literature, which was used to inform an update of a series of trial-based and network meta-analyses assessing the efficacy of dietary, pharmacological and psychological therapies in treating IBS. Specific recommendations have been made according to the Grading of Recommendations Assessment, Development and Evaluation system, summarising both the strength of the recommendations and the overall quality of evidence. Finally, this guideline identifies novel treatments that are in development, as well as highlighting areas of unmet need for future research.

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## Other treatments in development

In recent years, there has been considerable interest in the evaluation of faecal microbiota transplantation (FMT) for IBS. Unfortunately, a meta-analysis of 5 RCTs, containing 267 patients, demonstrated no significant benefit of FMT compared with placebo (RR 0.98; 95% CI 0.58 to 1.66), and in 2 pooled trials placebo capsules administered orally were superior to capsules containing donor stool (RR 1.96; 95% CI 1.19 to 3.20).<sup>294</sup> Criticisms of the trials, to date, have included small sample sizes, heterogeneity in IBS subtypes recruited, lack of standardisation of donor samples and suboptimal end points used. There is therefore a need for further, large, high-quality trials of FMT for IBS, perhaps targeting subgroups of patients with evidence of dysbiosis, who may be more likely to benefit. At present, therefore, there is insufficient evidence to recommend FMT for IBS outside of a research setting. **Enterogel**, an intestinal adsorbent approved for use in IBS-D and available over-the-counter in the UK is currently the subject of a multicentre RCT in IBS-D (Kemppinen A, Howell C, Allgar V, *et al*. Randomised, double-blind, placebo controlled multi-centre study to assess the efficacy, tolerability and safety of **Enterogel**® in the treatment of irritable bowel syndrome with diarrhoea (IBS-D) in adults. *Trials* 2020;21:122).

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