

Quality of Life Impact and Special Issues in Women with Inflammatory Bowel Diseases

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Research

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Abstract

Background: The impact of inflammatory bowel diseases (IBD) on quality of life (QoL) of patients is significant and has important social and professional consequences.

Methods: We aimed to describe the patients' perspective regarding the impact of IBD on their overall QoL and to evaluate the differences between men and women. An observational cross-sectional study, that included 180 patients with IBD in clinical remission, was conducted. All the patients completed a number of 3 questionnaires in order to evaluate the general aspects of their QoL. A separate questionnaire was created regarding gender-specific issues in women with IBD encounter. Also, particular features such as the incidence of anemia and osteoporosis among IBD patients were documented. The data obtained were analyzed and compared between the two gender-classified groups.

Results: According to the Short Inflammatory Bowel Disease Questionnaire (SIBDQ), patients had a general perception of a good QoL, but the impact was higher in women. Fatigue and tiredness were severely perceived almost to the same degree regardless of gender, whereas anxiety and unemployment were more present in men. No significant differences in women with IBD during active disease and during disease remission were found.

Conclusions: The overall quality of life of IBD patients is affected in many aspects, leading to the deterioration of their social and professional lives, for both men and women, but some aspects remain gender-specific and require a personalized standard of care.

1. Background

Inflammatory bowel diseases (IBD) have a great impact on quality of life (QoL), both in men and women, and represent a substantial burden on the patient and caregivers. The consequences can be observed in the increased rates of depression, anxiety, chronic fatigue and reduced workforce participation seen in patients with Crohn's disease or ulcerative colitis [1, 2].

The impact of IBD on QoL is multifaceted, from direct physical distress because of symptoms, in addition to psychological impairment, financial burden and low work productivity [1]. People with IBD frequently require multidisciplinary care, including mental health care, that can help in developing adaptive coping mechanisms in order to reduce psychological distress.

Several studies have demonstrated that QoL is impaired in these patients in comparison with the general population [3, 4], but only a few evaluated the differences between men and women. As women are regarded as being more sensitive to social stigma of having a chronic disease and prone to anxiety and depression [5], we tend to think that women with IBD should be more affected than men, but are they really?

In order to answer this question, we aimed to make a comparison between women and men regarding the impact of IBD on their overall QoL. We tried to raise physicians' awareness on the gender-specific aspects women with IBD may have and the impact on their self-esteem, resilience and overall coping mechanisms on the daily basis.

2. Materials And Methods

We conducted an observational cross-sectional study, that included patients with IBD who attended our Gastroenterology and Interventional Endoscopy Department from January 2017 until January 2020.

Patients previously diagnosed with IBD, either ulcerative colitis or Crohn's disease, who achieved clinical remission under treatment with biologic agents, were included. Patients' consent for data registration and subsequent use in clinical research was obtained at the time of inclusion in the current study, after the approval of the local ethics committee was obtained. Each patient received a number of questionnaires regarding their perception on overall QoL, using the Short Inflammatory Bowel Disease Questionnaire (SIBDQ) [6], the degree of chronic fatigue using the Functional Assessment of Chronic Illness Therapy-Fatigue Scale (FACIT-F) [7], the level of anxiety and depression caused by numerous hospital admissions (HADS) [8] and the diseases' impact on work productivity. Furthermore, a particular questionnaire was created for women with IBD specifically, concerning issues such as menstruation (age of menarche, variations of menstruation symptoms during remission and active disease) and the possibility of obtaining a full term pregnancy with or without continuing the biologic treatment, the Special Issues of Women with IBD Questionnaire (SIWIBDQ) (Table 1).

Nevertheless, the incidence of osteoporosis and anemia among our patients was also documented from data provided by the periodical biological examinations and bone density scans of each patient.

