ASIAN JOURNAL OF PHARMACEUTICAL AND BIOLOGICAL RESEARCH





Asian journal of Pharmaceutical and biological research 2231-2218 http://www.ajpbr.org/ Universal IMPACT factor 7 SJIF 2022: 4.465 Volume 12 Issue 2 **MAY-AUG. 2023 Editorial board** Dr. Madhu Bala Scientist 'F' and Joint Director, Institute of Nuclear Medicine and Allied Sciences (INMAS), India Dr. Sandip Narayan Chakraborty Research Asst, Translational Molecular Pathology, Ut Md Anderson Cancer Center, Life Sciences Plaza, Houston, TX 77030 Dr. Tushar Treembak Shelke Head of Department of Pharmacology and Research Scholar, In Jspms Charak College of Pharmacy & Research, Pune, India Dr. Subas Chandra Dinda Professor-cum-Director: School of Pharmaceutical Education & Research (SPER), Berhampur University, Berhampur, Orissa, India. Dr. Jagdale Swati Changdeo Professor and Head, Department of Pharmaceutics, MAEER's Maharashtra Institute of Pharmacy, S.No.124, MIT Campus, Kothrud, Pune-411038 Dr. Biplab Kumar Dev Principal, Department of Pharmacy, Assam downtown University, Sankar Madhab Path, Panikhaiti 781026, Guwahati, Assam, India Dr. Yogesh Pandurang Talekar Research Associate, National Toxicology Centre Dr. Indranil Chanda Assistant Professor, Girijananda Chowdhury Institute of Pharmaceutical Science, Hathkhowapara, Azara Guwahati-17, Assam, India. Dr. Sudip Kumar Mandal Department of Pharmaceutical Chemistry, Dr. B. C. Roy College of Pharmacy & AHS, Bidhannagar, Durgapur-713206, India. Sodikova Dilrabokhon Andijan state medical institute Dr., associate professor Kuryazova Sharofat Tashkent Pediatric medical institute Dr., Abdurakhmanova Nigora Nazimovna Tashkent Pediatric Medical Institute Abdullaeva Umida Bukhara state medical institute Dr. Neeraj Upmanvu Prof., Peoples Institute of Pharmacy & Research Center, Bhopal, MP, India. Dr. Mirrakhimova Maktuba Khabibullaevna Tashkent medical academy Uzbekistan Dr. Nishanova Aziza Abdurashidovna, Tashkent State Dental Institute Dr. Sadikova Minurakhon Adkhamovna Andijan State Medical Institute Kurbanova Sanobar Yuldashevna Tashkent State Dental Institute Zokirova Nargiza Bahodirovna Tashkent Pediatric medical institute Khabilov Behzod Nigmon ugli Tashkent State Dental Institute Dr. Domenico De Berardis Department of Mental Health, Azienda Sanitaria Locale Teramo, 64100 Teramo. Italy Dr. Azizova Rano Baxodirovna associate professor of the Department of neurology of the Tashkent Medical Academy Dr. Ishankhodjaeva Gulchekhra Tashkent Medical Academy

Institute of Nuclear Medicine and Allied Sciences (INMAS), India

Brig SK Mazumdar Marg, Timarpur, New Delhi, Delhi 110054 India

IRRITABLE BOWEL SYNDROME: STAGES OF DIAGNOSIS Makhmudova Lola Izzatilloyevna Bukhara state medical institute, Bukhara, Uzbekistan

A. Manning created the first set of formal criteria that made it possible to diagnose IBS with a certain degree of certainty without the need for an extensive expensive examination. And this set of symptoms formed the basis for the development of the Rome diagnostic criteria for IBS in the III edition that existed until recently [1].

In 1984, Kruis and colleagues reported a similar set of symptoms used to define IBS: abdominal pain; bloating; and altered bowel function. In contrast to the Manning criteria, the Cruise criteria paid more attention to the duration of symptoms and actually suggested a two-year duration. More importantly, the Cruis criteria emphasized the need to consider warning signs ("red flags") as well as rule out organic disease in combination with routine physical examination and basic laboratory tests (CBC and ESR). Ultimately, however, these criteria turned out to be too cumbersome to be used in clinical practice and lost their popularity [2, 24-28].

