Congenital Esophageal Stenosis: A Rare Case of Childhood Dysphagia

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Rec date: Jan 28, 2016; Acc date: Feb 4, 2016; Pub date: Mar 15, 2016

Abstract

Congenital esophageal stenosis (CES) is a rare cause of esophageal dysphagia. It is secondary to failure of complete separation of respiratory tract from primitive gut in early fetal life. Diagnosis may be delayed requiring high index of clinical suspicion and esophagogram studies. Final diagnosis of the subtype may require histopathological examination. Treatment may range from endoscopic dilatation in mild cases to esophageal resection in severe cases. In this article, we describe a rare case of congenital esophageal stenosis presenting in late childhood as dysphagia with characteristic barium esophagogram study. Very few cases have been reported in the medical literature in the past.

Keywords: Congenital; Stenosis; Dysphagia

Introduction

Congenital esophageal stenosis is a developmental anomaly occurring in approximately one out of 25,000-50,000 live births [1]. Fetal anoxia causing defective canalization of esophagus is a postulated mechanism [2]. Clinical presentation may range from dysphagia or recurrent vomiting/aspiration pneumonitis in early childhood to severe dysphagia, chest pain during deglutition or recurrent food impaction in early adulthood. Early diagnosis is the key to early management obviating the need for major esophageal surgery especially resection.

Case Report

An eleven-year old boy presented in our hospital outpatient department with a complaint of significant progressive dysphagia since last 2 years associated with regurgitation of food particles recently without history of significant weight loss or chest pain.

Figure 1: Radiograph of chest in PA projection shows right mediastinal soft tissue mass.

No history of dysphagia in early childhood or choking/respiratory symptoms could be elicited. There was no history of corrosive ingestion in the past. No history of any obvious abnormality is noted in antenatal fetal ultrasonographic examination. Laboratory tests were unremarkable. Chest radiograph revealed right mediastinal soft tissue mass with smooth lateral margin (Figure 1).

Barium swallow examination was then advised to look for the cause of dysphagia. Examination was performed under image intensifier and it revealed dilatation of proximal two-third esophagus (corresponding to right mediastinal soft tissue mass in chest radiograph) without shouldering or reflux esophagitis or hiatus hernia (Figure 2).

Evidence of irregular filling defect was noted at the site of stricture associated with eccentric jet-effect (Figure 3) and fairly-preserved longitudinal mucosal folds in the normal-appearing distal esophagus and morphologically normal gastro-esophageal junction (Figure 4).

Esophageal emptying was delayed with proximal hold-up of contrast in esophageal lumen for up to 60 minutes associated with slight oral regurgitation of barium.

Figure 2: Barium swallow views in frontal (a) and oblique (b) projection show partial, abrupt, smooth, luminal narrowing in the distal-third esophagus (white arrow) showing moderately dilated proximal esophagus corresponding to the right mediastinal soft tissue mass on plain radiograph without shouldering.
on barium esophagogram was due to food particles admixed with desquamated mucosa. No evidence of any obvious signs of malignancy was noted.

**Discussion**

Congenital esophageal stenosis may exist in one of the following three forms [2]:

- Thin esophageal membrane or web comprising of normal mucosa and submucosa,
- Fibromuscular stenosis caused by fibrotic changes in submucosa and muscularis propria, and
- Tracheobronchial remnants characterized by presence of cartilage, tracheal glands and respiratory epithelium.

CES secondary to tracheobronchial remnants is commonest involving preferentially lower-third esophagus while those secondary to web/membrane and fibromuscular etiologies are more common in the middle-third part of esophagus [3]. In general, higher stenoses present commonly with respiratory symptoms while lower ones present commonly with vomiting.

Esophageal web or membrane is 1-2 mm thick, shelf-like structure made of normal esophageal mucosa and submucosa causing partial or complete luminal obstruction showing central or eccentric orifice.

Though symptoms secondary to CES may start around the weaning period during infancy with dysphagia to solids yet in many cases minor degrees of stenosis is often overlooked, seeking medical attention only in young adulthood due to major food impaction or regurgitation [4]. Despite a long history of dysphagia, a high-clinical suspicion with characteristic esophagogram serves as clues for early diagnosis. Final diagnosis may require tissue examination in most cases as CES secondary to tracheobronchial remnants is the commonest etiology [5]. About one-third cases of CES may be associated with esophageal atresia [6,7].

Esophagogram in CES characteristically reveals short-segment, circumferential luminal narrowing in upper, middle or lower-third part of esophagus with smooth and tapering margins associated with proximal dilatation without obvious signs of shouldering [4,5]. However, normal esophagogram may be noted in initial cases while signs of reflux esophagitis may be noted in cases of lower-third involvement.

Multiple cases of CES in various forms have been reported in the literature especially tracheobronchial type [1-4,6-8] but very few cases secondary to esophageal web especially in late childhood had been reported in medical literature in the past [9-11].

Puntis et al. described a case of esophageal web causing dysphagia in young child involving the postcricoid region [9]. CES secondary to esophageal web may require luminal dilatation with endoscopic resection or laser division as luminal dilatation alone is associated with recurrence [10]. Altamimi also reported a case of childhood dysphagia caused by esophageal web in upper esophagus [11]. Thus our index case is not only rare but is unique in the fact that it involves lower-third esophagus.

In addition to CES, differential diagnosis of esophageal stricture includes the following:

- Reflux esophagitis usually seen with lower-level stenosis causing gastroesophageal reflux showing lack of distensibility with or
without transverse/converging, thickened folds secondary to inflammation,
• Achalasia which is rare in infants and children involving the esophagogastric junction showing a long, tapered, narrow segment resembling a "bird's beak" with disorganized peristalsis,
• Vascular ring due to double aortic arch seen as smooth posterior compression of middle-third esophagus unaffected by peristalsis and
• Esophageal neoplasm, very rare in infants causing asymmetric, irregular and abrupt stricture.

Conclusion

Congenital esophageal stenosis is a rare cause of childhood dysphagia occurring in various forms. Barium esophagogram plays a central role in diagnosis. Early diagnosis is very crucial for early management that may avoid surgical esophageal resection, reducing morbidity and mortality significantly in CES patients. Thus, CES should be strongly suspected as a cause of dysphagia in childhood and young adults.

References