

## **Inflammatory Bowel Disease: A Retrospective Study**

Qasim Razaq Radhi, DM, FICMS\* Jehad Radhi Al-Qamish, FRACP, MRCP\*\*

**Background:** Inflammatory bowel diseases (IBDs), Crohn's disease (CD) and ulcerative colitis (UC) are a group of chronic intestinal inflammatory conditions.

**Objective:** To evaluate inflammatory bowel diseases in Bahrain.

**Design:** Retrospective study.

**Setting:** Department of Medicine, Gastroenterology Unit, Salmaniya Medical Complex.

**Method:** One hundred patients with IBD were included in the study from July 2007 to January 2008. Data collected include characteristics of patients, clinical presentation, endoscopic findings, histopathology, radiologic studies, extent of disease, complications, exacerbations, hospital admissions, associated medical diseases and management, medical or surgical.

**Result:** IBD affected 55 (55%) females and 45 (45%) males. The presenting symptoms were: 64 (64%) had diarrhea, 69 (69%) had gastrointestinal bleed and 57 (57%) had abdominal pain. The main complication was intestinal bleeding in 47 (47%). Steroid dependency was seen in 21 (21%) patients. The following medications were used: Asacol (Mesalazine) in 83 (83%), Azathioprine (Imuran) in 50 (50%) and infliximab in 8 (8%) patients.

**Conclusion:** IBD affected more females than males. Diarrhea, gastrointestinal bleed and abdominal pain were the most common presenting symptoms. More than half of cases of UC involved the left side of colon, while in CD the majority of cases involved small and large bowels. Most common drugs used were Mesalazine, Azathioprine and Infliximab.

*Bahrain Med Bull 2011; 33(2):*

Inflammatory bowel disease (IBD) is a group of inflammatory conditions of the large and small intestine; it includes Crohn's disease (CD) and ulcerative colitis (UC)<sup>1-3</sup>.

IBD have significant long-term morbidity and healthcare resource consequences. IBD is a common cause of gastro-intestinal disease in the Western world, the combined prevalence is 100-200/100,000<sup>4</sup>. In the Middle East, no sufficient data is available.

---

\* Senior Resident

\*\* Consultant

Department of Medicine  
Gastroenterology Division  
Salmaniya Medical Complex  
Kingdom of Bahrain  
Email: kasimrra@hotmail.com

The exact cause of IBD remains unknown, but several theories have been proposed. Hygiene theory suggests that an alteration in the microbial environment of patients facilitates the evolution of chronic immune-mediated diseases and development of IBD. Recent studies showed that there is an increased incidence of IBD because of exogenous infections, use of antibiotics and diet. Medical treatment options have rapidly expanded in recent years.

Current medical therapy is facilitative and supportive rather than curative. The principles of medical treatment are approximately the same for ulcerative colitis and Crohn's disease. Treatment emphasizes, besides drugs, the individuality of the therapeutic response<sup>5</sup>.

Current disease management guidelines have therefore focused on the use of anti-inflammatory agents, aminosalicylates, corticosteroids, immuno-modulators and more recently biological drugs in addition to surgery.

The aim of this study is to evaluate inflammatory bowel diseases in Bahrain.

## METHOD

One hundred patients with IBD were reviewed, from July 2007 to January 2008. Data collected include: patients' characteristics (age, sex, nationality, occupation, age at diagnosis, duration of disease, smoking and family history of IBD), clinical presentation, endoscopic findings, histopathology, radiologic studies (abdominal ultrasound, Barium studies and abdominal CT studies), extent of disease, complications (intestinal and extra intestinal), exacerbations, hospital admission, associated medical diseases and treatment, medical or surgical.

