

Evolutionary Profile of Chronic Inflammatory Bowel Disease at Brazzaville University Hospital in 2023: A Case Series of 26 Patients

Ngami RS^{1,2*}, Mikolélé Ahoui Apendi PC^{1,2}, Mimiesse Monamou JF^{1,2}, Mongo-Onkouo A^{1,3}, Itoua-Ngaporo NA^{1,2}, Motoula Latou PM^{1,2}, Ngalessami Mouakosso M¹, Adoua Céline Sandra¹, Ibobi Mauria Gilga¹, Deby Gassaye^{1,2} and Atipo Ibara BI^{1,2}

¹Gastroenterology and Internal Medicine Department, Brazzaville University Hospital, Republic of Congo.

²Faculty of Health Sciences, Marien Ngouabi University, Republic of Congo.

*Correspondence:

Ngami RS, Gastroenterology and Internal Medicine Department, Brazzaville University Hospital, Republic of Congo.

Received: 10 Feb 2024; Accepted: 15 Mar 2024; Published: 22 Mar 2024

Citation: Ngami RS, Mikolélé Ahoui Apendi PC, Mimiesse Monamou JF, et al. Evolutionary Profile of Chronic Inflammatory Bowel Disease at Brazzaville University Hospital in 2023: A Case Series of 26 Patients. *Gastroint Hepatol Dig Dis.* 2024; 7(1): 1-4.

ABSTRACT

Chronic inflammatory bowel disease (IBD) is rarely described in sub-Saharan Africa. The aim of this study was to evaluate the evolutionary profile of a series of 26 cases of IBD at the Brazzaville university hospital in 2023.

Patients and Methods: This was a cross-sectional analytical study conducted in the Gastroenterology department of Brazzaville university hospital over a 13-year period. IBD was diagnosed on the basis of clinical (chronic diarrhoea, rectal discharge), morphological (intestinal parietal thickening), endoscopic (acute or chronic lesions) and histological criteria. Severe forms were defined by the fact of having a pancolitis and ano-perineal lesions. The Wilcoxon test was used to compare variables at a threshold of 0.05.

Results: A total of 26 patients were included, representing 0.2% of hospitalizations. The sex ratio was 1.9. The median age was 37 (IQR 24-51), and nine patients were under 30. IBD included Crohn's disease (CD), ulcerative colitis (UC) and indeterminate forms in 10, 8 and 8 cases respectively. The median duration of symptoms before diagnosis was 2.2 years (IQR 0.1-6). Twelve patients had a severe form, including two cases of severe acute colitis. Undernutrition was observed in 16 patients, including 13 cases of severe undernutrition. By 2023, 17 out of 26 patients had been lost to follow-up, 5 had died and 4 were regularly monitored. The median time to death was 3.3 months (IQR 1.4-3.8). The presence of ano-perineal lesions ($p=0.0143$) and disease severity ($p=0.019$) were factors associated with death.

Conclusion: Keeping IBD patients in the care circuit is a multidisciplinary challenge to reduce the proportion of patients lost to follow-up, and improving their management.

Keywords

Crohn's disease, Ulcerative colitis, Evolutionary profile.

Introduction

Chronic inflammatory bowel diseases are complex nosological entities of unknown cause, most often evolving in intermittent flare-ups [1]. They are more prevalent in northern European and American countries, with a north-south gradient. [2]. In sub-Saharan Africa, the prevalence of IBD is low, and the disease is

little known, particularly among the general population. The lack of diagnostic resources and trained personnel partly explain the problems of IBD management in Africa. In Congo, an initial study carried out in 2015 described the epidemiological, diagnostic and therapeutic parameters of 9 cases over 13 years [3]. However, little is known about patient outcomes. The aim of this study was to assess the evolutionary profile of chronic inflammatory bowel disease at Brazzaville University Hospital in 2023.