A total of 180 adult patients achieved sustained clinical and biochemical remission under biologic treatment. Data regarding symptoms and patient-reported outcomes were gathered using the previously stated questionnaires, after a minimum of 6 months of persistent clinical remission. Clinical remission was defined by the absence of symptoms: resolution of diarrhea and bleeding, abdominal pain and weight loss. However, laboratory testing was performed for each patient in order to correlate clinical with biochemical remission. Laboratory tests included complete blood count, inflammatory markers (C-reactive protein), albumin level and faecal calprotectin levels. All patients included in this study had no leukocytosis on complete blood count tests and C-reactive protein and albumin levels were within normal range. Also, faecal calprotectin levels were normal.

The exclusion criteria for participation in this study were pre-existing diagnoses of depression or anxiety disorder and continuous or intermittent treatment with anti-depressant or anti-anxiety medications.

From a total of 180 patients who met the inclusion criteria, 65 were women and 115 were men, with an average age of 37 years. Therefore, two groups were created based on gender classification.

All patients included in this study completed the following questionnaires: (1) Short Inflammatory Bowel Disease Questionnaire (SIBDQ); (2) Functional Assessment of Chronic Illness Therapy-Fatigue (FACIT-F) and (3) Hospital Anxiety and Depression Scale (HADS). Women also completed the (4) Special issues of women with IBD Questionnaire (SIWIBDQ).

The SIBDQ is a disease-specific health-related quality of life (HRQoL) measuring tool, meant to evaluate four domains – bowel, emotion, social and daily activities, and general well-being. Its importance lies in the capacity to address key points in one's life, such as social life, psychological well-being, the degree of fatigue, the presence of depression and body image. The questionnaire contains 10 items, and each item has a 7-point scale from 1 to 7. The absolute SIBDQ score varies from 10 to 70, with higher values suggesting better QoL.

The Functional Assessment of Chronic Illness Therapy - fatigue (FACIT-fatigue) scale is an instrument designed to assess fatigue and tiredness and its impact on daily activities and functioning in patients with chronic diseases. This score contains 13 items, each item being ranked from 0 to 4 points, with a total score of 52 points. The instrument includes aspects such as tiredness, weakness, lack of energy, listlessness and the impact of these feelings on daily activities (for example sleeping and social activities). A score under 30 points is representative for severe fatigue and the higher the score, the better QoL patients have.

In assessing the QoL of patients included in this study, a questionnaire consisting of a 7-questions anxiety assessment and 7-questions depression assessment, the HADS score, was also used. Scoring for each item varies from zero to three, with three indicating highest anxiety or depression level. A total subscale score of more than 11 points out of a possible 21 suggests considerable symptoms of depression or anxiety, between 8 and 10 is borderline abnormal, and less than 7 is normal.

In order to fully understand the women-specific issues, we created the Special Issues of Women with IBD Questionnaire (SIWIBDQ), consisting of 4 questions regarding menarche age, menstruation symptoms assessment during clinical remission in comparison to active disease phase and pregnancy (Table 1).

All data obtained were analyzed and compared between the two gender-classified groups.

Table 1
Special issues of women with IBD Questionnaire (SIWIBDQ)

1. What was your menarche age?	
• 12–13 years	0
• < 12 years	1
• ≥ 14 years	2
2. How do you perceive menstruation symptoms during clinical remission of IBD versus active disease phase?	
• Menstruation symptoms are less severe	0
• Menstruation symptoms are the same	1
• Menstruation symptoms are more severe	2
3. Did you have a pregnancy since the biological treatment for IBD was initiated?	
• No	0
• Yes	1
4. If the answer to question 3 was yes, did you discontinue the biological treatment during pregnancy?	
• No	0
• Yes	1

Tick the box beside the reply that describes best your current situation. Please answer all the questions.

Scoring:

Total score: _____

Normal = 0–3; Abnormal (case) = 4–6.

3. Results

A total of 180 patients were included in this study. From these, 115 patients were men and 65 women, aged between 19–47 years, thus balancing the scale towards men, with a percentage of 63.89%. The majority of the patients had no other comorbid diseases (91.67%). Overall, median SIBDQ, FACIT-F, HADS-anxiety and HADS-depression for males were 53.52, 17.04, 5.21, and 4.56, respectively, and for women, 52.30, 18.53, 4.69, and 4.07. Demographic data including patients' clinical characteristics are shown in Table 2.