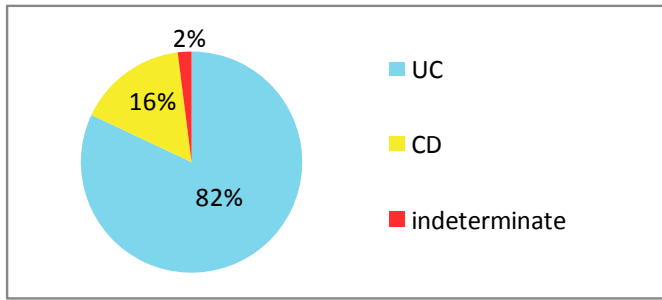
## RESULT

One hundred patients were reviewed. Patients' characteristics are shown in the table 1. Fifty-eight (58%) patients were 30-50 years old. IBD affected 55 (55%) females and 45 (45%) males. Eighty-five (85%) patients were Bahrainis. Forty-nine (49%) patients were not working. Most of the employed patients are semiskilled and 91 (91%) were indoor workers.

**Table 1: Characteristics of Patients with IBD**

Variables	Number and Percentage
<b>Age (Years)</b>	
Indoor Work	91 (91%)
Outdoor Work	9 (9%)
31-40	31 (31%)
41-50	27 (27%)
51-60	13 (13%)
61 and above	9 (9%)
<b>Sex</b>	
Male	45 (45%)
Female	55 (55%)
<b>Nationality</b>	
Bahraini	85 (85%)
Non-Bahraini	15 (15%)
<b>Occupation</b>	
Professional	6 (6%)
Skilled	16 (16%)
Semiskilled	23 (23%)
Worker	6 (6%)

Eighty-two (82%) patients had ulcerative colitis; 16 (16%) had Crohn's disease and 2 (2%) were undetermined, see Figure 1. Fifty-nine (59%) patients were diagnosed between 20-40 years. Forty-four (44%) patients had IBD between 0-5 years, 23 (23%) 6-10 years, twenty-four (24%) 11-15 years, 7 (7%) 16-20 years and 2 (2%) patients 20 years or more. Ninety-eight (98%) IBD patients were non-smoker. Most patients gave negative family history for IBD, see table 2.



**Figure 1: IBD Diagnosis**

**Table 2: Clinical Pattern of IBD**

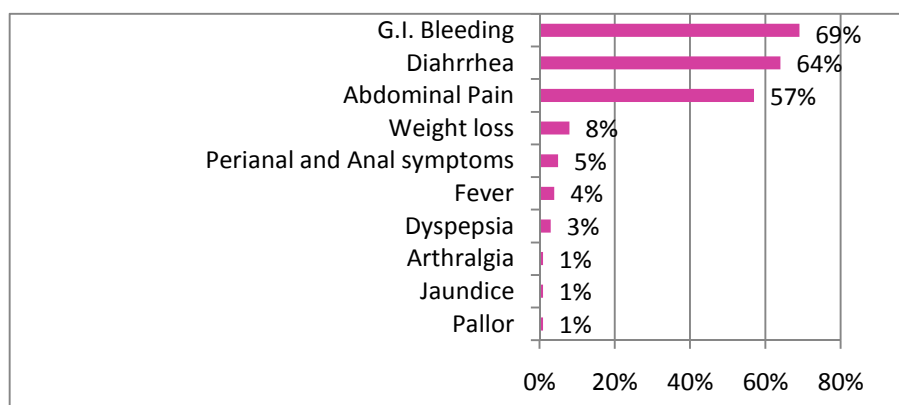
Variables	Number and Percentage
<b>Diagnosis</b>	
UC	82 (82%) M 39 (39%) F 43 (43%)
CD	16 (16%) M 5 (5%) F 11 (11%)
Indeterminate	2 (2%) M 1 (1%) F 1 (1%)
<b>Patient Age at Diagnosis</b>	
0-20	19 (19%)
21-30	29 (29%)
31-40	30 (30%)
41-50	16 (16%)
51-60	9 (9%)
61 and above	0 (0%)
<b>Duration of Disease (Years)</b>	
0-5	44 (44%)
6-10	23 (23%)
11-15	24 (24%)
16-20	7 (7%)
21 and above	2 (2%)
<b>Smoking</b>	
Smokers	2 (2%)
Average Duration of Smoking (yrs)	2.2 (2.2%)
<b>Family History of IBD</b>	
<b>Ulcerative Colitis</b>	
Father	1 (1%)
Mother	1 (1%)
Brother	2 (2%)
Sister	1 (1%)
<b>Crohn's Disease</b>	
Father	0 (0%)
Mother	0 (0%)
Brother	2 (2%)
Sister	0 (0%)

