

times with a history of previous bowel resection for CD.<sup>4,5,10,20</sup> IFP pathogenesis remains unknown and several authors shared the etiology of a florid granulation tissue proliferation in response to local trauma.<sup>1,3,10</sup> It has been proposed that either CD itself or previous abdominal surgery for the disease may be the local stimulus for the formation of this apparently reactive lesion, this local injury hypothesis being enhanced by the conjunction of an underlying inflammatory bowel disease. Histologically, other mass lesions could arising in the case of CD, such as lymphoid follicular hyperplasia, diffuse edema with lymphangiectasia, and also regenerative pseudopolyps which display an inflammatory mucosae with basophilic crypts and glands.<sup>4</sup> Of course, a malignant lesion should be excluded.

Concerning the management of intussusception, surgical resection of the bowel segment is required in most cases. Although IFP are benign lesions, 2 recurrence cases were reported in the literature after surgery.<sup>21,22</sup>

**L. Deschamps, MD\***

**F. Bretagnol, MD†**

**A. Couvelard, MD\***

**O. Corcos, MD‡**

**P. Bedossa, MD\***

**Y. Panis, MD†**

\*Department of Pathology

†Department of Colorectal Surgery

‡Department of Gastroenterology and Nutrition Support

Pôle des Maladies de l'Appareil Digestif (PMAD)

Beaujon Hospital  
Clichy, France

## REFERENCES

- Johnstone JM, Moroson BC. Inflammatory fibroid polyp of the gastrointestinal tract. *Histopathology*. 1978;2:349–361.
- Dilawari RA, Patterson WB. Inflammatory fibrous polyp (pseudotumor) of ileum, a rare cause of intestinal obstruction. *Am Surg*. 1976;4:920–922.
- Shimer GR, Helwig EB. Inflammatory fibroid polyps of the intestine. *Am J Clin Pathol*. 1983;81:708–714.
- Williams GR, Jaffe S, Scott CA. Inflammatory fibroid polyp of the terminal ileum presenting in a patient with active Crohn's disease. *Histopathology*. 1992;20:545–547.
- Parasi A, Triantafyllidis JK, Barbatzas C, et al. Coexistence of Crohn's disease and inflammatory fibroid polyp of the small bowel. *Ann Ital Chir*. 2005;76:395–399.
- Vanek J. Gastric submucosal granuloma with eosinophilic infiltration. *Am J Pathol*. 1949;25:397–411.
- Helwig EB, Ranier A. Inflammatory fibroid polyps of the stomach. *Surg Gynecol Obstet*. 1953;96:355–367.
- Korman U, Kuruoglu S, Haider S. Rare complication of intestinal Crohn's disease: giant fibroid polyp. *Eur Radiol*. 2000;10:435–437.
- Ozolek JA, Sasatomi E, Swalsky PA, et al. Inflammatory fibroid polyps of the gastrointestinal tract: clinical, pathologic, and molecular characteristics. *Appl Immunohistochem Mol Morphol*. 2004;12:59–66.
- Bays D, Anagnostopoulos GK, Katsaounos E, et al. Inflammatory fibroid polyp of the small intestine causing intussusception: a report of two cases. *Dig Dis Sci*. 2004;49:1677–1680.
- Agha F. Intussusceptions in adults. *AJR Am J Roentgenol*. 1986;146:527–531.
- Atalay F, Balci S, Kirimlioglu, et al. Intussusception due to inflammatory fibroid polyp of the ileum. A report of two cases from Turkey. *Hiroshima J Med Sci*. 1995;44:141–144.
- Zager JS, Shaw JP, Kaufman JP, et al. Three cases of small bowel intussusception in relation to a rare lesion: inflammatory fibrous polyps. *Dig Surg*. 2001;18:142–146.
- Benjamin SP, Kawk WA, Turnbull RB. Fibrous inflammatory polyps of the ileum and caecum: review of five cases with emphasis on differentiation from mesenchymal neoplasm. *Cancer*. 1977;39:1300–1305.
- Balci NC, Radjazi S, Polat H. Adult intussusception secondary to inflammatory fibroid polyp: demonstration of MRI. *Eur Radiol*. 2000;10:1708–1710.
- Isik Gonul I, Erdem O, Ataoglu O. Inflammatory fibroid polyp of the ileum causing intussusception: a case report. *Turk J Gastroenterol*. 2004;15:59–62.
- Suen KC, Burton JD. The spectrum of eosinophilic infiltration of the gastrointestinal tract and its relationship to other disorders of angiitis and granulomatosis. *Hum Pathol*. 1979;10:31–43.
- Goldman RL, Friedman NB. Neurogenic nature of so-called inflammatory fibroid polyps of the stomach. *Cancer*. 1967;20:134–143.
- Bayle S, Rossi P, Bagneres D, et al. Ileum inflammatory fibroid polyp revealed by intussusception. About one familial case. *Rev Med Intern*. 2005;26:233–237.
- Navas-Palacio JJ, Colina-Ruizdelgado F, Sanchez-Larrea MD, et al. Inflammatory fibroid polyps of the gastrointestinal tract. An immunohistochemical and electron microscopic study. *Cancer*. 1983;5:1682–1690.
- Anthony PP, Morris DS, Vowles KDJ. Multiple and recurrent inflammatory fibroid polyps in three generations of a Devon family: a new syndrome. *Gut*. 1984;25:854–862.
- McGreevy P, Doberneck RC, McLeay JM, et al. Recurrent eosinophilic infiltrate (granuloma) of the ileum causing intussusception in a two-year-old children. *Surgery*. 1967;61:280–284.