Patients and Methods

We carried out a cross-sectional analytical study from September 2010 to October 2023, a period of 13 years, in the Gastroenterology and Internal Medicine Department of the Centre Hospitalier Universitaire (CHU) de Brazzaville. The study population consisted of patients hospitalized or followed-up for IBD, diagnosed on the basis of a combination of epidemiological (age, lifestyle, non-steroidal anti-inflammatory drugs), clinical (diarrhea, bloody emissions, abdominal pain, anemia, undernutrition), morphological (abdominal ultrasound and CT scan), endoscopic (gastroscopy, rectoscopy or ileocoloscopy) and histological (diffuse lymphoplasmacytic infiltrate of the chorion, polynuclear infiltrate and loss of mucosecretion in UC, epitheliogigantocellular granuloma without central necrosis in CD. We included all patients aged 18 and over, diagnosed with IBD and followed up in the department during the study period. Files suggesting IBD but containing no endoscopic examination were excluded. Progression profile included endoscopic mucosal healing, follow-up status, mean follow-up time, death, mean time to death and risk factors for death. Data were collected on a paper questionnaire, entered into Microsoft Excel 19 and analyzed using Epi info 7.2. The Fisher's exact test and the Wilcoxon test were used from a threshold of 0.05.

Results

Over a 13-year period, we selected 26 cases of patients hospitalized and followed in the department for chronic inflammatory bowel disease (IBD) out of a total of 12,064 hospitalizations, representing a prevalence of 0.2%. There were 17 men and 9 women, giving a *sex ratio* of 1.9. The median age was 37 (IQR 24-51), and 9/26 patients were under 30. The IBD diagnosis was Crohn's disease (CD), ulcerative colitis (UC) and indeterminate forms in 10, 8 and 8 cases respectively. One case of co-morbidity of UC and human immunodeficiency virus infection was noted. In this case, histology was essential in the diagnosis of UC. The median duration of symptoms before diagnosis was 2.2 years (IQR 0.1-6). Twelve cases of severe colitis were diagnosed with deep anorectal ulcerations (photo 1), including two cases of severe acute colitis with profuse hematochezia.

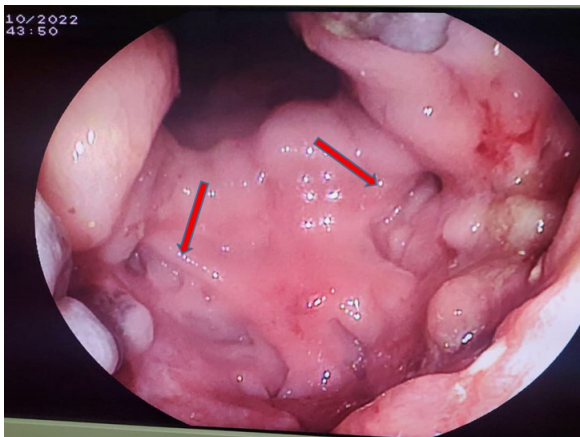


Photo 1: Anorectal hollowing ulcerations.

Source: Brazzaville University Hospital, department of Gastroenterology

Undernutrition was observed in 16 patients, including 13 cases of severe undernutrition and 3 cases of moderate undernutrition. Medical treatment consisted of Salazopyrine, Mesalazine, Prednisone and Azathioprine in 9, 7, 12 and 7 cases as initial therapy, and Salazopyrine, Mesalazine, Budesonide and Azathioprine in 9, 8, 2 and 7 cases as maintenance therapy. Patient follow-up was irregular. In three months of follow-up, 13 patients were regularly monitored, 11 were lost to follow-up and 2 died. In 2023, after 13 years of follow-up, 17 patients had been lost to follow-up, 5 had died and 4 were regularly monitored.