The presenting symptoms were: 64 (64%) had diarrhea, 69 (69%) had gastrointestinal bleed and 57 (57%) had abdominal pain. Less common symptoms were: 8 (8%) had weight loss, 5 (5%) had perianal and anal symptoms, 4 (4%) had fever and 3 (3%) had dyspepsia, see table 3 and Figure 2. Ninety-four (94%) patients had colonoscopy, 32 (32%) had esophagogastroduodenoscopy (OGD) and 18 (18%) had sigmoidoscopy.

**Table 3: Presentation of IBD**

Variables	Number and Percentage
-----------	-----------------------

Diarrhea	64 (64%)
GI Bleeding	69 (69%)
Abdominal Pain	57 (57%)
Others	23 (23%)
Weight Loss	8 (8%)
Perianal and Anal symptoms	5 (5%)
Fever	4 (4%)
Dyspepsia	3 (3%)
Arthralgia	1 (1%)
Jaundice	1 (1%)
Pallor	1 (1%)

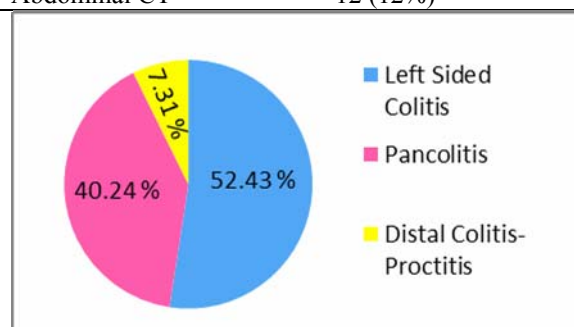


**Figure 2: Presentation of IBD**

Histopathology was done for 94 (94%) patients, barium study for 20 (20%), abdominal ultrasound for 17 (17%) and abdominal CT for 14 (14%), see table 4. Forty-three (52.4%) of UC patients involved the left side of the colon, 33 (40.2%) had pancolitis, and 6 (7.3%) had distal colitis, see figure 3.

**Table 4: Diagnostic Procedures**

Variables	Number and Percentage
<b>Endoscopic procedures</b>	
OGD	32 (32%)
Colonoscopy	94 (94%)
Limited Colonoscopy	6 (6%)
Sigmoidoscopy	18 (18%)
Histopathology Study	94 (94%)
Abdominal Ultrasound	17 (17%)
Barium Study	20 (20%)
Abdominal CT	12 (12%)

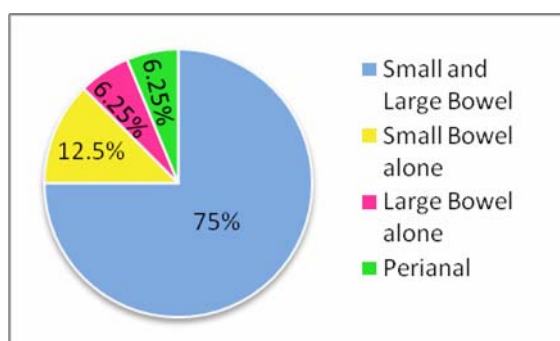


### Figure 3: Extent of UC

Twelve (75%) of CD involved small and large bowels, 2 (12.5%) involved small bowel alone, one patient (6.25%) involved large bowel alone and one patient (6.25%) involved perianal area, see table 5 and figure 4. Forty-seven (47%) had intestinal bleeding, 14 (14%) had sepsis, 10 (10%) had polyps, 6 (6%) had fistula, 4 (4%) had stricture and 3 (3%) had obstruction.

