Upper Gastrointestinal Bleeding: A Complication of “Inlet Patch Mucosa” in the Upper Esophagus

Heterotopic gastric mucosa has been reported along the alimentary tract, and its prevalence in the esophagus varies from 0.1% to 10% (1,2). Rare complications have been described, including dysphagia, stricture, esophagotracheal fistula formation, and adenocarcinoma development (1–3).

A 20-year-old male was admitted to our hospital due to two episodes of hematemesis with hypovolemic signs (98 bpm and blood pressure 100/60 mmHg) and hematoctrit of 38%. A complete emergency endoscopic examination showed two ulcers (1.5 x 1.0 cm each) located at a distance of 22 cm from the incisors (Figure 1). No signs of gastroesophageal reflux and no other possible sources of bleeding were noted in a complete esophagogastroduodenoscopy. The biopsy specimens taken from the esophageal ulcers showed squamous and foveolar, gastric-type mucosa, lined by surface columnar mucus. A mild lymphocytic infiltrate was present in the lamina propria (Figure 2). Omeprazole (40 mg daily) was administered, with no recurrence of the hemorrhage in the one-year follow-up. At the control endoscopic examination, a circumferential red patch was detected at 22 cm from the incisors, but no ulceration was seen.

Heterotopic gastric mucosa is most commonly found in the cervical portion of the esophagus, due to the widespread use of gastrointestinal endoscopy. These lesions appear as a reddish spot or patch that stands out against the pale squamous epithelium of the esophagus. Biopsies of these patches show fundic-type mucosa with chief and parietal cells (1,2). The clinical relevance of these lesions is controversial, and a relationship between the endoscopic finding and symptoms is rarely demonstrable (3–4). Murray et al. describe a case with repeated rectal bleeding because of gastric mucosa present in the rectum, controlled with H2-receptor antagonists (5), but no cases of upper gastrointestinal bleeding due to heterotopic mucosa in the cervical esophagus have been described.

R. Bataller1, J.M. Bordas1, J. Ordi1, J. Llach1, J.I. Elizalde1, F. Mondelo1
1 Endoscopy Section, Gastrointestinal Service, and
2 Department of Pathology, Clinic i Provincial Hospital, University of Barcelona, Barcelona, Spain

References

Corresponding Author
J. M. Bordas, M.D.
Section of Digestive Endoscopy
Hospital Clinic i Provincial
University of Barcelona
Villarroel 170
08036 Barcelona
Spain