A 44-year-old man with no relevant past medical history was referred for colonoscopy because of recurrent hematochezia and iron deficiency anemia. He reported no abdominal pain or weight loss. He had been on oral therapy with ferrous sulfate. A hypertrophic lower left limb was seen on physical examination. Colonoscopy (Video 1) revealed ectasia and congestion of the submucosal and mucosal venous vessels and hemangiomas affecting the rectum and the left colon in a 44-year-old man with Klippel–Trenaunay syndrome.

Endoscopic findings associated with Klippel–Trenaunay syndrome are highlighted.

Klippel–Trenaunay syndrome is a congenital malformation syndrome characterized by the presence of cutaneous capillary malformations, asymmetrical disturbed growth of soft tissues and/or bone, and venous and/or lymphatic malformations [1]. It is rare, with an unknown etiology and an incidence of approximately 1:100,000 live births [2]. Gastrointestinal involvement occurs in about 1%–13% of affected patients, most of whom are asymptomatic [3]. Gastrointestinal hemorrhage can occur, ranging from occult to life-threatening and severe in nature [2, 3]. In this patient, endoscopic and angiographic treatments were not feasible and the severity of symptoms forced an early surgical approach. The endoscopic findings associated with Klippel–Trenaunay syndrome are highlighted.
References


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