Table 2
Demographic and clinical characteristics of
patients with IBD.

Variables	N = 36
Median age, years	42 (19–47)
Males, n (%)	115 (63.89%)
Females, n (%)	65 (36.11%)
Other comorbid diseases, n (%)	
Diabetes mellitus	5 (2.78%)
Arterial hypertension	10 (5.55%)
None	165 (91.67%)
Unemployment	
Males, n (%)	40 (22.23%)
Females, n (%)	25 (13.89%)
Anemia	
Present, n (%)	15 (8.33%)
Absent, n (%)	165 (91.67%)
Osteoporosis	
Males, n (%)	0 (0%)
Females, n (%)	(8.33%)
Median SIBDQ	
Males	53.52
Females	52.30
Median FACIT-F	
Males	17.04
Females	18.53
Median HADS-anxiety	
Males	5.21
Females	4.69
Median HADS-depression	

Variables	N = 36
Males	4.56
Females	4.07
Median SIWIBDQ	1.85

The SIBDQ score

A very useful tool for measuring health-related QoL is the Short Inflammatory Bowel Disease Questionnaire (SIBDQ). When asked to fulfill the SIBDQ questionnaire, women included in this study (36.11% from the total of patients) obtained total scores ranging between 38 and 67 points, with a median score of 52.30 points. Most of the patients were men (63.89%) and in this subgroup similar results were observed: their scores varied from 27 to 68 points and the median score was 53.52 points. High median scores were identified in both subgroups, which means that patients had a general perception of good QoL, but the impact was higher in women who obtained lower scores than men.

The FACIT-F score

The impact of the disease on the perception of fatigue was evaluated for the 115 male subjects in the study: the scores in this category of patients varied from 7 to 42, with a median score of 17.04 points. Women who completed this questionnaire and graded their symptoms obtained scores from 9 to 33 points, the median score among them being 18.53 points.

Therefore, fatigue and tiredness were severely perceived by the two groups almost to the same degree.

The HADS score

The HADS score can be divided into two subgroups HADS-anxiety and HADS- depression. The majority of the patients in our study were within the normal range, without significant differences between men and women for both the HADS-anxiety (78.26% vs. 76.92%) and HADS-depression (78.26% vs. 84.61%). In the borderline category, we encountered more women than men (15.38% vs. 4.34%) in the HADS-anxiety subgroup, whereas in the HADS-depression subgroup there where only men (17.40%). In the abnormal category there were only 5 patients for the HADS-anxiety and 3 patients for HADS-depression. But here, in comparison, the gender scale is inverted, as more men than women felt severe anxiety (17.4% men, 7.70% women) (Table 3) and more women than men felt severe depression (15.39% vs. 4.34%) (Table 4).

Table 3
HADS-anxiety score.

	Women percentage (number of cases)	Men percentage (number of cases)
Normal	76.92% (50)	78.26% (90)
Borderline abnormal	15.38% (10)	4.34% (5)
Abnormal	7.70% (5)	17.4% (20)
Total	100% (65)	100% (115)

Table 4
HADS-depression score.

	Women percentage (number of cases)	Men percentage (number of cases)
Normal	84.61% (55)	78.26% (90)
Borderline abnormal	0% (0)	17.40% (20)
Abnormal	15.39% (10)	4.34% (5)
Total	100% (65)	100% (115)

Impacts on work activity and daily activities

Regarding the employment status and work productivity of IBD patients, we observed that more than one third were unemployed, more men than women (22.23% vs. 13.89%). Regarding the work productivity and number of hours missed from work due to bowel problems, the majority (56%) of the patients, all women, did not miss one single hour from work. Only men (8%) had hours missed from work because of bowel problems and overall malaise.