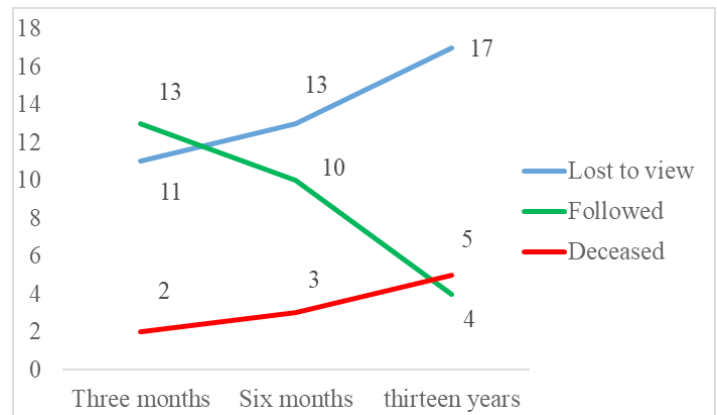


Figure 1: Evolutionary profile of IBD cases.

At 3 months, endoscopic mucosal healing (EMH) was observed in five out of 13 patients. In 2023, three of the four patients monitored had EMH. The median time to death was 3.3 months (IQR 1.4-3.8). Ano-perineal lesions and disease severity were associated with death (Table 2).

Table 1: Bivariate analysis between lost to follow-up status and IBD characteristics.

IBD features	Lost to view (n)		p	
	Yes	No		
Age (years)	< 30	4	4	0,2545
	≥ 30	13	5	
Gender	Female	4	5	0,1158
	Male	13	4	
APL*	Yes	1	3	0,1039
	No	16	6	
Severe IBD	Yes	11	3	0,1329
	No	6	6	
Undernutrition	Yes	9	7	0,2096
	No	8	2	
EIM**.	Yes	3	4	0,1585
	No	14	5	

* Ano-perineal lesion; ** Extra-intestinal manifestation.

Table 2: Bivariate analysis between death and IBD characteristics.

IBD features		Death		
		Yes	No	
Age (years)	< 30	2	6	0,4975
	≥ 30	3	15	
Gender	Female	2	7	0,5803
	Male	3	14	
PAL*	Yes	3	1	0,0143
	No	2	20	
Severe IBD	Yes	5	7	0,012
	No	0	14	
Undernutrition	Yes	3	13	0,6569
	No	2	8	
EIM**.	Yes	3	4	0,1014
	No	2	17	

* Ano-perineal lesion; ** Extra-intestinal manifestation.

Discussion

The prevalence of IBD in sub-Saharan Africa is difficult to assess. In Mali, 59 cases were reported in 3 years, representing 0.27 of hospitalizations and 6.1% of digestive pathologies, which was comparable to our study [4]. However, these authors included children in their study. Medhioub and colleagues noted an increase in the prevalence of IBD in patients over 60, but there was no difference in terms of clinical forms and surgery compared with younger patients [5]. Hassine and colleagues in Morocco and Siala et al. in Tunisia found that 13% and 14% respectively of people over 60 had IBD [6,7]. The average age was 48.2 ± 18.4 years, with extremes of 10 and 89 years, and a female predominance in Mali. The epidemiology of IBD is different in children, with a predominance of CD (65%) compared with UC (35%); a male predominance in CD and a female predominance in UC [8].

In 2015, nine cases of CD were reported in Congo, compared with ten cases in 2023 [3]. The hospital prevalence of CD does not appear to be increasing in our country. Bougouma and colleagues in Burkina Faso reported 20 cases of ulcerative rectocolitis, while Diouf and colleagues in Senegal reported 32 cases of UC out of 2667 recto-sigmoidoscopies and colonoscopies, i.e. 1.2% of low digestive endoscopic examinations. Moderate forms were mainly observed in the elderly, i.e. 82.3% of the population [7]. Half of our patients had moderate forms but were young.

Medication for IBD has been on the rise for several years [9]. In our country, available treatments are limited to conventional therapies, which limits the aggressive management recommended in severe forms, or in severe acute colitis [1,10]. Two patients underwent emergency surgery for small bowel perforation peritonitis during CD.

