A substantial embossment of the lesion and a marked vascular net in the proximity is observed with Fuji Intelligent Chromo Endoscopy (FICE), (Figure 2). These findings reinforce the belief of the vascular origin of the process. EUS (Figure 3 and 4) showed an hypoechoic lesion originated in the deeper mucosal layer that exhibited doppler signs inside and spared the submucosal and muscular layers. These findings are consistent with those described for esophageal haemangioma.2,8

The esophageal haemangioma is a rare entity, representing less than 3% of the benign tumors of the esophagus.1 When they involve the gastrointestinal tract, the most frequent sites are the small bowel, the colon and the stomach. The inferior portion is the site of the esophagus most frequently affected.3 When patients present with symptoms, dysphagia, bleeding, epigastric pain, thoracic pain and melena are the most common complaints.2 Hemangiomas are classified as cavernous, capillary and mixed types.4 The cavernous type is composed of large dilated vascular spaces filled with blood and the capillary type is composed of conglomerates of small, thin-walled vessels. The cavernous type, when large, may bleed seriously, with platelet consumption within the lesion being an aggravating factor. The bleeding is often episodic and initiates in childhood.5 The endoscopic feature is very suggestive and the biopsy may be avoided due the risk of hemorrhage.5 Atypical cases may be confounded with esophageal carcinoma.1
The treatment of choice for esophageal hemangioma has classically been described as surgical, but with the recent advances in therapeutic endoscopy, many reports have appeared in medical literature for the endoscopic treatment of esophageal hemangiomas, such as endoscopic band ligation, snare polypectomy, mucosectomy, sclerotherapy and laser fulguration.2,6-8

In the present report we describe a case of a young symptomless man with an incidental discovered esophageal lesion. We believe that we added important data to the diagnosis of esophageal hemangioma by means of little invasive procedures such as EUS and FICE. The endoscopic ultrasonography findings were consistent with those described in other publications for this pathology. We did not find previous reports in the literature using FICE for the diagnosis of gastrointestinal tract hemangiomas. It is our opinion that FICE reinforced the diagnosis of hemangioma in our case by enhancing the vascular structure and nature of the lesion. However, these results should be supported by more robust data.

On the other hand, we do not have notice about the natural history of esophageal hemangiomas in symptomless young patients. To our knowledge there are no papers comparing the morbidity and mortality between the natural history and the surgery. Therefore, in the absence of a more reliable report and after a meticulous explanation, the patient preferred an expectant behavior with follow-up of the lesion.

Referencias