

Mistakes in irritable bowel syndrome and how to avoid them

Robin Spiller

Around 11% of the worldwide population experience irritable bowel syndrome (IBS), making it one of the most frequent gastroenterological diagnoses.¹ The symptoms of IBS include abdominal pain associated with unpredictable bowel habits and variable changes in the form and frequency of stool.² While all patients with IBS suffer from recurrent bouts of abdominal pain, their bowel habits are varied: around one-third suffer predominantly with diarrhoea (IBS-D), one-fifth experience predominantly constipation (IBS-C) and half have an erratic mixed pattern of both diarrhoea and constipation (IBS-M).³ This very heterogeneous condition undoubtedly has multiple causes and an individualized approach to management and treatment is required.

Here I discuss the mistakes most frequently made when diagnosing and managing IBS. The mistakes and discussion that follow are based, where possible, on published data and failing that, many years of my own clinical experience.



Mistake 1 Failing to detect bile salt malabsorption

If excessive amounts of bile acids enter the colon, colonic secretion is stimulated and the amount of water incorporated in the stool increased, which causes frequent loose stools associated with a sensation of urgency, often accompanied by nocturnal diarrhoea. According to the findings of a meta-analysis, 10% of patients with IBS-D-like symptoms may have severe bile acid malabsorption, retaining <5% of bile acids at 7 days.⁴ A UK survey indicates that almost one in four IBS patients who are referred to secondary care with diarrhoea have bile acid diarrhoea.⁵

The most sensitive and specific test for bile acid malabsorption remains the 7-day retention of Selenium-75-labelled homocholic acid taurine (⁷⁵SeHCAT). If retention at 7 days is <5%, the test predicts a 100% response to colestyramine, while 5–10% retention predicts a response of around 37%.⁶ Since the ⁷⁵SeHCAT test is not available worldwide alternative blood tests have been suggested, as has the simpler therapeutic trial of colestyramine; however, such trials are less reliable as they are influenced by many other uncontrolled factors like diet and emotion. Alternative assessments include measuring serum levels of

7-alpha-hydroxy-4-cholesten-3-one (C4), which is a key intermediary in bile acid synthesis from cholesterol, faecal bile acids and serum FGF19, which is a signalling molecule that normally provides negative feedback to inhibit bile acid synthesis; however, these tests are only available in a few laboratories, though this may change in the future.^{7,8}

Mistake 2 Failing to recognize somatization, leading to multiple referrals to non-gastrointestinal specialists

Multiple medically unexplained symptoms are a common feature in patients who have IBS. This feature can easily be assessed using the Patient Health Questionnaire-12 somatic symptom (PHQ-12SS) scale, which asks about non-gastrointestinal symptoms such as bodily pains and symptoms. Less than 5% of healthy controls score more than 6 on the PHQ-12SS scale, while 67% of IBS patients do.⁹ High scores predict more visits to the primary care physician and are clinically useful. Low scores suggest that an alternative diagnosis needs to be excluded. Ignoring this feature results in multiple referrals to non-gastrointestinal specialists and is a very

likely cause of the excess of hysterectomies and cholecystectomies seen in IBS patients.^{10,11}

Mistake 3 Not telling the patient that they have a high probability of having IBS at the onset of investigation

Meeting IBS criteria in the absence of any alarm features is associated with a very high probability that investigations will yield normal results,¹² so it is important to make this clear to the patient at the onset. In this setting, when test results turn out to be normal the soundness of the diagnosis will be apparent to the patient. By contrast, if no prior diagnosis has been made then a negative test may simply lead to the demand for more tests, an all too common feature of many IBS patients' medical 'careers'.

Mistake 4 Failing to recognize the key features of bloating, leading to multiple negative investigations including CT and ultrasound

Bloating is a condition that is mysterious to many patients and physicians, and often leads to unnecessary investigations and considerable irradiation. Two types of bloating need to be recognized. The first involves a sensation of distension without any obvious change in girth and is thought to reflect increased visceral sensitivity.¹³ The second is characterized by visible distension that requires loosening of clothes and an increase in abdominal girth, something that usually worsens during the day and remits overnight.¹⁴ Until recently it was unclear how even a mouthful of food could induce a sudden distension of the abdomen. We now recognize, however, that this very characteristic and diagnostically helpful feature is due to a combination of

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Robin Spiller is at the Nottingham Digestive Diseases Centre, University of Nottingham, Queens Medical Centre, Nottingham, United Kingdom.