In 1988, a group of international experts met in Rome to discuss functional gastrointestinal disorders (FGI). The main goal was to classify FGID using a symptom-based classification scheme, emphasizing the fact that patients report symptoms despite the absence of chemical, radiological, or physiological abnormalities. This culminated in the publication of the Rome Criteria in 1992 (later known as Rome I), which raised the medical community's awareness of FGID. Bloating, the main symptom of many patients with IBS, was no different from abdominal pain. The IBS criteria have been easily incorporated into research studies but have proven cumbersome for clinical practice.

A few years later, the Rome Committee met again to revise the original Rome I criteria based on feedback from clinicians, investigators, regulators, and new information gleaned from the scientific literature. The revised Rome II criteria were

published in 1999 [4,2]. Like Rome I, Rome II required symptoms to have been present for at least 12 weeks out of the preceding 12 months, although the times need not be consistent. The term "discomfort" was added to the definition, and a new criterion was added, noting that two of the three criteria for abdominal pain should have been required for the diagnosis of IBS to guarantee the presence of altered bowel habits. At that time, patients were not divided into specific subtypes based on bowel habits [5].

The Rome III criteria were introduced in 2006, with the most significant change being the classification of IBS into subtypes. Subtypes were based on stool consistency rather than stool frequency and included IBS-C (constipation), IBS-D (diarrhea), IBS-M (mixed) and IBS-U intermittent (the present stool consistency disorders are not sufficient to apply the criteria of the first three IBS options) option [6]. Another significant change was that the symptom of bloating as the main symptom was removed from the definition. This change was based on the notion that bloating as a symptom is so common that it is neither sensitive nor specific to IBS. In a validation study conducted by Ford and colleagues in patients with IBS symptoms who underwent colonoscopy, the sensitivity of the Rome III criteria was 68.8%,

Since the release of the Rome III criteria in 2006, research on IBS has increased dramatically. Creative research work in the basic and clinical sciences has led to the identification of new etiologies of IBS and to a better understanding of the complex pathophysiology underlying the onset of IBS symptoms. Many new drugs have come on the market that have focused on specific subtypes of IBS, based in part on a better understanding of the underlying pathophysiology. These advances in knowledge, along with a desire to make the Rome criteria more clinically useful, led to several key changes to the Rome criteria when the fourth iteration was released in 2016 [2,7].

Rome IV determinedirritable bowel syndrome (IBS) as a functional bowel disorder in which recurrent abdominal pain is associated with defecation or a change

in bowel habits. Bowel disorders (eg, constipation, diarrhea, or a combination of constipation and diarrhea) are usually present, as are symptoms of bloating/bloating. Symptoms must have appeared at least 6 months before diagnosis, and symptoms must have been present for the past 3 months.

The criteria of Rome IV differ from the criteria of Rome III) in several distinct ways. First, the term "discomfort" has been removed from the current definition and diagnostic criteria because some languages do not have the word "discomfort" or have different meanings in different languages. In addition, based on a study of IBS patients who reported significant differences in understanding of these terms, it is not clear whether the difference between pain and discomfort is qualitative or quantitative. Secondly, the frequency of abdominal pain was increased from 3 days per month to one day per week on average. Although this change appears small, it was based on a large population-based study to increase the sensitivity and specificity of the criteria. Third, bloating and distension are now recognized as common symptoms. This highlights the prevalence of these symptoms in patients with IBS and other FGIDs (Chronic constipation, functional dyspepsia) and reinforces the earlier results of Kruis et al. Fourth, the previous criteria included a somewhat ambiguous phrase about the presence of disordered defecation. This has now been clarified with the phrase "... intestinal disorders (constipation, diarrhea or a mixture of constipation and diarrhea) are usually present." Finally, it is now expressly stated that IBS subtypes are based on predominant bowel habits on days with abnormal bowel movements. The Rome Committee, using data from a large population-based study (Rome Regulatory Survey of Gastrointestinal Symptoms; unpublished), determined that analysis of days without bowel movement did not increase bowel subtype specificity, and analysis of only days with abnormal bowel movements increased specificity [8,2, 29-32].

