Various systemic and gastrointestinal diseases cause mucosal alterations of the duodenum. These include celiac disease, Crohn’s disease, congenital enteropathies (e.g., microvillus inclusion disease, tufting enteropathy), and Whipple’s disease [1–3]. Thus careful endoscopic inspection and targeted biopsies are strongly recommended to achieve a correct diagnosis and avoid false reassurance.

Here we describe the case of a 45-year-old man who presented with diarrhea, weight loss, and mild abdominal pain. In addition, he complained of strong joint pains. Physical examination was unremarkable. Laboratory investigations showed hemoglobin of 8.2 g/dL (reference 13–17), iron of 25 μg/dL (40–160), and C-reactive protein of 84 mg/L (< 5). Upper-gastrointestinal endoscopy revealed pale yellow shaggy mucosa with intermittent, superficial, erythematous eroded patches of the duodenum (Fig. 1a).

Optical chromoendoscopy with narrow-band imaging (NBI) (Olympus, Tokyo, Japan) showed the intestinal villi to appear edematous and slightly flattened. Although no changes of the microvasculature were observed, the villous lymphatics were clearly enhanced (i.e. villous lymphangiectasia) (Fig. 1b). Histopathological examination of the intestinal biopsies showed multiple periodic acid–Schiff (PAS)-positive macrophages within the lamina propria, characteristic of Whipple’s disease (Fig. 1c). Our patient was put on oral antimicrobial therapy with co-trimoxazole for 2 years and remained in remission at 6-month follow-up.

Whipple’s disease is a rare multisystemic bacterial infection caused by Tropheryma whippelii that can involve the gastrointestinal, nervous, and cardiac system and the skin [1–3]. Previously, we have shown that high-magnification endoscopy can clearly help to visualize the mucosal characteristics typical for Whipple’s disease, including edematous and engorged duodenal villi filled with a white material with the appearance of lymph [2]. Although various studies have described the potential of advanced endoscopic imaging in the small bowel, there are currently no data available on the usefulness of chromoendoscopy (either vital or virtual) for the evaluation and diagnosis of Whipple’s disease [3,4].

Our case is interesting for several reasons, because it shows for the first time the findings of Whipple’s disease using NBI. NBI can aid in the characterization of the mucosal alterations present in Whipple’s disease, including villi architecture disarrangement and altered lymphatic microvasculature. In addition, our case highlights clinical and histological characteristics of Whipple’s disease. Physicians should be aware of the differential diagnosis and should also consider duodenal biopsies in patients submitted because of extraintestinal symptoms such as joint pains and pericarditis, and increased inflammatory reactions such as elevated sedimentation rate and thrombocytosis [2–4].

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References