Correspondence to: robin.spiller@nottingham.ac.uk
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relaxation of the abdominal wall and lowering of the diaphragm.¹⁵ This neural response can occur within seconds. Bloating thus does not involve any acute change in abdominal contents. As an increasing amount of abdominal fat is a frequent cause of a slow, progressive increase in abdominal distension, recent weight gain should be specifically enquired for in such patients. Ovarian cancer can also present with progressive distension but in this case the day-to-day variability characteristic of IBS is lacking.

Mistake 5 Using opiates to control IBS pain

Although the pain in patients with IBS is often described as extremely severe and opiates are undoubtedly effective, most clinicians strongly advise against their use because receptor desensitization occurs rapidly leading to rapid dose escalation. High doses of opiates are associated with troublesome side effects, including nausea and vomiting, as well as profound constipation and drug dependence.¹⁶ While IBS symptoms are usually intermittent, opiate use is constant. In a subgroup of susceptible patients who often have psychological comorbidities, opiate use may result in 'narcotic bowel syndrome', in which the opiates appear to actually aggravate the pain. Opiate withdrawal is difficult owing to psychological dependence, but can result in marked remission of pain.¹⁷

Mistake 6 Misdiagnosing Crohn's disease as IBS-D

All new cases meeting the Rome III criteria for IBS-D should have, as a minimum, a full blood count, serological test for coeliac disease and a faecal calprotectin measurement to exclude inflammatory bowel disease (IBD). An ileocolonoscopy should be performed for those with abnormal results or for other reasons, such as a family history of IBD or weight loss. If symptoms are chronic and unchanged since a previous normal colonoscopy this need not be repeated unless there is evidence of systemic inflammation (raised CRP levels or platelet count) or elevated faecal calprotectin.

Referred patients probably have a greater risk of having Crohn's disease. Indeed, a large study in Canada suggested that 8.6% of patients referred to secondary care who met the Rome III criteria turned out to have Crohn's disease.¹⁸ Community studies indicate that patients who have colonic Crohn's disease can have symptoms for many years prior to diagnosis¹⁹ and are often labelled as having

IBS since they lack the key alarm features of rectal bleeding and weight loss. Faecal calprotectin has high sensitivity and specificity for IBD,²⁰ as may a full blood count showing an elevated platelet count or microcytosis.²¹

Mistake 7 Performing cholecystectomy for right upper quadrant pain without gallstones

The pain in patients with IBS is poorly localized, but may in some cases be right upper quadrant pain, which can lead to confusion with biliary pain. Relief on defaecation may help distinguish the two. The pattern of pain is also helpful: biliary pain is typically very episodic with weeks of freedom, whereas IBS pain is associated with only a few days free from pain before the next flare occurs. Postcholecystectomy pain may reflect the presence of pre-existing, unrecognized IBS.

Mistake 8 Performing a hysterectomy/laparoscopy and division of adhesions for IBS pain

As previously mentioned, IBS patients have an increased risk of undergoing gynaecological procedures, which is most likely due to the attribution of IBS symptoms to gynaecological disease. Paying careful attention to the Rome criteria, especially relief on defaecation or association of pain with changes in bowel habit, should help distinguish IBS from other causes of lower abdominal pain. Likewise, multiple somatic complaints should also point towards a diagnosis of IBS²² rather than a specific gynaecological cause. Once surgery has been performed there is a very real risk of developing adhesions, further confusing the diagnosis and hindering management.

Mistake 9 Testing for lactose intolerance when a patient consumes <240ml of milk or its equivalent per day

Taking a careful dietary history is important before any dietary recommendations are made. Many patients already restrict their consumption of dairy products and there is little point in doing a lactose tolerance test on someone who consumes <240ml of milk or its equivalent per day. A randomized blinded trial showed that this amount of milk could not be distinguished from a lactose-free placebo, even in those with true lactose malabsorption.²³ More recent studies demonstrate that symptoms developing after lactose challenge are dose dependent: only 3% of those with genetically determined lactose malabsorption developed symptoms with

a 10 g lactose challenge, rising to 21.7% of patients challenged with 20 g lactose and 73.3% of patients challenged with 40 g lactose.²⁴ IBS patients, however, show more symptoms after each dose regardless of its size, indicating a degree of visceral hypersensitivity. It is also worth noting the strong nocebo²⁵ effect of challenging IBS patients with foods they believe they are intolerant of. Thus, until underlying beliefs have been changed, little progress can be expected.

