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Editorial Board Member of World Journal of Gastroenterology, Dirk Jacob van Leeuwen, MD, PhD, FAASLD, Adjunct Professor, Section of Gastroenterology and Hepatology, Geisel School of Medicine at Dartmouth College, Hanover, NH 03756, United States. dirk.j.vanleeuwen@gmail.com

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LETTER TO THE EDITOR

Thiopurines are an independent risk factor for active tuberculosis in inflammatory bowel disease patients

Flora Maria Lorenzo Fortes, Raquel Rocha, Genoile Oliveira Santana

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Flora Maria Lorenzo Fortes, Genoile Oliveira Santana, Ciências da Vida, Universidade do Estado da Bahia, Salvador 41150000, Brazil, and Programa de Pós-Graduação em Medicina e Saúde, Universidade Federal da Bahia, Salvador 40110060, Brazil

Raquel Rocha, Ciência da Nutrição, Universidade Federal da Bahia, Salvador 40110160, Brazil

Corresponding author: Raquel Rocha, DSc, Adjunct Associate Professor, Ciência da Nutrição, Universidade Federal da Bahia, Av Araujo Pinho, Salvador 40110160, Brazil. raquelrocha2@yahoo.com.br

Abstract

The use of thiopurines is an independent risk factor for active tuberculosis in patients with inflammatory bowel disease.

Key Words: Tuberculosis; Inflammatory bowel disease; Thiopurines; Therapy; Risk

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Core Tip: Inflammatory bowel disease (IBD) patients recommended for anti-tumor necrosis factor (anti-TNF) therapy need to be tested for latent tuberculosis (TB) prior to treatment. Azathioprine monotherapy is also an independent risk factor for active TB in patients with IBD. However, the recommendations of the Brazilian Public Health Guideline for Tuberculosis Prevention do not include patients who are receiving immunosuppressive therapy in the risk group for screening for latent TB. We evaluated 301 patients with IBD, and the use of azathioprine treatment increased the risk by 6.87fold compared to patients without this treatment. The use of anti-TNF therapy had a 10.34-fold increased risk of TB, and the combination of both increased the risk by 17.81-fold.

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TO THE EDITOR

It is known that immunosuppression increases the risk of tuberculosis, especially in countries with a high frequency of active tuberculosis. We read with interest the article published by Fortes et al[1], who performed a retrospective cohort study among Inflammatory bowel disease (IBD) patients at a reference center in Brazil, which is a country with a moderate incidence of TB. A total of 301 IBD patients were evaluated; 61.8% had ulcerative colitis, and 38.2% had Crohn's disease. Twenty-seven (9.0%) patients received anti-tumor necrosis factor (anti-TNFα) as a monotherapy, 31 (10.3%) patients received anti-TNF α associated with azathioprine, 3 (1.0%) patients received anti-TNF α treatment associated with methotrexate, and 70 (23.3%) patients received only azathioprine. The use of azathioprine treatment increased the risk by 6.87-fold in comparison to patients without this treatment. The use of anti-TNF therapy showed a 10.34-fold increased risk of TB in this sample, and the association of both increased the risk to 17.81.

Advances in the treatment of IBD have been adopted worldwide. Some post marketing adverse events have been reported, including active tuberculosis (TB) during anti-TNF therapy. It has already been established that the incidence of active TB in this scenario is associated with the TB burden in the geographic region of the study. Brazil is one of the 20 countries in which TB presents a high incidence along with countries from Africa and Asia[2].

IBD patients with a recommendation for anti-TNF therapy need to test for latent TB before treatment. The TNF alpha blocking mechanism, which is critical in stabilizing granulomas during TB infection, would explain this increase in risk. An unanswered question is whether azathioprine in monotherapy is an independent risk factor for active TB in IBD patients[1,3]. A case report published by van Wijngaarden et al[4] already drew attention to the development of pleural tuberculosis in a patient with Crohn's disease while receiving azathioprine as the sole immunosuppressive treatment.

Considering that transplant recipients need substantial immunosuppression and azathioprine is one of the drugs used, studies among transplant recipients receiving immunosuppressive therapy helped guide physicians in the care of IBD patients. A Spanish group evaluated the risk factors for active TB after lung transplantation and concluded that the use of azathioprine was identified as an independent risk factor[5].

The recommendations of the Brazilian Public Health Guideline for Tuberculosis Prevention, reviewed in 2020, did not include patients receiving immunosuppressive therapy in the risk group for screening of latent TB[6]. However, consensus from endemic countries suggests investigation and treatment of latent TB before starting immunosuppressive therapy[7-9].

These findings suggest that in areas with a high burden of TB, the use of thiopurines is an independent risk factor for active TB in IBD patients. This evidence needs to be considered when using this therapy for these patients, especially those from countries with a high TB burden. We suggest giving attention to and treating patients with latent tuberculosis and guiding prevention with possible contacts with active tuberculosis. New studies reporting the risk of active TB among IBD patients receiving immunosuppressive therapy from countries with different incidence rates of TB are needed.

FOOTNOTES

Author contributions: Fortes FML designed the study and performed the data analysis; Rocha R reviewed the manuscript and provided technical and material support; and Santana GO contributed to the study design, manuscript revision, supervision of the study, had full access to all of the data in the study and was responsible for the integrity of the data.

Conflict-of-interest statement: Genoile O Santana is on the Advisory Board for Janssen; has received speaking fees from Abbvie, Ferring, Janssen, Takeda, Pfizer and UCB Pharma; and has received research grants from Janssen, Lilly, Pfizer, Roche and Takeda. The other authors declare that they have no conflicts of interest.

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Country/Territory of origin: Brazil

ORCID number: Flora Maria Lorenzo Fortes 0000-0002-9897-5337; Raquel Rocha 0000-0002-2687-2080; Genoile Oliveira Santana 0000-0001-5936-9791.

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