Special issues of women with IBD and biologic agents' safety during pregnancy

Regarding specific issues of women with IBD such as menstruation, the mean age of menarche was between 12 and 13 years (61.55%), the same as the general population [9]. Moreover, we did not find significant differences in women with IBD between dysmenorrhea during active disease and during disease remission, as 84.62% (55 patients) felt equally painful periods (Table 5).

Although 77% of women have never had a pregnancy prior to diagnosis, 23% were able to deliver a full-term baby without complications, of which 66.60% continued the biologic treatment during pregnancy and only 33.40% discontinued it.

Table 5
Menarche and the evaluation of dysmenorrhea during active disease

Age of menarche	Number of patients (percentage)
11 years	15 (23.07%)
12 years	25 (38.48%)
13 years	15 (23.07%)
14 years	10 (15.38%)
Dysmenorrhea during active disease	
More painful	5 (7.69%)
Equally painful	55 (84.62%)
Less painful	5 (7.69%)
Pregnancy	
No pregnancies	50 (77%)
At least one pregnancy	15 (33%)
Continued biological treatment during pregnancy	10 (66.60%)
Discontinued biological treatment during pregnancy	5 (33.40%)

Anemia and osteoporosis among IBD patients

The topic of anemia and osteoporosis in patients with IBD receives little attention when analyzing extraintestinal disease complications. In our study, we observed that 92% of our patients have no anemia, which shows that the disease is controlled and the patients respond well to iron treatment, and only 9% of patients, all women, had osteoporosis.

4. Discussion

The results of this study focused on the psychological impact of IBD, demonstrating that although both men and women obtained high median scores according to the SIBDQ questionnaire, therefore having a general perception of an optimum QoL, the impact was higher for women, who obtained lower scores in comparison with men.

A study published in 2020 reports similar results: a lower QoL among women, compared to men diagnosed with IBD [10]. This finding corresponds to data observed in the general population, more precisely a higher rate of absenteeism from work and a more frequent use of health care services among women compared to men [10, 11]. Another important aspect that contributes to a reduced QoL is the

presence of physical asthenia. Compared to men, women report a higher frequency and intensity of physical asthenia. This symptom is independent from the activity of the disease and the presence or absence of anemia, one of the causal factors [12–14].

However, a surprising feature pointed out by this study was that more than one third of the patients (36.1%) are unemployed, because of various reasons. Even though they are predominantly young adults, living with IBD is definitely a challenge because work productivity and employment prospects are diminished, both directly by the symptoms and complications of the disease, and, indirectly, by the psychological distress patients have to cope with, making it difficult to get and, especially, to keep a workplace.

A study conducted in Norway reports higher rates of unemployment and sick leave among patients diagnosed with IBD when compared to the general population [15]. Moreover, the unemployment rate was slightly higher in women compared to men (12.4% vs. 11.1%) [16]. The unemployment rate was higher in patients diagnosed with ulcerative colitis (13.2%) compared to those diagnosed with Crohn disease (8.7%) [15]. The mean unemployment age in patients with ulcerative colitis was 55.3 years, compared with 42.2 years in those with Crohn disease [15].

Even if fatigue and tiredness are more and more present in the general population due to a constant need to keep up with each other, especially in the working environment, for IBD patients it represents a daily burden, regardless of their gender [16]. According to the FACIT-F score, both women and men were severely affected, obtaining similar scores.

Numerous studies show that QoL is impaired in these patients in comparison with the general population. Also, psychological distress doesn't always correlate with the activity and severity of the disease, because even during clinical remission patients may experience impairment on QoL [1–4].

Disease chronicity requires development of self-management and coping skills over time, therefore it is not surprising that IBD patients have an increased risk of mental health issues, including depression, which is an important problem patients often experience; therefore, it is essential for the caregiver to recognize these diseases in early phases. Recent studies show that the relationship between depression and chronic disease is bidirectional, with poor health leading to poor self-management, as well as poor self-management leading to poor health [17, 18].

The ongoing problem depression and anxiety pose in IBD patients is outlined by this study using the HADS anxiety and depression questionnaire, which showed that anxiety is more present in men than women, whereas depression was encountered more in women.

