Solitary Esophageal Varix Simulating a Neoplasm

F.P. AGHA, M.D.
Department of Radiology
University of Michigan, Medical Center, Ann Arbor, Michigan 48109

A solitary esophageal varix simulating an esophageal tumor is reported. To my knowledge this rare entity has only been described once previously.

CASE REPORT

A 57-year-old woman was referred to the University of Michigan Hospital in July 1984 for evaluation of iron deficiency anemia. She was transfused with five units of blood and then underwent complete evaluation of her gastrointestinal tract. During double contrast examination of her upper gastrointestinal tract a 2.0 x 1.3 cm lobulated well demarcated polypoid lesion was found in the mid esophagus. This lesion was unaffected by esophageal distension (Figure 1). It was felt to be a benign lesion such as a fibrovascular polyp or a lipoma, however, malignancy could not be completely excluded. The remainder of the upper gastrointestinal tract and small bowel follow through was normal except for a small hiatal hernia. An air contrast barium enema was unremarkable. Endoscopic examination of the esophagus revealed a sausage shaped lobulated 2.1 x 1 cm firm submucosal mass in the mid esophagus with normal overlying mucosa. It had the bluish tinge of an ectatic venous structure. The diagnosis of a thrombosed esophageal varix was made. To completely exclude the possibility of a neoplasm a mucosal biopsy was performed which was normal at histopathologic examination. Colonoscopy revealed large internal hemorrhoids and no other abnormalities. A chest CT examination did not reveal any mediastinal abnormalities. The esophageal varix was not seen. There was no evidence of portal hypertension or liver disease.

It was felt that source of this patient’s chronic gastrointestinal blood loss was large internal hemorrhoids. She was discharged and subsequently underwent an elective hemorrhoidectomy. Her clinical course was satisfactory. Follow-up endoscopic examinations at 6 and 12 months have revealed no change in the appearance of the solitary thrombosed esophageal varix.

DISCUSSION

The typical esophageal varices are characterized by their multiplicity, orad orientation in the distal half of the esophagus and the clinical setting of portal hypertension. On the basis of etiology esophageal varices may be classified into congenital or acquired. In a detailed review of the subject, Jorup2 found that only two or three authentic cases of congenital esophageal varices have been reported in the world literature up to 1948. Harinck, et al3 in 1971 reported congenital esophageal varices in identical twins without portal hypertension.

The acquired varices are most commonly due to portal hypertension. The common intrahepatic causes of portal hypertension consist of alcoholic cirrhosis, cryptogenic cirrhosis, hepatic cirrhosis associated with hepatocellular carcinoma, congenital hepatic fibrosis, chronic active hepatitis, biliary cirrhosis, Wilson’s disease, hepatic vein thrombosis and obstruction, hemochromatosis and galactosemia. The extrahepatic causes of portal hypertension consist of portal vein atresia, stenosis, cavernous malformation and thrombosis, compression of portal vein by tumor, cyst or nodes, arterioporal fistula related to trauma and pancreatitis. Some rare causes include schistosomiasis, alpha I — antitrypsin deficiency and methotrexate induced hepatic fibrosis4. In most esophageal varices due to portal hypertension, the blood flow is “uphill” towards the superior vena cava and azygos vein. Esophageal varices are occasionally found in patients without portal hypertension.

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Address for reprints:
Farooq P. Agha, M.D.
Department of Radiology — Box 013
University Hospital
1405 East Ann Street
Ann Arbor, Michigan 48109

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hypertension. These are so called "downhill" varices found in the upper esophagus and are most commonly due to obstruction of the superior vena cava or azygos vein by neoplastic processes of the mediastinum, retrosternal goitre, thymoma and mediastinal fibrosis.

A solitary esophageal varix without underlying portal hypertension, superior vena cava or azygos venous obstruction is extremely rare. Its presentation as esophageal mass has only been reported once previously.

The typical esophageal varices are distinguished radiographically as multiple serpiginous filling defects which can be effaced by distension of the esophagus during double contrast esophagram. This feature may be seen in a solitary varix without any thrombosis, however, a thrombosed varix does not show obliteration by distension of the esophageal lumen. Of the two cases reported previously, one did and the other did not, show obliteration of the varix when the esophagus was distended. In the present case the polypoid lobulated mass did not change in size, shape or configuration during double contrast esophagram.

The incidence of solitary esophageal varix is not known. The congenital esophageal varices and idiopathic esophageal varices are also reported to be quite rare. Garrett and Gall in a series of 18,000 autopsies found only 3% incidence of idiopathic esophageal varices. None of the standard texts and major reviews pertaining to esophageal diseases and portal hypertension mention about solitary esophageal varix.
SOLITARY ESOPHAGEAL VARIX SIMULATING A NEOPLASM

The mechanism of development of a solitary varix is not known. It has been postulated that congenital weakness of the wall of venous channels might be responsible. In three cases of idiopathic solitary esophageal varix reported thus far (including the present case), the varix was located in the mid esophageal region, ranging in size from 1-2 cm with typical submucosal location. Two were thrombosed and one without thrombosis.

The most common differential diagnosis of solitary esophageal varix should include benign intramural tumors, esophageal cyst (retention cyst or foregut respiratory cyst) and localized hematoma. The intramural and submucosal tumors of the esophagus which constitute the intramural and submucosal lesions are usually asymptomatic until they are large enough to compromise the lumen to produce dysphagia. Although carcinoma of the esophagus, esophageal lymphoma and metastatic tumors to the esophagus may present as submucosal mass, the associated clinical setting and other radiographic finding are helpful in arriving at a correct diagnosis.

The present case is quite unique, because despite its unchanged radiographic appearance at double contrast esophagram, the endoscopy revealed characteristic submucosal location, bluish tinge and ectatic venous structure. The diagnosis of a non-thrombosed solitary varix can be suggested with confidence at esophagram. The thrombosed varix presents a diagnostic challenge and difficult to diagnose short of skillful endoscopy or surgery. Awareness of this entity and familiarity with its radiographic and endoscopic features are a prerequisite for arriving at a correct diagnosis.

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