A 30-year-old woman with history of common variable immunodeficiency disease (CVID), Von Willebrand disease, and factor VII hemophilia presented to the emergency department with complaints of diarrhea and weight loss for 2 weeks. On admission her vital signs were stable. She reported 6–8 watery bowel movements per day, which were not improved with Imodium, but had no hematochezia or melanotic stools. The findings from the physical examination were noncontributory. She had work-up for diarrhea, which was negative for infectious etiology. A positive fecal fat test suggested malabsorption.

Of note, she had undergone a colonoscopy in 2010, which had been indicated for chronic diarrhea, and biopsy results had revealed nodularity in the ascending colon and rectal mucosa consistent with lymphoid hyperplasia. She also underwent esophagogastroduodenoscopy during her present admission in 2014, which revealed no irregularities, and a biopsy, which was negative for celiac sprue. Subsequently, she underwent capsule endoscopy (Fig. 1) and small-bowel enteroscopy (Fig. 2), which revealed grossly nodular lymphoid hyperplasia throughout the distal small bowel. Small-bowel enteroscopy revealed areas with multiple 4–7-mm sessile polyps and no bleeding in the proximal and distal jejunum (Fig. 2). The patient responded well to budesonide and metronidazole and underwent follow-up with a gastrointestinal (GI) specialist.

CVID is a primary immunodeficiency disorder that is characterized by impaired B-cell differentiation and defective immunoglobulin production [1]. GI manifestations of CVID include diarrhea, malabsorption, weight loss, acute infections, and inflammatory bowel disease. Diffuse
nodular lymphoid hyperplasia is a rare lymphoproliferative disorder of the GI tract with uncertain etiology [2,3]. It is believed that exposure to infectious agents leads to the repetitive stimulation of lymphoid follicles and eventual hyperplasia [4,5].

**Competing interests:** None

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