**Table 5: Extent of Disease**

Disease	Number and Percentage
<b>Ulcerative Colitis</b>	82 (82%)
Distal Colitis -Proctitis	6 (7.3%) of UC
Left Sided Colitis	43 (52.4%) of UC
Pancolitis	33 (40.2%) of UC
<b>Crohn's Disease</b>	16 (16%)
Small Bowel alone	2 (12.5%) of CD
Large Bowel alone	1 (6.25%) of CD
Small and Large Bowel	12 (75%) of CD
Perianal	1 (6.25%) of CD



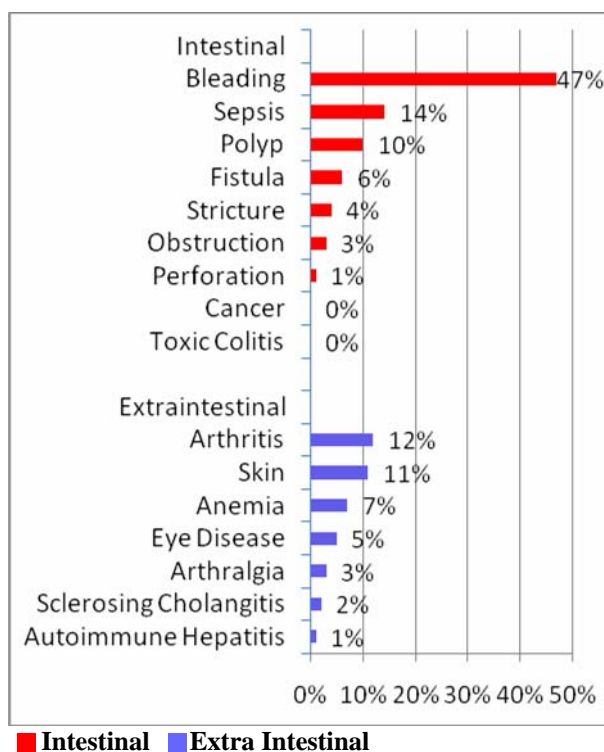
**Figure 4: Extent of CD**

Twelve (12%) patients had arthritis, 11 (11%) had skin lesions and 7 (7%) had anemia. Sclerosing cholangitis was seen in 2 (2%) patients and autoimmune hepatitis was seen in one patient (1%), see table 6 and figure 5.

**Table 6: Complications of IBD**

Intestinal	Number and Percentage
Toxic Colitis	0 (0%)
Obstruction	3 (3%)
Stricture	4 (4%)
Bleeding	47 (47%)
Perforation	1 (1%)
Fistula	6 (6%)
Sepsis	14 (14%)
Cancer	0 (0%)

Polyp	10 (10%)
<b>Extraintestinal</b>	
Sclerosing cholangitis	2 (2%)
Autoimmune hepatitis	1 (1%)
Arthritis	12 (12%)
Skin	11 (11%)
<b>Other</b>	
Eye disease	5 (5%)
Arthralgia	3 (3%)
Anemia	7 (7%)



**Figure 5: Complication of IBD**

The average exacerbation of IBD was once per year in 27 (27%) patients, two exacerbations per year were seen in 13 (13%) patients, once in two years in 19 (19%) and once every 3 to 5 years in 20 (20%). Fifty-two (52%) patients required admission to hospital, see table 7. The following diseases were associated with IBD: diabetes Mellitus was found in 11 (11%) patients, hypertension in 10 (10%), G6PD deficiency in 6 (6%), sickle cell trait in 6 (6%), hyperlipidemia in 4 (4%), and sickle cell disease in one patient (1%), see table 8.

