

Boerhave Syndrom Revealed by Pneumodiastinum and Subcutaneous Emphysema: About One Case

Hajar Arfaoui¹, Asmaa Fahmi^{1*}, Hasna Jabri¹, Wiam Elkhatabi¹, Hicham Afif¹

¹Pneumology Professor, Hospital 20 Aout, 6 Rue Lahcen Al Aarjoune, Casablanca 20250, Morocco

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*Corresponding author: Asmaa Fahmi

Pneumology Professor, Hospital 20 Aout, 6 Rue Lahcen Al Aarjoune, Casablanca 20250, Morocco

Abstract

Case Report

The Boerhaave syndrome or spontaneous rupture of the esophagus is a rare pathology that represents 15% of all esophagus ruptures, caused by a sudden increase in intraluminal pressure due to vomiting efforts. The prognosis depends on the early instatement of the adapted treatment. We report the case of a 35-year-old patient who showed provoked repetitive vomiting since one month. He consulted at the pneumology service because of brutal thoracic pain since 4 days associated to a violent epigastric pain, complicated three hours after a respiratory distress. The clinical exam oriented us to the esophagus rupture hypothesis (Boerhaave syndrome) confirmed by the imaging that showed the presence of a pneumomediastinum, a hydropneumothorax, an important emphysema dissecting soft tissue, and a clear distension of the lower border of the thoracic esophagus. In front of the aggravation of the respiratory difficulty, the patient was transferred to the intensive care service.

Keywords: Pneumomediastinum; boerhaave syndrome; emphysema.

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INTRODUCTION

The Boerhaave syndrome or the spontaneous rupture of the esophagus is a rare clinical entity caused by the intraluminal pressure rising following the vomiting efforts. In the absence of typical symptoms, the diagnosis is difficult. The prognosis depends on the early stage of medical support [1]. We report a case of a pneumomediastinum associated to a hydropneumothorax and a subcutaneous emphysema happened to a young man due to a vomiting effort and we discuss the diagnosis difficulties, medical support modalities and the prognosis of this pathology.

CASE REPORT

MR. H.S, 35 years old, without particular pathologies antecedents in particular no recent trauma, he reported repetitive provoked vomiting since one month in a psychological context (bulimia probably). He consulted at the pneumology service due to brutal thoracic pain since 4 days and dyspnea at any effort associated to an epigastric pain complicated three hours later by a respiratory distress which needs hospital care.

The clinical exam showed a dyspneic patient, tachycardic, 88% saturation at air ambient. The

pulmonary exam showed a mixed effusion syndrome. We found at the neck exam a subcutaneous cervical emphysema extended to the superior members and to the left eye. The abdominal exam showed a contracted abdomen, painful at the epigastrium.

The initial biological exams showed an increased leukocyte count of 29 000 leukocytes/mm³ at PNN predominance, a CRP at 348 mg/l, a renal failure stage 3A with a DFG at 56 ml/min/1.73.

The esophagus rupture hypothesis after vomiting efforts (or Boerhaave syndrome) brought us to radiological exams without delay. The standard chest X-ray confirmed the existence of a pneumomediastinum associated to a subcutaneous cervical and thoracic emphysema and a homogeneous dense opacity at the inferior base (Figure 1). The abdomen without preparation didn't show peritoneal pneumonia. The Thoracic computed tomography showed right hydropneumothorax with a large pneumomediastinum extending along the cervical thoracic region realising an important emphysema dissecting, infiltration of mediastinal fat, we note a frank distention of the thoracic esophagus at heterogeneous content lower border (Figure 2) orienting, with the absence of any

other etiology, to the syndrom of Boerhaave. The ingestion of the hydrosoluble couldn't be possible in front of the aggravation of the respiratory distress.

The patient was immediatly transfered to the intensive unit care where he did a thoracal drainage. After being stabilized, he was sent for emergency abdominal surgery, during which he died.



Figure 1: Chest x-ray face (A): pneumomediastinum (black arrow) and subcutaneous cervical and thoracal emphysema (white arrow)