Asian journal of Pharmaceutical and biological research 2231-2218

http://www.ajpbr.org/ Universal IMPACT factor 7

SJIF 2022: 4.465

Volume 12 Issue 2

MAY-AUG. 2023

| Table 2. Stages of diagnostic criteria for IBS | | | | |
|--|---|-------------------------------|-------------------------------|--|
| A. Manning | Rome I | Rome II | Rome III | Rome IV (2016) |
| (1978) | (1989) | (1999) | (2006) | |
| Two or more of the | At least 3 months of | At least 12 weeks in the last | At least 3 days per month for | Recurrent abdominal pain, |
| following | continuous or recurrent pain in | 12 months of continuous or | the last 12 weeks of | on average at least |
| symptoms: | stomach: | recurrent abdominal | continuous or recurrent | 1 day/week in the past 3 |
| bloating | • relief from bowel movements | pain or discomfort | abdominal pain or discomfort | months associated with 2 or |
| relief of pain during | or | At least 2 of the following | At least 2 of the following | more of the following: |
| bowel movements | association with changes in | symptoms: | symptoms: | associated with defecation |
| frequent stools with | stool consistency | • relief from bowel | • relief from bowel | • associated with changes in |
| pain | At least 2 of the following | movements | movements | stool frequency |
| loose stools at the | symptoms on at least 25% of | • change in stool frequency | • change in stool frequency | associated with a change |
| beginning of pain | days: | • changing the shape of the | • changing the shape of the | in the shape of the stool |
| passage of mucus | • change in stool frequency | chair | chair | Criteria are valid if they |
| feeling of incomplete | • changing the shape of the | Onset of symptoms more than | Onset of symptoms more than | have been available within |
| emptying | chair | 12 months before diagnosis | 6 months before diagnosis | the last |
| | • changing the passage of the | | | 3 months with symptom |
| | chair | | | onset at least 6 months ago! |
| | • passage of mucus | | | |
| | bloating or distension | | | |

6 11 ... • • • TDC

Stages of diagnostic criteria for IBS are presented in Table. 2.

IBS is subdivided into 3 main subtypes according to the predominant type of colonic disorder: IBS with constipation (IBS-C), IBS with diarrhea (IBS-D), and mixed subtype of IBS (IBS-M) (Table 3).

Patients who meet the diagnostic criteria for IBS, but in whom it is difficult to accurately identify one of the three types, should be categorized as unclassified IBS [9,10]. Difficulties in choosing 1 of the 3 main subgroups to which the patient belongs may arise as a result of frequent changes in diet and the use of various drugs that affect transit through the gastrointestinal tract. The choice of subtype should be based on the predominant type of intestinal contractility disorder. The Bristol Stool Shape Scale should be used to assess stool consistency.

| Subtype | Characteristic | | |
|------------------|---|--|--|
| SRK-C | More than a quarter (25%) of all acts of defecation - 1st or Type 2 (hard or fragmented stools) on the Bristol Stool Shap Scale and less than a quarter (25%) - 6th or 7th type (liquid or watery) | | |
| SRK-D | More than a quarter (25%) of all acts of defecation - the 6th or 7th type (liquid or watery) according to Bristol stool shape scale and less than one quarter (25%) - Type 1 or 2 (hard or fragmented stools) | | |
| SRK-M | More than a quarter (25%) of all acts of defecation of the 1st or type 2 (hard or fragmented stools) Bristol stool shape scale and more than a quarter (25%) - 6th or 7th type (liquid or watery) | | |
| Unclassified IBS | Patients who fit the diagnostic criteria for IBS, but a precise definition for which | | |

one of the three types is difficult

The Bristol Chair Shape Scale (BCSS) was developed in the 1990s at the Bristol Royal Infirmary in England [11]. The authors described seven types of stool, which are noted below:

