

Letter to the Editor

SARS-CoV-2 Infection in the Follow-Up of a Population With Inflammatory Bowel Disease

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To the Editors:

We read the manuscript titled "COVID-19 and Outcomes in Patients with Inflammatory Bowel Disease: Systematic Review and Meta-Analysis" by Tripathi et al¹ with great interest. It included 23 studies and 18 case reports incorporating 51 661 patients with inflammatory bowel disease (IBD), 1467 of whom had confirmed coronavirus disease 2019 (COVID-19). We agree with the most important messages that the prevalence of COVID-19 in IBD patients is low, anti-tumor necrosis factor therapy is not associated with an increased risk of adverse outcomes, and patients should continue their maintenance medications.

However, clinical reports with longer follow-up are lacking. A recent observational study with 1 year of follow-up suggested that neither IBD nor immunosuppressant use is associated with an increased risk of severe COVID-19.2 There are scarce data about COVID-19 and outcomes in Latin American IBD patients. Therefore, we decided to describe our experience of Chilean IBD patients during the first 20 months of the COVID-19 pandemic. A cross-sectional study was performed between January 2020 and August 31, 2021. Of 393 IBD patients being followed up, 22 (6%) were diagnosed with COVID-19 and 6 (27%) had comorbidities (4 had hypertension and 2 had diabetes). Demographic and clinical characteristics are presented in Table 1. Thirteen patients had COVID-19 before the vaccination period, 8 patients had COVID-19 after receiving 1 vaccine dose, and 1 patient had COVID-19 after receiving 2 vaccine doses. Four patients required hospitalization and no patients died.

Our findings are in accordance with the study by Tripathi et al and other publications.³ Data acquired during 20 months of follow-up showed that neither IBD nor immunosuppressant use was associated with an increased risk of COVID-19. Young patient age, a low frequency of comorbidities, and a high percentage of patients in clinical remission at the time of SARS-CoV-2 (severe acute respiratory syndrome coronavirus 2) infection may be associated with a lower risk of COVID-19, consistent with a previous study.²

Table 1: Demographic and clinical characteristics of Chilean patients with inflammatory bowel disease

	No SARS- CoV-2 Infection (n = 371)	SARS-CoV-2 Infection (n = 22)	P
Diagnosis			
Ulcerative colitis	243 (65)	14 (64)	NS
Crohn disease	117 (32)	8 (36)	
No classifiable IBD	11 (3)	0	
Age, y	41 (14-85)	36 (22-52)	NS
Sex			NS
Female	238 (64)	15 (68)	
Male	133 (36)	7 (32)	
Treatment			
5-ASA	229 (62)	13 (59)	NS
Immunosuppressant	105 (28)	7 (32)	NS
Biologic therapies	112 (30)	8 (36)	.042
Anti-TNF (ADA/IFX/ Goli)	63/17/27	2/2/1	_
p40 inhibitors (IL-12/23)	5	2	_
Anti-integrins	0	1	_
Small molecules			_
Tofacitinib	1	0	_
Prednisone	1 (<1)	0 (0)	_
Budesonide	16 (4)	1 (4)	_
Nontreatment	27 (7)	0 (0)	_
COVID-19 symptoms			
General symptoms	_	10 (45)	
Respiratory symptoms	_	9 (41)	
Digestive symptoms	_	2 (9)	
Asymptomatic	_	1 (5)	
Inflammatory activity after SARS-CoV-2 infection	_	22 (100)	

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Table 1. Continued

	No SARS- CoV-2 Infection (n = 371)	SARS-CoV-2 Infection (n = 22)	P
Remission	_	0 (0)	
Flare	_	_	

Values are N (%), median (range), or n.

Abbreviations: 3-ASA, 5-aminosalicylic acid; ADA, adalimumab; COVID-19, coronavirus disease 2019; Goli, golimumab; IBD, inflammatory bowel disease; IFX, infliximab; IL, interleukin; NS, not significant; SARS-CoV-2, severe acute respiratory syndrome coronavirus 2; TNF, tumor necrosis factor.

Attenuated immunogenicity to a single dose of a COVID-19 vaccine has been observed in IBD patients treated with infliximab.⁴ In our study, 50% of patients diagnosed with COVID-19 after receiving the first vaccine dose were on biological therapy. Another vaccine dose or booster vaccination is necessary in these patients. In the meantime, patients must maintain selfcare measures.⁵

In conclusion, our data suggest that patients with IBD are not at increased risk of COVID-19. Adherence to the immunization program is required by all IBD patients.

Author Contributions

P.N., R.Q., and L.F. contributed equally to drafting and preparation of this manuscript. All authors reviewed and approved the final manuscript.

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Conflicts of Interest

None.

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