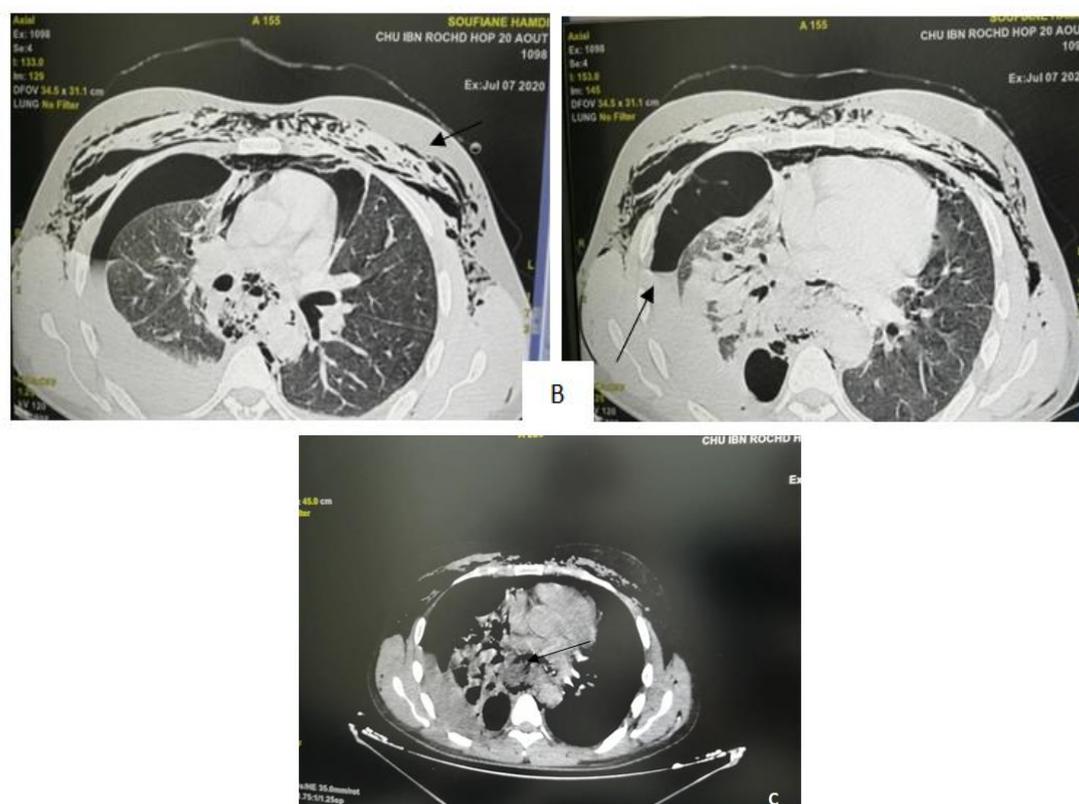


Figure 2: Thoracic computed tomography in parenchymal window (B) confirmed right hydropneumothorax with pneumomediastinum and subcutaneous emphysema Mediastinal window (C) distanciation of the thoracal esophagus

DISCUSSION

Boerhaave syndrome consists of spontaneous longitudinal transmural rupture of the esophagus, usually in its distal part. It generally develops during or after persistent vomiting as a consequence of a sudden

increase in intraluminal pressure in the esophagus [2]. It's a rare pathology that represents 15% of all esophagus ruptures [3]. It affects mainly people between 40 and 70 years with a man majority [4]. Our patient was a 35 years old.

The clinical manifestations of an Boerhaave syndrome are variables and make the diagnosis difficult. In 50% of the cases, it is manifested by Mackler's triad: vomiting, lower thoracic pain and subcutaneous emphysema [5]. Signs of acute respiratory distress will dominate the clinical board testifying a pleural invasion [5].

Some other symptoms such as dyspnea, fever are usual and may be confused with other diseases such as myocardial ischemia, aortic dissection, pericarditis, spontaneous pneumothorax, pneumonia, perforated peptic ulcer and pancreatitis [6]. In our patient case, good anamnesis and clinical exam, allowed to guide our diagnosis to the spontaneous esophagus rupture, so the realisation of non invasive other complementary exams that allowed to eliminate the differentials diagnosis.

The standard chest x-ray lead the diagnosis showing pneumomédiastinum or hydropneumothorax and subcutaneous emphysema.

The Thoracic computed tomography confirmed the presence of air in the médiastinum and subcutaneous tissue were not visible at the radiography and show sometimes a pneumothorax, a pneumoperitoneum, or a pneumopericardium, [7, 8]. Esophagography (before the 48 th hour) is an important imaging examination for confirming the diagnosis and the location of perforation because it shows extravasation of contrast into the pleural space [9].

The pronostic depend of the early stage of care support. Every delay of diagnosis rise the mortality wich may reach 100% [10].

The treatment for Boerhaave syndrome is both conservative, endoscopic, and surgical. The choice between theses treatments depending on the time that has elapsed since development of the rupture and its recognition and treatment, clinical stability and the scope the thoracical contamination [10]. The chirurgical treatment is preferred when septicemia is installed, while the endoscopical approche is preferred when there is no sign of septicemia and/ or a minimal contamination of the pleural cavity and médiastinum. A conservative support consistent on an antibiotic treatment and a drainage of abscesses [11].

CONCLUSION

Spontaneous rupture of the esophagus is a rare clinical entity with a high mortality rate. Early clinical suspicion will lead to timely diagnosis and maximize the survival chances for the patient.

Conflict of interest: No conflict of interest.

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