**Table 7: Average Attacks of Exacerbation and Admission of IBD Patients**

<b>Exacerbation</b>	<b>Number and Percentage</b>
No exacerbation (0)	17 (17%)
one attack every 6-10 years	1 (1%)
once every 3-5 years	20 (20%)
once every two years	19 (19%)
1 per year	27 (27%)
2 per year	13 (13%)
3 per year	3 (3%)
<b>Admission per year</b>	
Not Admitted (0)	48 (48%)
One every 6-10 years (0.1)	5 (5%)
Once 3-5 years (0.25)	15 (15%)
Once every two years (0.5)	5 (5%)
1 per year (1)	23 (23%)
2 per year (2)	2 (2%)
3 per year (3)	2 (2%)

**Table 8: Associated Diseases**

<b>Disease</b>	<b>Number and Percentage</b>
Diabetes Mellitus	11 (11%)
Hypertension	10 (10%)
Ischemic Heart Disease	3 (3%)
Rheumatoid Arthritis	1 (1%)
<b>Others</b>	
G6PD decreased activity	6 (6%)
Sickle Cell Trait	6 (6%)
Hyperlipidemia	4 (4%)
Irritable Bowel Syndrome	3 (3%)
Hypothyroidism	3 (3%)
Osteoporosis	3 (3%)
Sickle Cell Disease	1 (1%)

Twenty-one (21%) patients were steroid dependent (steroid dependence was defined as a requirement for steroid therapy  $\geq 10$  mg/day during the preceding six months, with at least two attempts to discontinue the medication)<sup>6</sup>. Six (6%) patients had GI surgery, 3/6 patients (50%) had small bowel surgery, 3/6 patients (50%) had perianal surgery. Eight (8%) patients had endoscopic polypectomy.

The following drugs were used: Asacol (Mesalazine) in 83 (83%) patient, Imuran (Azathioprine) in 50 (50%), Steroid in 21 (21%), Asacol suppository in 17 (17%), Pentasa (Mesalazine) in 13 (13%), Infliximab in 8 (8%) and Folic acid in 71 (71%), see table 9.

**Table 9: Management**

<b>Steroid Dependency</b>	<b>Number and Percentage</b>
Steroid Dependent	21 (21%)
Steroid Non-Dependent	79 (79%)
<b>Endoscopic Polypectomy</b>	8 (8%)
Surgery(All surgeries)	24 (24%)
GI Surgery	6 (6%)
No Surgery	76 (76%)
<b>Type of Surgery</b>	
Colectomy	0 (0%)
Small Bowel Surgery	3 (3%) (50% of GI surgeries)
Perianal Surgery	3 (3%) (50% of GI surgeries)

Other Surgery ( Not GIT)	18 (18%)
<b>Types of Drugs Used in Treatment</b>	
Asacol (Mesalazine)	83 (83%)
Pentasa (Mesalazine)	13 (13%)
Steroid	21 (21%)
Immunosuppressive - Imuran	50 (50%)
Infliximab	8 (8%)
Folic Acid	71 (71%)
<b>Other</b>	
Omeprazole	14 (14%)
Iron	11 (11%)
Librax	5 (5%)
Asacol Suppository	17 (17%)
Salazopyrin	5 (5%)
Steroid Enema	1 (1%)
Anti TB	2 (2%)

## DISCUSSION

This study showed that age ranged 30 to 50 years, a mean of 40.69 years. A review study, the median age of onset of IBD was 35 years<sup>7</sup>. In another study, the age of patients showed biphasic distribution with two peaks between 20 and 30 and 50 and 70 years<sup>8</sup>.

In this study, females are affected more than males, for both types of IBD, UC and CD. Other studies showed that men and women at similar risk for UC. In CD, males are affected more than females, while in Western population females are affected more than males<sup>7</sup>. In a study in Lebanon, more males have been affected than females in CD (69.3%) and in UC (61.4%)<sup>9</sup>.

Fifty-one (51%) patients were employed; ninety-one (91%) were indoor workers. In a study, the mortality was higher in sedentary indoor workers compared to farmers and construction workers<sup>10</sup>. Sonnenberg suggested that employment involving outdoor air and physical activity is protective against IBD<sup>11</sup>.

In this study, the age at diagnosis of IBD was between 20-40 years, other studies showed that the peak age was 15-30 years old, although IBD can occur at any age<sup>10</sup>.

In this study, most patients were non-smoker; this is similar to other studies where smoking prevalence was significantly lower in UC patients than in general population (9% versus 28%)<sup>12</sup>.