Mistake 10 Encouraging food exclusion without reinforcing the need to reintroduce foods to confirm apparent intolerance, leading to ever more restricted diets and malnutrition

Some patients develop an eating disorder and lose weight because they exclude more and more foods as they try to link flares of IBS symptoms with particular foods. It is vital to explain to patients that flares should only be attributed to foods if the response can be reproduced on more than one occasion. It is also important to test these foods again after an interval—in many cases double blind challenge later shows these foods do not cause symptoms and that the flare was due to other uncontrolled factors. As previously mentioned, a strong nocebo effect²⁵ can lead to these beliefs being self-perpetuating, so supervision of such exclusion diets by a dietician is helpful to avoid patients developing a nutritionally inadequate diet.

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References

1. Lovell RM and Ford AC. Effect of gender on prevalence of irritable bowel syndrome in the community: systematic review and meta-analysis. *Am J Gastroenterol* 2012; 107: 991–1000.
2. Longstreth GF, Thompson WG, Chey WD, et al. Functional bowel disorders. *Gastroenterology* 2006; 130: 1480–1491.
3. Tillisch K, Labus JS, Naliboff BD, et al. Characterization of the alternating bowel habit subtype in patients with irritable bowel syndrome. *Am J Gastroenterol* 2005; 100: 896–904.
4. Wedlake L, A'Hern R, Russell D, et al. Systematic review: the prevalence of idiopathic bile acid malabsorption as diagnosed by SeHCAT scanning in patients with diarrhoea-predominant irritable bowel syndrome. *Aliment Pharmacol Ther* 2009; 30: 707–717.
5. Aziz I, Mumtaz S, Bholah H, et al. High prevalence of idiopathic bile acid diarrhea among patients with diarrhea-predominant irritable bowel syndrome based on Rome III criteria. *Clin Gastroenterol Hepatol* 2015; 13: 1650–1655.e2.
6. Williams AJK, Merrick MV and Eastwood MA. Idiopathic bile acid malabsorption—A review of

- clinical presentation, diagnosis, and response to treatment. *Gut* 1991; 32: 1004–1006.
7. Vijayvargiya P, Camilleri M, Shin A, et al. Methods for diagnosis of bile acid malabsorption in clinical practice. *Clin Gastroenterol Hepatol* 2013; 11: 1232–1239.
 8. Walters JR, Tasleem AM, Omer OS, et al. A new mechanism for bile acid diarrhea: defective feedback inhibition of bile acid biosynthesis. *Clin Gastroenterol Hepatol* 2009; 7: 1189–1194.
 9. Spiller RC, Humes DJ, Campbell E, et al. The Patient Health Questionnaire 12 Somatic Symptom scale as a predictor of symptom severity and consulting behaviour in patients with irritable bowel syndrome and symptomatic diverticular disease. *Aliment Pharmacol Ther* 2010; 32: 811–820.
 10. Hasler WL and Schoenfeld P. Systematic review: Abdominal and pelvic surgery in patients with irritable bowel syndrome. *Aliment Pharmacol Ther* 2003; 17: 997–1005.
 11. Longstreth GF and Yao JF. Irritable bowel syndrome and surgery: A multivariable analysis. *Gastroenterology* 2004; 126: 1665–1673.
 12. Vanner SJ, Depew WT, Paterson WG, et al. Predictive value of the Rome criteria for diagnosing the irritable bowel syndrome. *Am J Gastroenterol* 1999; 94: 2912–2917.
 13. Houghton LA and Whorwell PJ. Towards a better understanding of abdominal bloating and distension in functional gastrointestinal disorders. *Neurogastroenterol Motil* 2005; 17: 500–511.
 14. Houghton LA, Lea R, Agrawal A, et al. Relationship of abdominal bloating to distension in irritable bowel syndrome and effect of bowel habit. *Gastroenterology* 2006; 131: 1003–1010.
 15. Villoria A, Azpiroz F, Burri E, et al. Abdomino-phrenic dyssynergia in patients with abdominal bloating and distension. *Am J Gastroenterol* 2011; 106: 815–819.
 16. Tuteja AK, Biskupiak J, Stoddard GJ, et al. Opioid-induced bowel disorders and narcotic bowel syndrome in patients with chronic non-cancer pain. *Neurogastroenterol Motil* 2010; 22: 424–430.
 17. Kurlander JE and Drossman DA. Diagnosis and treatment of narcotic bowel syndrome. *Nat Rev Gastroenterol Hepatol* 2014; 11: 410–418.
 18. Ford AC, Bercik P, Morgan DG, et al. Validation of the Rome III criteria for the diagnosis of irritable bowel syndrome in secondary care. *Gastroenterology* 2013; 145: 1262–1270.e1.
 19. Burgmann T, Clara I, Graff L, et al. The Manitoba Inflammatory Bowel Disease Cohort Study: prolonged symptoms before diagnosis—how much is irritable bowel syndrome? *Clin Gastroenterol Hepatol* 2006; 4: 614–620.
 20. Tibble JA, Sigthorsson G, Foster R, et al. Use of surrogate markers of inflammation and Rome criteria to distinguish organic from nonorganic intestinal disease. *Gastroenterology* 2002; 123: 450–460.
 21. Cabrera-Abreu JC, Davies P, Matek Z, et al. Performance of blood tests in diagnosis of inflammatory bowel disease in a specialist clinic. *Arch Dis Child* 2004; 89: 69–71.
 22. Spiller RC, Humes DJ, Campbell E, et al. The Patient Health Questionnaire 12 Somatic Symptom scale as a predictor of symptom severity and consulting behaviour in patients with irritable bowel syndrome and symptomatic diverticular disease. *Aliment Pharmacol Ther* 2010; 32: 811–820.
 23. Suarez FL, Savaiano DA and Levitt MD. A comparison of symptoms after the consumption of milk or lactose-hydrolyzed milk by people with self-reported severe lactose intolerance. *N Engl J Med* 1995; 333: 1–4.
 24. Yang J, Deng Y, Chu H, et al. Prevalence and presentation of lactose intolerance and effects on dairy product intake in healthy subjects and patients with irritable bowel syndrome. *Clin Gastroenterol Hepatol* 2013; 11: 262–268.
 25. Biesiekierski JR, Peters SL, Newnham ED, et al. No effects of gluten in patients with self-reported non-celiac gluten sensitivity after dietary reduction of fermentable, poorly absorbed, short-chain carbohydrates. *Gastroenterology* 2013; 145: 320–328.

