A 76-year-old man with previous history of kidney transplantation in 2003 (due to chronic kidney injury of unknown etiology), arterial hypertension and a thoracoabdominal aortic aneurysm was admitted to our center for a thoracic endovascular repair. His usual medication was tacrolimus, mycophenolate mofetil, prednisolone, captopril and carvedilol. After surgery the patient was referred for an abdominal computed tomography (CT) due to epigastric pain and vomiting with associated hemoglobin drop to 8 g/dl (from a previous value of 14 g/dl). CT scan revealed a massive retroperitoneal hematoma (12 × 6.5 cm). Blood transfusions were undertaken and due to iron deficiency anemia, oral ferrous sulfate was initiated.

One week later, the patient was referred for an upper endoscopy due to dysphagia.

Upper endoscopy did not show esophageal lesions; however multiple erosions in the gastric body were detected (Fig. 1) with biopsies revealing extracellular crystalline iron deposition the superficial lamina propria beneath eroded epithelium and in the lumen of the gastric glands, with some focal intracellular granular iron deposition in glandular cells, confirmed by Perls’ iron (Prussian blue) stain (Fig. 2). In the upper endoscopy was also evident a 12 mm ulcer with clean base in the anterior wall of the second part of duodenum (Fig. 3). Biopsies of duodenal mucosa revealed an area of ulceration with extracellular crystalline and fibrillar iron deposition (Perls’ positive) over granulation tissue associated with foreign body reaction (Fig. 4).

Cytomegalovirus (CMV) infection was ruled out after negative results in the immunohistochemical and molecular biology analysis. Also Helicobacter pylori infection was not found with Giemsa stain study.

Oral ferrous sulfate is commonly used to treat iron deficiency anemia. Iron pill-induced gastroduodenopathy is a possible, but rarely described complication of oral iron supplementation. However it is likely that its exact prevalence is largely underestimated. Abraham et al. reported a prevalence of focal erosive mucosa injury from iron therapy in 0.7% of gastrointestinal biopsies.1 Kaye et al. also prospectively evaluated the effects of

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iron supplementation on patients with iron deficiency anemia and encountered visible erosions in 6 of the 16 patients with confirmed iron deposition on biopsy, a considerable higher proportion when compared to patients without iron supplementation (14/141).2

Histologically, iron deposition in the upper gastrointestinal tract can occur in three distinctive patterns, as described by Marginean et al.3-5 Type A (“nonspecific”) pattern is characterized by intracellular iron deposition in macrophages, stromal and epithelium. It is frequently associated with mucosal inflammation and may represent iron deposition from prior mucosal hemorrhage. Type B (“iron-pill induced mucosal injury”) pattern is characterized mostly by extracellular deposition and focally, by intracellular deposition in blood vessels, macrophages, and epithelium. This type is typically associated with oral iron-containing medications. In this type iron deposition may be associated with fibrosis, inflammation, and foreign body reaction. In the type C (“glandular siderosis”) pattern there is diffuse iron deposition predominantly in deeper glands of the mucosa. It is often associated with systemic iron overload or hemochromatosis.

In our case, the iron deposition in both gastric and duodenal mucosa fit in the pattern B, suggesting that the mucosal injury was probably triggered by the oral ferrous sulfate intake.

However, since the patient had multiple comorbidities it is impossible to know if ferrous sulfate was the main responsible or exacerbated previously induced gastroduodenal ulceration.

Reports of iron-induced gastroduodenal mucosal injury with endoscopic and histologic evidence are scarce. Histology is essential to render this diagnosis. In the presence of ulcers with atypical location or with suspicious endoscopic findings, biopsies should always be obtained. If this condition is suspected, iron pills should be suspended and iron should be administered intravenously.6
Conflicts of interest

The authors declare no conflicts of interest.

References