In this study, the common presenting symptoms were Diarrhea, gastrointestinal bleed and abdominal pain, less common symptom was weight loss. The cardinal symptom of UC is bloody diarrhea; symptoms of colicky abdominal pain, urgency or tenesmus may be present. Symptoms of CD are more heterogenous, but typically include, abdominal pain, diarrhea and weight loss<sup>13</sup>.

In this study, the following investigative procedures were performed: colonoscopy 94 (94%), sigmoidoscopy 18 (18%) and histopathology 94 (94%). Compared with other studies diagnostic criteria were endoscopy, histopathology and radiology<sup>14,15</sup>.

In this study, Forty-three (52.4%) of UC patients involved the left side of the colon, 33 (40.2%) had pancolitis, and 6 (7.3%) had distal colitis; our result is comparable with other studies<sup>14</sup>.



In this study, CD involved small and large bowel in 12 (75%), small bowel alone in 2 (12.5%), large bowel alone in one patient (6.25%) and perianal area in one patient (6.25%). Butt et al, in a study of CD showed that ileo-colonic disease affected 52%, ileal disease 24% and colonic 24%<sup>16</sup>.

Intestinal complications of IBD in this study were: forty-seven (47%) had intestinal bleeding, 14 (14%) had sepsis, 10 (10%) had polyps, 6 (6%) had fistula, 4 (4%) had stricture and 3 (3%) had obstruction. In a study of patients who had Crohn's disease, gastrointestinal complications particularly abdominal sepsis, intestinal ischemia, and intestinal hemorrhage accounted for the excess of mortality<sup>17</sup>. Twelve (12%) patients had arthritis, it was the most common extraintestinal complication seen in our study, 11 (11%) had skin lesions and 7 had (7%) anemia, compared with other studies, joints manifestations were the most common<sup>18,19</sup>.

In our study, average exacerbation of IBD was once per year in 27 (27%) patients, while in another study relapses was seen in 32.9%, that study suggested that stressful life events do not trigger exacerbations in patients with IBD<sup>20</sup>. Elderly patients with IBD using mesalazine and corticosteroid lead to adverse effects including osteoporosis, bone fractures, changes in mental status, diabetes and hypertension<sup>21</sup>.

In this study, steroid dependency was seen in 21 (21%) patients with IBD, which is comparable to other studies<sup>22,23</sup>. In patients with steroid dependency, it might be advisable to consider the use of biological agent (such as infliximab) for better control of IBD.

## CONCLUSION

**IBD affects females more than males. Eighty-two (82%) patients had Ulcerative Colitis, 16 (16%) had Crohn's disease and 2 (2%) were undetermined. Diarrhea, gastrointestinal bleed and abdominal pain were the most common presenting symptoms. More than half of the cases of UC involved left side of colon, while in CD, the majority of cases involved small and large bowels. 21 (21%) patients were steroid dependent. The most common drugs used was Asacol (Mesalazine), Imuran (Azathioprine) and Infliximab. This study is limited and does not reflect the whole problem of IBD in Bahrain.**

---

**Author Contribution:** All authors share equal effort contribution towards (1) substantial contributions to conception and design, acquisition, analysis and interpretation of data; (2) drafting the article and revising it critically for important intellectual content; and (3) final approval of the manuscript version to be published. Yes