In addition to the huge impact that IBD has on QoL, people living with IBD face numerous other challenges. These include social stigma of having a chronic disease, affordability of medications, diminished employment prospects, difficulty with leisure time and travelling, limited community-based

supports and inequitable access to health care specialists and services [16, 18]. Reducing these difficulties should be a priority for physicians and health care providers, to improve QoL of their patients.

During flares, symptoms of IBD (both gastrointestinal and systemic symptoms) disrupt immediate activities and cause worry and embarrassment to patients. These immediate effects result in a cascade of impacts on a patient's life and psychological well-being, which continue to have consequences even during clinical remission of the disease, but to a lower grade [19].

A 2016 systematic review analyzed 171 studies including 158,371 individuals and showed a prevalence rate of anxiety of approximately 20% and a prevalence rate of depression of 15% in patients with IBD [20]. Both mental disorders had a higher prevalence during the disease activity period compared to the remission period. The estimated anxiety prevalence in patients with IBD during the disease activity was 75.6%, compared to 20.5% in the general population. It is noteworthy that the anxiety prevalence has been shown to be higher in patients with active IBD compared to patients with diabetes or chronic obstructive pulmonary disease [20–22]. The estimated depression prevalence in patients with active disease was 40.7%, compared to 15.2% in the general population. Depressive disorders have also been diagnosed more frequently in patients with Crohn disease compared to those with ulcerative colitis [20].

Interestingly, we found that although both men and women obtained similar scores, men were more affected by anxiety, encountered more difficulties getting and, especially, keeping a workplace, which had a huge impact on their social life as well.

A key strength of this study was the evaluation of special issues women with IBD have, despite maintenance of remission under biologic agents. Moreover, by using accessible and easy to understand and answer questionnaires, we were able to obtain clinical data, as it is felt by each patient. An important limitation of this study is the small number of patients included, but the data obtained can be a starting point for further larger prospective multicentre studies.

Although women are more affected by depression and social stigma than men, they are more resilient and develop better coping mechanism, therefore obtaining a better QoL than men.

These results correspond to the existing data that highlight the development of better coping mechanisms in women, compared to men diagnosed with IBD [23, 24]. The data so far support a greater negative impact on the QoL of women with IBD compared to men diagnosed with the same conditions [25–27].

Nevertheless, we must highlight that, even during sustained clinical remission, these patients, regardless of gender, experience QoL impairment.

5. Conclusions

Our study results show that the QoL of patients with IBD is affected in many aspects, leading to the deterioration of their social and professional lives, regardless of their gender, even if the majority of

patients regarded themselves as having a good QoL. Moreover, data shows that differences between men and women are present; when asking the question if women are really more affected than men, we observed that, although women encounter more difficulties, they have an overall better QoL than men. Nevertheless, future longitudinal studies on the psychological impact of patients with IBD are required in order to help us better understand their gender-specific needs, create better measurement tools and improve their QoL.

6. Declarations

Ethics approval and consent to participate:

Ethical approval for this study was obtained from the ethics committee of the Emergency Clinical Hospital of Bucharest. All subjects included in the study were informed about the purpose of the study and its involvement, subsequently signing an informed consent.

Consent for publication:

Not applicable

Availability of data and materials:

The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Competing interests:

The authors declare that they have no competing interests.

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Authors contribution:

Conceptualization, G.C., M.I. and B.P.; methodology, A.B., E.R., O.P, G.G.; software, V.S. A.M.O.; validation, C.C.D., G.C. and M.I.; formal analysis, V.S., A.M.O.; investigation, E.R., B.P.; resources, G.G., A.M.O.; data curation, A.B., E.R., O.P; V.S.; writing—original draft preparation, A.B, E.R.; writing—review and editing, C.C.D, G.G.; visualization, C.C.D.; supervision, M.I. and C.C.D; project administration, G.C., M.I; funding acquisition, B.P. All authors have read and agreed to the published version of the manuscript.

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