Your IBS briefing

Online courses

- 'Irritable Bowel Syndrome' from UEG [<https://www.ueg.eu/education/online-courses/irritable-bowel-syndrome/>].

UEG Week sessions

- 'Practical management of patients with irritable bowel syndrome (IBS)' at UEG Week 2015 [<https://www.ueg.eu/education/session-files/?session=1353&conference=109>].
- 'Irritable bowel syndrome: What can science tell us' at UEG Week 2015 [<https://www.ueg.eu/education/session-files/?session=1393&conference=109>].
- 'From guidelines to clinical practice: IBS management' at UEG Week 2015 [<https://www.ueg.eu/education/session-files/?session=1410&conference=109>].
- 'Altered intestinal microbiota composition in IBS: Does it affect clinical practice?' At UEG Week 2014 [<https://www.ueg.eu/education/session-files/?session=1143&conference=76>].
- 'Therapy update: How to be successful when you treat IBS' at UEG Week 2014 [<https://www.ueg.eu/education/session-files/?session=1224&conference=76>].

education/session-files/?session=1224&conference=76].

Standards and Guidelines

- NICE guidelines [CG61]. Irritable bowel syndrome in adults: diagnosis and management. Updated February 2015 [<https://www.nice.org.uk/guidance/cg61>].
- Weinberg DS, Smalley W, Heidelbaugh JJ, et al. American Gastroenterological Association Institute Guideline on the Pharmacological Management of Irritable Bowel Syndrome. *Gastroenterology* 2014; 147: 1146–1148 [<http://www.gastro.org/guidelines/2014/09/14/pharmacological-management-of-ibs>].
- World Gastroenterology Organisation Global Guidelines. Irritable Bowel Syndrome: a Global Perspective. Updated September 2015 [<http://www.worldgastroenterology.org/guidelines/global-guidelines/irritable-bowel-syndrome-ibs/irritable-bowel-syndrome-ibs-english>].
- Lacy BE, Mearin F, Chang L, et al. Bowel Disorders. *Gastroenterology* 2016; 150: 1393–1407.e5 [[http://www.gastrojournal.org/article/S0016-5085\(16\)00222-5/abstract](http://www.gastrojournal.org/article/S0016-5085(16)00222-5/abstract)]