Follow-up of IBD patients is difficult. In our series, 17/26 cases of loss of sight were observed. The reasons for this were not evaluated in this study. For some authors, the reasons for poor compliance included the relationship established with the physician, depression, subjective quality of life, health status evaluation and the therapeutic benefit/treatment burden ratio [11,12].

Mortality is high in IBD patients compared with the general population. Gismera and colleagues found a mortality of 5%, and mortality risk factors vary from study to study, but death was more often related to CD than UC for Nocerino and colleagues [13,14]. Ano-perineal lesions and disease severity were the risk factors for death observed in our series. Dia and colleagues in Dakar reported two cases of severe acute colitis treated by conventional means and with lethal evolution [15]. The lack of biotherapy could be considered a risk factor for death in our context. However, the single-center nature of this study may be a source of patient selection bias.

Conclusion

Keeping IBD patients on the care circuit is a multidisciplinary challenge to reduce the proportion of patients lost to follow-up, and improving the management of severe forms with non-conventional treatments should change the prognosis of IBD at Brazzaville University Hospital.

References

1. Reynt V. Crohn's disease and its treatment. *Actual Pharm.* 2019; 58: 44-48.
2. Klotz C, Barret M, Dhooge M, et al. Hemorrhagic rectocolitis: diagnostic and therapeutic management. *Presse Médicale.* 2015; 44:144-149.
3. Mimiesse JF, Deby-Gassaye, Atipo-Ibara BI, et al. Crohn's disease: first description at Brazzaville University Hospital. *J Afr Hépatogastroentérologie.* 2015; 9: 73-75.
4. <https://www.bibliosante.ml/handle/123456789/4056>
5. Medhioub M, Khsiba A, Mahmoudi M, et al. Characteristics of chronic inflammatory bowel disease in the elderly. *Rev Médecine Interne.* 2023; 44: A234.
6. Hassine A, Hammami A, Ben Ameer W, et al. Chronic inflammatory bowel diseases of the elderly: epidemio-clinical, therapeutic and evolutionary profile. *Rev Médecine Interne.* 2022; 43: A205.
7. Siala A, Boudabous M, Chtourou L, et al. Epidemio-clinical profile of chronic inflammatory bowel disease in the elderly. *Rev Médecine Interne.* 2016; 37: A148.
8. Gowze-Rousseau C, Fumery M, Savoye G, et al. John Libbey Eurotext - Hepato-Gastro & Digestive Oncology - Epidemiology and natural history of chronic inflammatory bowel disease in children. *Hepato-Gastro Oncol Dig.* 2018; 25: 895-902.
9. Reenaers C, Louis E. New developments in the management of chronic inflammatory bowel disease. *Rev Médicale Liège.* 2022; 77: 323-329.
10. Beaugerie L. Chronic Inflammatory Bowel Diseases (CIBD): what place for conventional treatments? *Acta Endosc.* 2008; 38: 359-374.
11. Tahri N. Therapeutic compliance and chronic inflammatory bowel diseases. *Presse Médicale.* 2007; 36: 1236-1243.

-
12. Banovic I, Gilibert D, Olivier M, et al. Compliance and some of its determinants in chronic inflammatory bowel disease (IBD). *Prat Psychol.* 2010; 16: 157-172.
 13. Saro Gismera C, Lacort Fernández M, Argüelles Fernández G, et al. Mortality and causes of death in patients with chronic inflammatory bowel disease in Gijón, Asturias (Spain). *Rev Esp Enferm Dig.* 1999; 91: 199-208.
 14. Nocerino A, Feathers A, Ivanina E, et al. Mortality Risk of Inflammatory Bowel Disease: A Case-Control Study of New York State Death Records. *Dig Dis Sci.* 2019; 64: 1604-1611.
 15. Dia D, Cisse M, Diouf G, et al. Severe acute colitis: Two fatal cases in Dakar. *Med Sante Trop.* 2014; 24: 438-440.