A 39-year-old woman presented for colon cancer screening colonoscopy. She had no abdominal complaints. The entire colonoscopy was normal, except that the standard retroflexed view in the rectum demonstrated a 2-cm diverticulum without raised margins 5 cm from the anal verge (Fig. 1a, b). Endoscopically the mucosa appeared fairly edematous, but without malignant characteristics. The histological examination of the biopsies obtained revealed rectal mucosa abutting ectopic gastric mucosa without Helicobacter pylori (Fig. 2).

Heterotopic gastric mucosa can be observed anywhere along the human gastrointestinal tract. It is rarely seen in the rectum: only about 30 cases have been reported since the first description in 1939 [1]. In the clinical setting, gastric heterotopia typically presents as either an ulcerative lesion or abnormal mucosa in the rectum [2]. The origin of gastric mucosa in the rectum is believed to be failure of developmental descent of the embryonic hindgut. The occurrence of gastric heterotopia within a hindgut developmental anomaly such as rectal duplication is exceptional [3]. In clinical practice, a so-called rectal Meckel’s diverticulum may present with acute profuse rectal bleeding, especially in children and young adults [4]. In these cases surgical resection is advised. In patients with peptic ulceration, treatment with H2-blockers or proton pump inhibitors should be prescribed first. A rectal Meckel’s diverticulum may also be asymptomatic, as it was in our patient. We choose not to perform preventive resection or regular endoscopic screening, so long as the course is uneventful.

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