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Infection (Covid-19) in Patients with Chronic Inflammatory Bowel Disease: A Case Series

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Abstract

COVID-19 is a Severe Acute Respiratory Syndrome caused by the Coronavirus (SARS-CoV-2) that was described in China in late 2019 and led to a global pandemic.

The devastating effect of this infection is due to its high contagion, although mild forms predominate, severe cases and mortality are very high especially in patients with comorbidities and immunosuppression, raising the question of infection in patients with IBD who are often impaired and on immunosuppressants or anti TNF. Due to the recent nature of the infection and the shortage of studies, we considered it interesting to share our IBD experience through a series of eight IBD patients who presented with coronavirus infection.

Keywords: COVID-19 · SARS-CoV-2 · Chronic inflammatory bowel diseases

Introduction

Severe Acute Respiratory Syndrome Coronavirus (SARS-CoV-2) infection was first detected in Wuhan province of Hubei State, China in December 2019 [1]. Subsequently, the number of cases increased dramatically in a short period of time to become a covid pandemic.

Coronaviruses are single-stranded RNA viruses that can rapidly mutate, change tissue tropism, and adapt to different epidemiological conditions [2,3]. COVID-19 symptoms are more severe in patients at risk and with comorbidities [4] including those with (IBD) [5]. To date, there are limited data on these patients. Therefore, there is a need for studies to assess the risk and clinical characteristics.

Our aim is to share the experience of IBD patients infected with covid by describing their clinical, therapeutic and evolutionary characteristics.

Case Series

Our series includes ten IBD patients infected with confirmed covid-19, including nine women and one man, with a mean age of 40 years \pm 10 years [26-53 years], five patients followed for UC and five patients followed for Crohn's disease. The median duration of evolution was 6 years [4-13].

The diagnosis of COVID-19 was established by Polymerase Chain Reaction (PCR) analysis on nasopharyngeal swabs and chest CT scans.

Seven out of ten patients were treated as outpatients, it was a non-severe coronavirus infection without respiratory repercussions (normal oxygen pressure, no pneumonia or acute respiratory distress syndrome) and all were in remission from their IBD disease including two patients in remission under 5ASA: a 53-year-old patient followed for 12 years for pancolitic UC

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Received: 05 December, 2024, Manuscript No. jibdd-24-154360; **Editor** assigned: 07 December, 2024, Pre QC No. P-154360; **Reviewed:** 19 December, 2024, QC No. Q-154360; **Revised:** 24 December, 2024, Manuscript No. R-154360; **Published:** 31 December, 2024, DOI: 10.37421/2476-1958.2024.9.225 and a 32-year-old patient followed for five years for ileocolic Crohn's disease inflammatory phenotype with anoperineal lesions, two patients in remission under azathioprine this is a 36-year-old patient followed for 10 years for ileal Crohn's disease stenosing phenotype with anoperineal manifestations operated (ileal resection and flattening of an anal fistula) and a 45-year-old patient followed since 2007 for pancolitic UC under azathioprine for 03 years, a fifth patient followed for 10 years for corticosteroid-dependent pancolitic UC in remission under 6MP and 5ASA, a sixth patient in remission under adalimumab is a 48-year-old patient followed for six years for pancolitic UC, and a seventh patient in remission under azathioprine and infliximab combo therapy, this is a 43-year-old patient followed for 5 years for ileocecal Crohn's disease, stenosing phenotype with anoperineal manifestation.

Results

Three patients were hospitalized, two of them due to severe respiratory symptoms, a 45-year-old patient followed for 5 years for pancolitic UC with extradigestive axial joint manifestation in remission under azathioprine and a 45-year-old patient followed since 2013 for Crohn's disease in remission under 5ASA, both patients were treated with hydroxychloroquine and antibiotics, the evolution was favorable with resumption of azathioprine and 5ASA after negative PCR covid. The third patient was hospitalized due to a profound deterioration in general condition and a manifest inflammatory syndrome, this is a 26-year-old patient followed for 5 years for ileocolic Crohn's disease fistulizing phenotype with anoperineal manifestation in remission under azathioprine. 5ASA, immunosuppressive and anti-TNF treatments were stopped except in two out of ten patients: one was taking azathioprine and the second was on infliximab Tables 1 and 2.

Discussion

During the COVID-19 pandemic, the management of IBD patients poses diagnostic and therapeutic challenges.

Indeed, patients with COVID-19 may present gastrointestinal symptoms such as diarrhea, abdominal pain, nausea/vomiting, anorexia that could wrongly suggest an IBD flare-up with possible inappropriate initiation of treatment that may include corticosteroids [6-9].