- Type 1: Separate hard lumps like nuts (hard to convey)
- Type 2: Sausage but lumpy
- Type 3: Like a sausage but cracked on the surface
- Type 4: like sausage or snake, smooth and soft
- Type 5: Soft drops with clear edges (easy to transfer)
- Type 6: Fluffy pieces with carved edges, soft stool
- Type 7: watery, no solids, completely liquid

The authors classified type 1 and 2 stools as associated with constipation, while type 6 and 7 stools were associated with diarrhea (and type 5 stools to some extent). Stool types 3 and 4 were considered normal stool. BSFS is a convenient way of describing the bowel habits of patients and is commonly used in clinical trials. In addition, at two extremes (Bristol stool types 1 and 2 or types 6 and 7), stool shape serves as a crude surrogate marker for colonic transit. Patients with IBS-C have >25% of their bowel movements associated with BCSS 1 or 2, while patients with IBS-D have >25% of their bowels associated with BSFS 6 or 7. Those with a mixed subtype of intermittent constipation and diarrhea (IBS-C) has >25% of their bowel movements associated with BCSS 1 or 2 and >25% of their bowels associated with BSFS 6 or 7 [2].

Thus, based on the described scale, taking into account the frequency of occurrence of one type or another, the doctor establishes the form of IBS. At the same time, it must be understood that the same patient during the natural course of their disease can move from one type of IBS to another.

References

1. Lacy BE, Fermin Mearin, Lin Chang et al. Bowel Disorders. Gastroenterology 2016; 150:1393–407.

2. Lovell RM, Ford AC. Global prevalence of and risk factors for irritable bowel syndrome: a meta-analysis. Clin Gastroenterol Hepatol 2012; 10:712–21.

3. I. V. Maev, S. V. Cheremushkin, and Yu. Irritable Bowel Syndrome. Roman criteria IV. About the role of visceral hypersensitivity and methods of its correction. Toolkit. M., 2016.

4. Hawrelak JA, Myers SP. Effects of two natural medicine formulations on irritable bowel syndrome symptoms: a pilot sutdy. J Altern Complement Med. 2010; 16(10):1065–71.

5.Tibble, JA; Sigthorsson, G.; Foster, R.; Forgacs, I.; Bjarnason, I. Use of surrogate markers of inflammation and Rome criteria to distinguish organic from nonorganic intestinal disease. Gastroenterology 2002, 123, 450–460.

6. Thompson, WG; Longstreth, G.F.; Drossman, D.A.; Heaton, KW; Irvine, EJ; Müller-Lissner, SA Functional bowel disorders and functional abdominal pain. Gut 1999, 45, II43–II47.

7. Brian E. Lacy and Nihal K. Patel. Rome Criteria and a Diagnostic Approach to Irritable Bowel Syndrome. J.Clin. Med. 2017, 6, 99.

8.Maev I.V., Cheremushkin S.V., Kucheryavy Yu.A., Cheremushkina N.V. irritable bowel syndrome. Roman criteria IV. Consilium Medicum. 2016; 18(8): 79-85.

9. Myazin R.G. Irritable bowel syndrome: from diagnosis to treatment. Medical advice. 2016-9-130-133.

10. Andreev D.N., Dicheva D.T. Optimizing the treatment of patients with irritable bowel syndrome: focus on improving compliance. Medical advice. 2019-3-118-124.

Dicheva D.T., Andreev D.N., Shcheglanova M.P., Partsvania-Vinogradova
 E.V. Irritable Bowel Syndrome. In light of the Rome Criteria IV Revision (2016).
 Medical advice. 2018-3-60-66.

12. Makhov V.M., Balakhonov A.A., Isaikina M.A., Doronina Yu.A. Inflammation factor in the clinical picture and therapy of irritable bowel syndrome. Medical Council. 2018-14-67-72.

