Introduction

Congenital diaphragmatic hernia (CDH) occurs in less than 5/10,000 births [1,2] with morbidity and mortality mainly related to its pulmonary effects secondary to muscular hyperplasia of the pulmonary vasculature and decreased arterial branching [3]. CDH is frequently syndromic with other malformations, most commonly of the gastrointestinal and genitourinary systems in 25% of cases, while the incidence of a coexisting pulmonary sequestration is much lower at around 5% according to the Congenital Diaphragmatic Hernia Study Group registry data[4]. While the majority of pulmonary sequestrations are found as intralobar malformations, approximately 25% manifest as extralobar sequestrations found within the thoracic or peritoneal cavity with their own pleural membranes and complete anatomic separation from the normal lung [5]. Due to increased availability of prenatal care and imaging, CDH is primarily diagnosed during the prenatal period or during early childhood [6]. Here we present a case of a healthy 24-year-old female presenting with gastric volvulus secondary to an undiagnosed CDH and the incidental finding of associated extralobar pulmonary sequestration mimicking gastric perforation.

Case Report

This 24 year-old previously healthy Somalian woman presented to the emergency department with 12 hours of worsening diffuse abdominal pain with nausea and emesis, initially attributed to reflux disease by patient. She was previously evaluated for similar symptoms in 2014 and was discharged after improvement with antiemetics and pain medication. On exam, the patient was tachycardic with abdominal distension and tenderness to palpation in left upper quadrant and epigastric region without rebound tenderness or involuntary guarding. Laboratory values were significant for lactic acidosis of 3.5 mmol/L and leukocytosis of 12,300/mm³. Chest X-ray showed thoracic herniation of the stomach (Figure 1), and computed tomography of the abdomen and pelvis demonstrated a mesenteroaxial gastric volvulus in the left thoracic cavity and a small adjacent fluid collection with internal foci of air concerning for a contained gastric perforation (Figure 2a-c).

Abstract

Congenital diaphragmatic hernias (CDH) carry significant morbidity and mortality risk due to compressive effects in the thorax during organogenesis and impaired development of lung tissue and vasculature. Prenatal screening now often allows for diagnosis of CDH in utero and for subsequent coordination of neonatal medical and surgical therapies to limit the early mortality risk. Here we discuss delayed presentation in a healthy 24-year-old female with diffuse abdominal pain secondary to gastric volvulus. Intraoperatively, this patient was found to have a previously undiagnosed congenital diaphragmatic hernia and concurrent extralobar pulmonary sequestration which mimicked gastric perforation on preoperative imaging. Concurrent congenital abnormalities of organogenesis with CDH and pulmonary sequestration have been described. These concurrent pathologies must be in the differential for surgeons operating on adults with late presentation of CDH. Sequestered lung tissue has a risk of malignant transformation and chronic infection necessitating thorough exploration of the hernia sac for identification of this concurrent pathology and careful resection of sequestered pulmonary tissue during CDH repair.

Keywords: Congenital Diaphragmatic Hernia; Extralobar Pulmonary Sequestration; Gastric Volvulus

Figure 1: Diaphragmatic herniation of stomach, pre-operative chest x-ray.
The patient was promptly taken to the operating room for laparoscopic reduction of her presumed type IV posterolateral lumbar paraesophageal hernia with gastric volvulus. Intraoperatively, a 14 cm left sided diaphragmatic hernia containing the volvulized stomach and spleen was visualized. Reduction of hernia and reversal of the volvulus was easily achieved laparoscopically and neither the stomach nor spleen appeared ischemic. Due to the size of the defect and extent of the intrathoracic hernia sac the decision to convert to open laparotomy for complete sac excision and hernia repair was made. Careful exploration and excision of the complete hernia sac within the chest revealed no obvious gastric perforation, however, an extralobar pulmonary sequestration was discovered in the area concerning for fluid collection on the CT (Figure 3).

The pulmonary sequestration was then resected with ligation of the anomalous arterial supply originating from the supra-celiac abdominal aorta and venous drainage to the left inferior pulmonary vein (Figure 4). No anatomic communication between the sequestered lung tissue and the normal bronchial tree or adjacent lung was present. The diaphragmatic defect was too large for a primary closure, so the small amount of tissue surrounding the defect was first plicated and then repaired with a Gortex dual mesh on lay (Figure 5).

This hernia was due to a congenital absence of the patient’s entire left diaphragm and not due to trauma, so the hiatal crus was able to be repaired to a normal size and the patient’s gastroesophageal junction became fixed in the abdomen. Because of this, gastropexy or an anchoring gastrostomy tube, although considered, were not deemed necessary. In addition to the mesh, the spleen abutted this repair.
reducing the risk of recurrence of the hernia. A chest tube and abdominal drain were placed and the abdomen was closed. The patient was extubated post-operatively without complications and monitored in the ICU for pain control and respiratory status. Post-operative day one chest x-ray demonstrated stable repair of the diaphragm with recruitment of the normal left lung and left mediastinal shift related to flattening of the diaphragm with repair and removal of the previous volume of sequestered lung tissue (Figure 6). The patient was discharged in stable medical condition on room air on post-operative day six. At subsequent one month follow up, patient is recovering well with mild dyspnea while supine that has been improving.

In this report, we present a rare case in which a congenital diaphragmatic hernia with extrapulmonary sequestration was discovered in an adult presenting as gastric volvulus. Due to early detection screening and the high mortality associated with CDH, 59% of cases are diagnosed within 24 hours of birth [10,11]. Late detection of congenital diaphragmatic hernias can present as gastric volvulus and wandering spleen, both of which were present in this patient. This is thought to be due to abnormal laxity of the diaphragm and/or abnormal or absent of peritoneal attachment to the stomach and spleen [12,13].

This patient’s asymptomatic medical history may be explained by the combination of limited prenatal screening and the presence of her pulmonary sequestration. It is hypothesized that pulmonary sequestrations may have a protective effect on congenital diaphragmatic hernias because the presence of extra-lobar tissue prevents herniation of abdominal contents into the chest cavity [14]. Additionally, the presence of a hernia sac in CDH cases is a risk-reducing prognosticator, with greater fetal lung growth, less visceral herniation, and better outcomes seen in infants with a hernia sac compared to those without [15]. This case highlights the need for providers to maintain a high index of suspicion for these concurrent pathologies and for complete surgical exploration of the intrathoracic hernia sac in patients presenting with possible congenital diaphragmatic hernia. These pulmonary sequestrations also warrant surgical excision even if asymptomatic and found incidentally. Extralobar pulmonary sequestration carries the risk of chronic infection, hemothorax, torsion, infarction, and potential for malignant degeneration [16-19]. Though the risk of infection of an extralobar pulmonary sequestration (14%) is lower than that of an intralobar pulmonary sequestration (91%), the presence of chronic inflammatory cells in an ELS is 35%. More than half of extralobar pulmonary sequestrations have co-existing congenital cystic adenomatous malformations, and 8.6% of malignant pleuropulmonary blastomas in infants and bronchoalveolar carcinomas in adults were associated with congenital cystic lung malformations [20]. Careful dissection must be used as the systemic vasculature supply can be quite large and may originate from either the thoracic or abdominal aorta posing risk for significant operative risk and blood loss if not precisely identified and controlled [21-23].

**References**


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