# Laproscopic management of esophageal duplication cysts presenting in adults.

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### **Abstract**

Gastrointestinal duplications are commonly seen in children involving the esophagus in about 20% of all cases. However, esophageal duplication cysts are a rare cause of dysphagia in adults. We report a case of oesophageal duplication cysts in adult. This case report is presented mainly to describe that these predominantly congenital lesions may be symptomatic in adults and if so, must be treated expediously by surgical resection to prevent further complications.

Keywords: Cyst, Endoscopic ultrasound, Dysphagia, Computed tomography.

Accepted on October 25, 2018

### Introduction

Gastrointestinal duplications are commonly seen in children involving the esophagus in about 20% of all cases. Esophageal duplication cysts (EDCs) are congenital malformations of the posterior primitive foregut and often within the thoracic esophagus. Duplication cysts are congenital cystic malformation of the alimentary tract consisting of a duplication of the segment to which it is adjacent. Esophageal duplication cysts are the result of faulty intrauterine recanalization of the oesophagus and may account for 20% of all the gastrointestinal duplication cysts. Resection of oesophageal cysts is the treatment of choice, where excision of the cyst must take place with preserving the existing oesophageal mucosal and muscle integrity, and vagal nerves. In this case report, we describe the case of a 45-year-old female patient presented with progressive dysphagia, the diagnosis and its treatment.

## Case Report

A 45-year-old female patient presented with progressive dysphagia for 1 month. And history of regurgitation for 15 days. OGD revealed an extraneous impression in the posterior aspect of the terminal oesophagus just at the GE junction. Barium swallow showed a filling defect with sharp border in distal esophagus with slight displacement of the esophagus. CT chest showed well defined enhancing mass lesion of size 4.6  $\times$  $4.3 \times 4.2$  cm in the lower end of the esophagus extending into gastroesophageal junction. Dilatation of proximal esophagus is seen. Multiple endoscopic biopsies were negative and inconclusive. Intraoperatively, Lesion of size 4 cm noted in the distal esophagus adherent to the lower oesophagus along the left lateral and posterior aspects. laparoscopic transhiatal extramucosal excision done by a longitudinal incision made over the distal end of the esophagus and the lesion enucleated. The mucosa of the oesophagus was found to be intact and the seromuscular defect closed with 2-0 vicryl, the lesion retrieved and sent for HPE which was suggestive of oesophageal duplication cyst where the Cyst wall was lined by pseudostratified columnar eithelium with underlying thick fibrocollagen and of circular and longitudinal smooth muscle

and patient is on follow up. On 3<sup>rd</sup> POD Gastrografin study was done - No e/o leak and oral fluids was started (Figures 1 and 2).

Esophageal duplication cysts are the result of faulty intrauterine recanalization of the oesophagus and may account for 20% of all the gastrointestinal duplication cysts [1]. Based on their shape, they can be cystic (most common) or tubular and are seen most commonly involving the lower part esophagus (60%) [2]. Although most cystic duplication cysts are non-communicating, tubular duplications may occasionally communicate directly with the esophageal lumen [3].

## **Discussion**

The esophageal duplication cysts are usually asymptomatic and are rarely symptomatic. These rare symptoms may include dysphagia, chest pain, and are seen in children [4]. Other anomalies in these patients include concurrent vertebral anomalies [5] and other duplication cysts elsewhere in the alimentary tract also [6]. The differentials for esophageal duplication cyst may include specific mediastinal cysts like bronchogenic cysts, pericardial cysts or cystic degeneration



Figure 1. CECT (sagittal view) showing cystic lesion in the lower oesophagus on the left side.

of mediastinal tumors or other cysts including hydatid cyst, muellerian cysts [7].

The diagnosis is usually suspected on computed tomography, which classically may present as homogenous lesion with regular margins [1]. Hypodense or heterodense lesions may occur in the presence of pus, blood or thick insipid and result in diagnostic confusion [8,9]. EUS specifically characterize these lesions and distinguishes them from bronchogenic cysts by the following specific features namely the proximity to the esophagus, double muscle layer around cyst and due to nonvisualisation of cartilage in these cysts [10] (Figure 3).

Complications in oesophageal duplication cysts are rare causing haemorrhage [11], perforation, infection, malignant transformation [12,13] and even rupture [14] (Figures 4 and 5). EUS guided FNA has been used to confirm the diagnosis but EUS FNA may lead on to infection which may be refractory to the usage of prophylactic antibiotics [15]. Therefore, caution must be borne when planning EUS FNA in patients with esophageal duplication cysts.

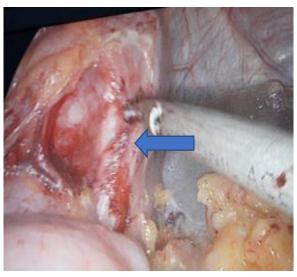


Figure 2. Laparoscopic dissection of the lower oesophagus showing lesion along the lateral aspect extending to the posterior aspect (arrow).

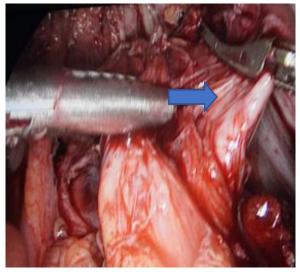


Figure 3. Dissection of the cyst (arrow) away from the muscle fibres of the lower oesophagus.



Figure 4. HPE specimen showing lesion of size 4 cm.



Figure 5. Post-operative barium swallows showing not showing any indentation or leak in the lower end of oesophagus.

## Conclusion

Resection of oesophageal cysts is the treatment of choice, where excision of the cyst must take place with preserving the existing oesophageal mucosal and muscle integrity, and vagal nerves [15]. Pathologically, these cysts are lined by mucous membrane with smooth muscle in their wall [6]. Ectopic gastric mucosa may be found in as many as 43% of patients [16]. Presence of other ectopic tissue like that of the pancreas is quite rare [17].

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