A Case of Spontaneous Pneumomediastinum without Direct Cause

Sam Beom Lee, Jung Ho Kim, Byung Soo Do
Department of Emergency Medicine
College of Medicine, Yeungnam University, Daegu, Korea

Introduction

Spontaneous pneumomediastinum is a rare, self-limiting benign disease that usually occurs in young patients without an obvious precipitating factors (1). Emergency physicians may be encountered in diagnosis and treatment in the emergency department, and is occasionally be able to be considered as an emergency condition in some cases.

We would present our experience in treating a case of spontaneous pneumomediastinum without direct precipitating cause, he was a 15-year-old boy treated with conservative therapy.

Case

A 15-year-old boy presented with left chest discomfort for 12 hours not associated with pleuritic chest pain or dyspnea. He did not cough or vomit vigorously just before this presentation. But it was aggravated by deep inspiration and radiated to the left shoulder. He did not suffer from other symptoms. He lately visited in emergency care center.

He did not have significant past medical history such as tuberculosis. There was no history of recent trauma, upper respiratory infection, or foreign body swallowing.

Vital signs were within the normal limit, blood pressure 130/80 mmHg, heart rate 60/minute, respiratory rate 18/minute, and body temperature 36.7°C.

Physical examinations were essentially normal except faintly audible mediastinal crunching sound at the left mid-precordium (Hamman’s sign).

He was checked on blood gas analysis, chest X-ray, electrocardiogram, and computerized tomography (CT). Blood gas analysis showed pH 7.434, PaCO₂ 35.4 mmHg, PaO₂ 109 mmHg, base excess 0.8 mmol/L, HCO₃⁻.
Fig. 1. Chest PA showed mediastinal air and “continuous diaphragm sign” and small amount of air in left pericardial space.

Fig. 2. Chest CT showed pneumomediastinum and pneumopericardium.
24 mmol/L, and O2 saturation 98.4%. Chest X-ray showed the pneumomediastinum and small amount of pneumopericardium, but rib fracture, pneumothorax, and other visible active lung disease were not seen on this study (Fig. 1). Electrocardiogram showed no evidence of ischemia or infarction such as ST changes, T inversion. Chest CT drip infusion study was taken by immediately and showed the pneumomediastinum with extending to interstitium, but visible bullae or pneumothorax were not seen (Fig. 2).

And also he immediately underwent a esophagoscopy, which did not demonstrate the evidence of esophageal perforation.

He was placed on oxygen by nasal catheter, and the following day, he did not complain such symptoms any more and the follow-up chest X-ray showed that the pneumomediastinum had began to resolve. So he was discharged that day without admission.

Discussion

Spontaneous pneumomediastinum is rare disease or condition and usually occurs in young patients without an apparent precipitating factor or disease (1).

Hamman (2) first described spontaneous pneumomediastinum in 1939. His description of audible crepitation occurring with the heartbeat on chest auscultation is known as “Hamman’s sign”, which was present in 9 of 22 cases (1).

Pneumomediastinum results from the rupture of terminal alveoli into the lung interstitium and the dissection of air along the pulmonary vasculature toward the hilum, with eventual extravasation into the mediastinum (3). This theory could explain our case which not confirmed by precipitating cause.

Spontaneous pneumomediastinum is often reported to be the result of asthma, labor, diabetic ketoacidosis, Hodgkin’s disease after irradiation, chemotherapy, coughing or forceful straining during exercise, Valsalva maneuver, etc, in the many literatures (4–10).

Classic symptoms include retrosternal chest pain, dyspnea, cough, cervical pain, but are not nonspecific. In our case, he only complained chest discomfort. Other symptoms include general weakness, dysphagia, back pain, sore throat, etc (11–15). Physical finding may include subcutaneous emphysema on the chest, back, or neck area (14). The precordial crunching sound described by Hamman (2) in 1939 can be audible at substernal area (Hamman’s sign), which can be confused with the pericardial friction rub. Abolnik said that the precordial crunching sound was heard in 40% of his cases (16).

The presence of pneumomediastinum was confirmed by chest X-ray films, and CT scan was performed to exclude any underlying lung disease such as bullous lung disease in our case. Esophagoscopy or esophagogram
are essential to demonstrate esophageal perforation. Emergency physicians initially have to study this methods for differential diagnosis including pleural, pulmonary, cardiac, esophageal, and musculoskeletal causes (1,17, 18).

No special treatment is required in spontaneous pneumomediastinum and the patient can be discharged after short-term observation without admission. Supplemental oxygen therapy may be useful, and careful observation, bed rest, analgesics can be needed in most cases. Recurrence and complications are unusual because spontaneous pneumomediastinum is self-limiting condition in most cases (1,14,17).

References


—A Case of Spontaneous Pneumomediastinum without Direct Cause—