The predictive value of fruit juice in the esophagus-pleural fistula

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ABSTRACT

Esophageal-pleural fistula is a rare and challenging condition to diagnose and requires strong clinical suspicion in order to be recognized promptly. Chest computed tomography (CT) with contrast medium for oral contrast medium (OS) is the gold standard for diagnosis. The definitive therapy is purely surgical, except for a few selected cases that benefit from endoscopic therapy. Our case involves a 45-year-old woman who came to the Emergency Department with dyspnea and thoracalgia. Chest X-ray and high-resolution CT showed empyema and pleural effusion to the left hemithorax. The lack of improvement despite the therapy and a subsequent clinical finding gave rise to the suspicion of esophageal-pleural fistula, confirmed with CT with contrast medium for OS. In this case, we opted for endoscopy correction of the esophageal defects. The diagnostic delay and the pre-existing comorbidities (previous kidney transplant for chronic kidney disease from lupus nephritis, high blood pressure, familiarity with Ischemic cardiomyopathy) could justify the inauspicious course of our case.

Introduction

Esophageal-pleural fistula is a rare and challenging condition to diagnose. The etiology is varied, most esophageal-pleural fistulas are iatrogenic (mainly from surgical or endoscopic complications); spontaneous rupture (or Boerhaave syndrome) is the most frequent non iatrogenic cause, more rarely they are due to ingestion of extraneous bodies, mainly in the pediatric patient, barotrauma and penetrating trauma, neoplasms, esophagitis, ulcers, and infections.1-4 The clinical analysis is not typical and may be characterized by chest pain, loss of appetite and vomiting, dyspnea, signs and symptoms of pneumomediastinum, and/or pleural effusion. The gold standard for diagnosis is chest computed tomography (CT) with contrast medium for oral contrast medium (OS) or diagnostic-operative endoscopy in unstable patients. X-ray and chest ultrasound can confirm a clinical suspicion, with pneumomediastinum relief, subcutaneous emphysema, pleural effusion, and hydropneumothorax (PNX). Laboratory tests can show leukocytosis; the analysis of pleural fluid can be useful, with the detection of traces of food, high level of salivary amylase, and the presence of squamous epithelial cells. Complications of unrecognized or early treated esophageal–pleural fistula include empyema, mediastinitis, aspiration pneumonia, and cardiac tamponade.1,5-6 Medical therapy should be reserved for patients with limited perforation and in the cervical or mediastinal area, in the absence of obstructions and sepsis. It provides oral fasting and nutrition through a nasogastric tube or total parenteral nutrition, proton-pump inhibitors, and broad-spectrum and antifungal antibiotics, eventual drainage of the collected fluid. Surgical therapy with the repair of the defect by thoracoplasty is the therapy of choice, endoscopic treatment with stent positioning is reserved for patients with important comorbidities who could not tolerate surgery.7,8
Case Report

A 45-year-old woman came to ED with chest pain, dyspnea, and dyspepsia. In anamnesis, she had a kidney transplant for chronic kidney disease from lupus nephritis in antirejection and steroid therapy, high blood pressure in drug therapy, familiarity with ischemic cardiomyopathy. The vital signs showed Glasgow Coma Scale 15, temperature 36.0°C, tachydyspnea with 90% oxygen saturation, arterial pressure 100/60 mmHg with 85 bpm in β-blocker therapy. There was objectively shallow breathing with vesicular murmur reduced to the left field, rhythmic heart tones, treatable and indolent abdomen. The electrocardiogram did not show anything relevant. Laboratory tests showed mild anemia with moderate to severe thrombocytopenia, acute renal failure, elevated cardiac enzymes, indices of inflammation. The chest X-ray (Figure 1) showed a large left hydro-pleural layer with a mediastinal shift. The thoracic tube was positioned with draining of air and 1500 cc of turbid liquid. Due to rapid hemodynamic deterioration, the patient was intubated, supported with volemic and amine filling, and hospitalized in intensive care. During the 29 days of hospitalization in intensive care, further radiological and laboratory tests were carried out. All viral serologies and microbial cultures (blood, urinary) were negative, except for the growth of *Streptococcus salivarius* in the pleural fluid taken from thoracic drainage. From the thoracic drainage, remained a continuous and constant leakage of serum-turbid liquid. The chest-abdomen CT with intravenous contrast showed the evolution of the picture with the appearance of hydro-pleural layer with a mediastinal shift. The thoracic aorta for which we proceeded to the surgical toilet with mini-thoracotomy. Despite hemodynamic support therapy, broad-spectrum antibiotic therapy, and surgical correction of the empyema, the patient remained critical, and the markers of sepsis persisted in the laboratory data. During a period of weaning from mechanical ventilation, the patient assumed for OS a fruit juice, and the medical staff noticed the leakage of the same in the thoracic drainage in place. This event aroused a new clinical suspicion, and chest CT was performed with contrast medium OS, which confirmed the diagnostic suspicion of an esophageal-pleural fistula with spreading of the contrast medium before the thoracic aorta from continuity solution of the lateral esophageal wall in the left pleural cavity (Figure 2). Afterward, operative esophagogastroduodenoscopy was performed with the positioning of a metal stent and effective closure of the breach. However, the clinical conditions did not show improvement, and the progression of multiple organ failures (MOFS) was expected with the development of different antibiotic resistances in the various crops. The patient was transferred to a third level center specialized in the treatment of MOFS: another chest CT with contrast medium OS showed persistent esophageal-pleural fistula that was subject to surgical correction before metal esophageal stent removal. Residual esophageal stenosis was treated with endoscopic balloon dilatation and stent insertion: this procedure was complicated by hematemesis (major bleeding) and related type II myocardial infarction. The patient was discharged after 5 months with percutaneous endoscopic gastrostomy in place for nutritional support.