**Potential Conflicts of Interest:** No

**Competing Interest:** None, **Sponsorship:** None

**Submission date:** 15.03.2011 **Acceptance date:** 16 April 2011

**Ethical approval:** Approved by the GI unit

## REFERENCES

1. Baumgart DC, Carding SR. Inflammatory Bowel Disease: Cause and Immunobiology. *Lancet* 2007; 369(9573): 1627-40.
2. Baumgart DC, Sandborn WJ. Inflammatory Bowel Disease: Clinical Aspects and Established and Evolving Therapies. *Lancet* 2007; 369(9573): 1641-57.
3. Xavier RJ, Podolsky DK. Unraveling the Pathogenesis of Inflammatory Bowel Disease. *Nature* 2007; 448(7152): 427-34.
4. Calkins B, Mendeloff A. The Epidemiology of Idiopathic Inflammatory Bowel Disease. In: Krisner JB, Shorter RG, eds. *Inflammatory Bowel Disease*. Baltimore: Williams and Wilkins, 1995; 31-68.
5. Kirsner JB. Inflammatory Bowel Disease. Part II: Clinical Therapeutic Aspects. *Dis Mon* 1991; 37(11): 669-746.
6. Ardizzone S, Maconi G, Russo A, et al. Randomised Controlled Trial of Azathioprine and 5-Aminosalicylic Acid for Treatment of Steroid Dependent Ulcerative Colitis. *Gut* 2006; 55: 47-53.
7. Suk-Kyon Yang, Edward VL Jr, William JS. Epidemiology of Inflammatory Bowel Disease in Asia. *Inflammatory Bowel Disease* 2001; 7(3): 260-70.
8. Tozun N, Atug O, Imeryuz N, et al. Clinical Characteristics of Inflammatory Bowel Disease in Turkey: A Multicenter Epidemiologic Survey. *J Clin Gastroenterol* 2009; 43(1): 51-7.
9. Abdul-Baki H, ElHajj I, El-Zahabi LM, et al. Clinical Epidemiology of Inflammatory Bowel Disease in Lebanon. *Inflamm Bowel Dis* 2007; 13(4):475-80.
10. Hanauer SB. Inflammatory Bowel Disease: Epidemiology, Pathogenesis, and Therapeutic Opportunities. *Inflamm Bowel Dis* 2006; 12 (Suppl 1): S3-9.
11. Sonnenberg A. Occupational Distribution of Inflammatory Bowel Disease among German Employees. *Gut* 1990; 31(9): 1037-40.
12. Miquel R, Kevin EK, Onki C, et al. Cigarette Smoking and Age at Diagnosis of Inflammatory Bowel Disease. *Inflamm Bowel Dis* 2005; 11(1): 42-7.
13. Carter MJ, Lobo AJ, Travis SPL. Guidelines for the Management of Inflammatory Bowel Disease in Adults. *Gut* 2004; 53(suppl 5): v1-16.
14. Saro GC, Lacort FM, Arguelles FG, et al. Epidemiology of Chronic Inflammatory Bowel Disease in Gijon, Asturias. *Gastroenterol Hepatol* 2001; 24(5): 228-35.
15. Saro GC, Riestra MS, Sanchez FR, et al. Epidemiology in Inflammatory Bowel Disease in Five Areas of Asturias, Spain. *An Med Interna* 2003; 20 (5): 232-8.
16. Butt MT, Bener A, Al-Kaabi S, et al. Clinical Characteristics of Crohn's Disease in Qatar. *Saudi Med J* 2005; 26(11): 1796-9.
17. Francisco M, Mohammed AQ, Bret AL. Severe Complications of Inflammatory Bowel Disease. *Med Clin N Am* 2008; 671-86.
18. Mendoza JL. Extraintestinal Manifestations in Inflammatory Bowel Disease: Differences between Crohn's Disease and Ulcerative Colitis. *Med Clin (Barc)* 2005; 125(8): 297-300.
19. Tozun N. Clinical Characteristics of Inflammatory Bowel Disease in Turkey: a Multicenter Epidemiologic Survey. *J Clin Gastroenterol* 2009; 43(1): 51-7.
20. Vidal A. Life Events and inflammatory Bowel Disease Relapse: A Prospective Study of Patients Enrolled in Remission. *Am J Gastroenterol* 2006; 101(4): 775-81.
21. Picco MF, Cangemi JR. Inflammatory Bowel Disease in the Elderly. *Gastroenterol Clin North Am*. 2009; 38(3):447-62.
22. Fish D, Kugathasan S. Inflammatory Bowel Disease. *Adolesc Med Clin*. 2004; 15(1):67-90, ix. Review.
23. Katz JA. Treatment of Inflammatory Bowel Disease with Corticosteroid. *Gastroenterol Clin North Am* 2004; 33(2): 171-89.