13. Sakhautdinova G.M., Asanbaeva K.E., Nagaeva R.R. Modern ideas about the etiology of irritable bowel syndrome. Medical advice. 2019-3-152-155.

14. Sheptulin A.A., Vize-Khripunova M.A. New in the etiology and pathogenesis of irritable bowel syndrome. Clinical medicine. 2016-94-2-92-96.

15. Rose LS Soares. Irritable bowel syndrome: A clinical review. World Journal of Gastroenterology. 2014-20-34-12144-12160.

16. Vahedi H., Ansari R., Mir-Nasseri MM., Jafari E. Irritable bowel syndrome: A review article. Middle East Journal of Digestive Diseases. 2010-2-2-66-77.

17. Kristen Ronn Weaver, Gail D'Eramo Melkus, Wendy A Henderson. Am J Nurse. 2017-117(6): 48-55.

18. Lekha Saha. World Journal of Gastroenterology. 2014-20-22-6759-6773.

19. Pogromov A.P., Mnatsakanyan M.G., Tashchyan O.V. The prevalence of irritable bowel syndrome. Clinical medicine. 2016-94-11-869-874.

20. Stepanov Yu.M., Budzak I.Ya. The role of visceral hypersensitivity in the development of irritable bowel syndrome. Gastroenterology. 2018:52(2):104-108.

21. Vasiliev Yu.V. irritable bowel syndrome: modern aspects of diagnosis and therapy. Medical advice. 2014-4-72-77.

22. Svistunov A.A., Osadchuk M.A., Osadchuk A.M., Butorova L.I. Rome criteria for irritable bowel syndrome IV revision (2016): what's new? Clinical medicine. 2017-95-11-987-993.

23. Karaulko I.V. Irritable Bowel Syndrome. Journal of Grodno State Medical University. 2011-4-85-89.

24. Serebrova S.Yu., Prokofiev A.B., Zhuravleva M.V., Eremenko N.N. Rome IV criteria for functional diseases of the gastrointestinal tract in assessing the interchangeability of drugs: the view of a clinical pharmacologist. 2017-4-221-227.

25. Abdullayev R. B., Makhmudova L.I. Features of Chemical Elements in Various Forms of Irritable Bowel Syndrome // Annals of R.S.C.B., ISSN:1583-6258, Vol. 25, Issue 2, 2021, Pages. 2993 – 3000.

26. Abdullayev R.B., Makhmudova L.I. Micro elemental imbalance in irritable bowel syndrome and IBS correction. Academicia. Vol. 11, Issue 5, May 2021:655-662.

27. Abdullayev R.B., Makhmudova L.I., (2021). Assessment Of Clinical And Psychological Status And Quality Of Life Of Patients In Different Forms Of Irritable Bowel Syndrome. The American Journal of Medical Sciences and Pharmaceutical Research, 3(02), 127-134.

28. Makhmudova L.I, Akhmedova N.Sh. Irritable bowel syndrome: a new look at the problem // Academicia. 10.5958/2249-7137.2020.00983.0. 433-38.

29. Makhmudova L.I., Akhmedova N.Sh., Ergashov B.B. Clinical manifestation of irritable bowel syndrome. Art of medicine. International medical scientific journal. Vol. 1, Issue 2. 2021:24-33.

30. Makhmudova L.I., Ismatova M.N., Mukhamedjanova M.H., Sulaymonova G.A. Evaluation of microelement status and IBS correction with irritable bowel syndrome. New day in medicine. 2(34) 2021:325-331.

31. Makhmudova L.I., Shazhanova N.S., Akhmedova N.Sh., (2021). Clinical Features Of Irritable Intestinal Syndrome. The American Journal of Medical Sciences and Pharmaceutical Research, 3(04), 154-159.

32. Makhmudova L.I., Sharipov J.N. State of intestinal microflora in irritable bowel syndrome. Tematics journal microbiology. Vol.6, Issue 1. 2022:104-109.