Discussion

Our patient presented at the Emergency Department with nonspecific symptoms, such as

![Figure 1. Chest x-ray.](image1)

![Figure 2. Chest computed tomography.](image2)
Case Report

is in itself rare.1,2

the correct diagnosis and treatment could also explain

literature, other cases have been reported with the

not uncommon for this type of pathology and, in

endoscopically self-expanding metal stent was

already in place, as input for diagnostic suspicion, is

in the diagnosis of the esophagus-pleural fistula is near

100%. On the other hand, a negative predictive value
cannot be defined, since the possibility of false-
negative test is almost unlimited.

Even the finding of an oropharynx pathogen, such as

S. salivarius, in the pleural fluid of thoracic drainage,

should have raised the clinical suspicion of a

continuous connection between the gastroenteric tract and the pleural cavity.3

In the absence of clinical suspicion, the use of

standard diagnostics, such as ultrasound, radiology,

and CT with intravenous contrast medium, did not

help the doctor in the definitive diagnosis because there were no pathognomonic findings of esophageal-

pleural fistula.

As per guidelines, chest CT with contrast medium for OS allowed, instead, a correct diagnosis. In our case, the therapeutic process was conditioned by the patient’s clinical instability, for which a surgical correction was not proposed. On the contrary, an endoscopically self-expanding metal stent was positioned considering the absence of the most frequent contraindications to this intervention (multiple breaches, lesion near the gastroesophageal junction, breach >6 cm long) and the absence of neoplastic tissue and/or obstructions of the bowel.7,8

What makes our clinical case interesting is the location of the esophageal-pleural fistula. Anatomically, the esophagus is in direct contact with pleura for a large area in the right hemithorax, while aorta is interposed between the esophagus and pleura on the left. The presence of a left esophageal-pleural fistula, therefore, represents a rarity in a pathology that is in itself rare.1,2

The esophageal-pleural fistula in this patient could recognize a multi-pathogenetic mechanism associated with her autoimmune disease. Patients with systemic lupus erythematosus often have an esophageal motility disorder, an inflammatory reaction of the esophageal muscle tunic due to the deposition of immune complexes and an ischemic alteration of the Auerbach plexus related to vasculitis. The risk of infectious esophagitis associated with immunosuppressive therapy is also increased.9

Conclusions

Esophageal-pleural fistula is a rare disease that requires high clinical suspicion to diagnose and establish the proper treatment promptly. Chest CT with contrast medium for OS represents the gold standard for diagnosis. Surgical correction of the breach is the treatment of choice, reserving endoscopy for a few selected cases or more unstable patients. If recognized and treated early, the esophageal-pleural fistula has a mortality of <10%, while if the diagnosis is delayed beyond 48 h the prognosis becomes severe.2-3 For the clinician in general, but also for the doctor of the emergency department, it is therefore useful to know the fundamentals of this rare pathology to be able to insert the esophageal-pleural fistula into the differential diagnosis scheme, and based on clinical suspicion, to prepare the most correct and timely diagnostic-therapeutic management.

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