Thus, it is proposed to endoscopically document any IBD flare and test patients with diarrhea or bloody mucus emission [10,11] This will help distinguish an IBD flare from diarrhea due to SARS-CoV-2 infection and avoid

File Number	Gender	Age	MICI	MC 'Montreal' Phenotype	RCH Extension	Duration of Illness (years)	Comorbidities	Treatment of IBD
1	F	53	RCH	-	pan colitic	12	HTA	5 ASA
2	F	48	RCH	-	pan colitic	6	-	Adalimumab
3	Μ	45	RCH	-	pan colitic	5	-	Azathioprine
4	F	45	RCH	-	pan colitic	14	-	Azathioprine
5	F	48	RCH	-	pan colitic	10	-	Purinethol 5 ASA
6	F	36	MC	A2L1B2P operated	-	10	-	Azathioprine
7	F	43	MC	A3L3B2p	-	5	-	Azathioprine
8	F	45	MC	A3L3B1	-	7	-	Infliximab 5 ASA
9	F	26	MC	A2L3B3p	-	-	-	Azathioprine
10	F	32	MC	A2L3B1p	-	-	-	5 ASA

Table 1. Characteristics of our IBD patients infected with covid-19.

Table 2. Clinical characteristics, complications and treatments of our IBD patients with coronavirus.

File Number	Symptoms	Complications	Therapies for COVID-19	Evolution
1	Fever, cough Headache	No	Antibiotics Zinc, vit C	Outpatient Good progress
2	Fever, cough	No	Hydroxychloroquine Antibiotics	Outpatient Good progress
3	Fever, myalgia, sore throat	dyspnea	Hydroxychloroquine Antibiotics	Hospitalized Good progress
4	asymptomatic	No	Antibiotics Zinc, vit C	Outpatient Good progress
5	Bloody, mucous diarrhea	No	Antibiotics Zinc, vit C	Outpatient Good progress
6	Fever, cough, myalgia	No	Hydroxychloroquine Antibiotics	Hospitalized Good progress
7	Cough, myalgia	No	Antibiotics Zinc, vit C	Outpatient Good progress
8	Sd severe pseudo flu	dyspnea	Hydroxychloroquine Antibiotics	Hospitalized Good progress
9	AEG	No	Hydroxychloroquine Antibiotics	Hospitalized Good progress
10	Fever, myalgia, sore throat	No	Hydroxychloroquine Antibiotics	Outpatient Good progress

inappropriate use of corticosteroids, immunosuppressants and biological treatments in order to prevent the progression of COVID-19 and multiply the risks of opportunistic and pulmonary infections, especially in polytherapy [12,13].

For therapeutic management, recommendations are changing and evolving rapidly in order to provide up-to-date information on best practices for optimal management of infected IBD patients with the aim of treating active disease and maintaining remission [14,15].

Regarding corticosteroids that are often used as first-line treatment to induce remission and manage flares, current evidence suggests that they may pose a significant risk for IBD patients with COVID-19 and that their use is associated with increased mortality and morbidity (admission to intensive care unit, ARDS, shock) mainly for patients taking high doses [16].

According to very recent studies, it is proposed to stop/reduce corticosteroids as much as possible and not to continue prednisone at doses higher than 20 mg/day in SARS-CoV-2 positive IBD patients, whether symptomatic or not [17].

For budesonide and beclomethasone data are currently not available.

Preliminary data from the SECURE-IBD registry have identified 5-Aminosalicylic Acids (5-ASA) as a risk for severe COVID-19 infection [18], in the absence of more detailed information, it is proposed to discontinue them in patients with confirmed infection [19].

In theory, immunosuppression may reduce viral clearance, but may also reduce the cytokine storm involved in Acute Respiratory Distress Syndrome [ARDS]. However, current evidence to date does not suggest that IBD patients on immunomodulatory therapy fare worse than the general population, either in terms of risk of acquiring the virus or disease severity.

On the other hand, the risk of opportunistic infection seems to be increased in these patients because these drugs block the intracellular signals necessary for the host to fight pathogens, and their use has been associated with lymphopenia (which is a sign of a poor outcome of COVID-19). Therefore, it is thought that patients with inflammatory bowel disease may be at increased risk of SARS-CoV-2 infection with a risk of severe clinical outcome under immunomodulatory treatment.

Therefore, we believe that immunomodulators, anti-tumor necrosis factors, anti-TNF, anti-interleukins, anti-integrins and JAK inhibitors should be maintained during this pandemic and should be stopped in case of suspected

or confirmed infection, they can be resumed after complete resolution of COVID-19 symptoms or, ideally, after negative PCR tests.

Conclusion

The current outbreak of SARS-CoV-2 infection poses new challenges in the diagnosis and management of IBD patients. We propose to test all patients with diarrhea, to maintain treatment in uninfected subjects and to interrupt them in case of COVID infection.

Acknowledgment

None.

Conflict of Interest

The authors declare no competing interests.

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How to cite this article: Gharbi, Khalid, Khadija Krati and Lamiaa Essaadouni. "Infection (Covid-19) in Patients with Chronic Inflammatory Bowel Disease: A Case Series." J Inflam Bowel Dis Disorde 9 (2024): 225.