## Mesalazine-Induced Jaundice, Eosinophilia, and Thrombocytopenia

### To the Editor:

A 45-year-old man was admitted as an acute surgical emergency with a 6-day history of right upper quadrant pain, jaundice, and pale stools. There were no identifiable risk factors for liver disease. There was an 8-year history of ulcerative colitis (currently quiescent) for which he was taking mesalazine 1600 mg (total duration of therapy = 100 months). The patient was jaundiced with no peripheral stigmata of chronic liver disease. The blood results were as follows: hemoglobin = 16.8 g/L; white cell count =  $60.4 \times 10^9/L$  (normal range = 4–11) with a eosinophilia of  $49.5 \times 10^9/L$  (normal range = 0.00–1.0); and platelet count =  $89 \times 10^9/L$  (normal range = 150–400). Bilirubin = 137 mmol/L (normal <20 mmol/L); alkaline phosphatase = 170 IU/L (normal range = 5–30); alanine transferase = 553 IU/L (normal range = 30–130). The serum albumin and clotting profile were normal.

The patient had been under annual review for the previous 7 years and the blood tests were consistently normal during this follow-up period. Abdominal ultrasound revealed a normal liver parenchyma, a thickened gall bladder, but no ductal dilation. Colonic biopsies revealed a moderately severe ulcerative colitis. Mesalazine was stopped on admission. However, the blood tests worsened and therefore a liver biopsy was performed. This revealed prominent eosinophilia involving the sinusoids, parenchyma, and, in particular, the central veins and portal tracts consistent with drug-induced hepatitis. The blood

Copyright © 2008 Crohn's & Colitis Foundation of America, Inc.

DOI 10.1002/ibd.20445

Published online 1 May 2008 in Wiley InterScience (www.interscience.wiley.com).

tests improved over the next week and eventually returned to normal. In the 3 years since presentation his liver function tests and full blood count have remained normal and his colitis has been well controlled on azathioprine.

Despite the long duration of therapy with mesalazine prior to this reaction, we conclude that the jaundice, eosinophilia, and thrombocytopenia were adverse reactions of mesalazine therapy. We have not rechallenged our patient with mesalazine because of the fear of inducing a severe reaction.

Hypersensitivity reactions to sulfasalazine including the development of hepatotoxicity and eosinophilia are well recognized.<sup>1</sup> These effects were initially blamed on the sulfa pyridine moiety and mesalazine was thought to be free of these effects. However, there have been separate reports of eosinophilia,<sup>2</sup> hepa-

totoxicity,<sup>3,4</sup> and thrombocytopenia<sup>5</sup> consequent to mesalazine therapy and reports of patients hypersensitive to sulfasalazine developing a severe reaction when given mesalazine.<sup>6</sup> In published case reports the duration of therapy prior to an adverse reaction has varied from a few days to 2 years. We believe this is the first reported case where a patient has experienced hepatotoxicity, thrombocytopenia, and eosinophilia concurrently as a consequence of 5-ASA therapy with onset many years after starting the drug. 5-ASA-induced toxicity should therefore be considered in any patient taking 5-ASA irrespective of the duration of therapy.

**Manu Nayar, MRCP**  
**William Cunliffe, FRCPS**  
**Paul Cross, FRCPath**  
**Kofi Oppong, FRCP**  
 Freeman Hospital

Gastroenterology  
 High Heaton  
 Newcastle upon Tyne, UK

## REFERENCES

1. Ransford RA, Langman MJ. Sulphasalazine and mesalazine: serious adverse reactions re-evaluated on the basis of suspected adverse reaction reports to the Committee on Safety of Medicines. *Gut*. 2002;51:536.
2. Bitton A, Peppercon MA, Hanrahan JP, et al. Mesalamine-induced lung toxicity. *Am J Gastroenterol*. 1996;91:1039–1040. Review.
3. Marteau P, Nelet F, Le Lu M, et al. Adverse events in patients treated with 5-aminosalicylic acid: 1993-1994 pharmacovigilance report for Pentasa in France. *Aliment Pharmacol Ther*. 1996;10:949–956. Review.
4. Deltenre P, Berson A, Marcellin P, et al. Mesalazine (5-aminosalicylic acid) induced chronic hepatitis. *Gut*. 1999;44:886–888.
5. Farrell RJ, Peppercon MA, Fine SN, et al. Mesalamine-associated thrombocytopenia. *Am J Gastroenterol*. 1999;94:2304–2306.
6. Hautekeete ML, Bourgeois N, Potvin P, et al. Hypersensitivity with hepatotoxicity to mesalazine after hypersensitivity to sulfasalazine. *Gastroenterology*. 1992;103:1925–1927.