Spontaneous Pneumomediastinum and Subcutaneous Emphysema: A Two Cases Reports

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Abstract

Spontaneous pneumomediastinum (SPM) is an uncommon, mostly benign disorder that usually occurs in young adult males without any apparent pre-existing factor or disease. SPM responds extremely well to conservative treatment, without recurrence in the great majority of cases. We report two cases of SPM and pneumothorax in two young males following a forceful vomiting and a bout of sudden coughing with no underlying lung lesion.

Keywords: Coughing, vomiting, spontaneous pneumomediastinum, subcutaneous emphysema, pneumothorax.

INTRODUCTION

Pneumomediastinum is the presence of air in the mediastinum, it is considered as secondary when a causative factor is identified. Spontaneous pneumomediastinum (SPM) is a rare condition which occurs in the absence of pulmonary disease or any other inducing factors [1, 2]. We report two cases of SPM and pneumothorax in two young males following a forceful vomiting and a bout of sudden coughing with no underlying lung lesion.

CASE REPORTS

Case 1

A 41-year-old man, working as a builder and nonsmoker but exposed to second hand smoke for 20 years at work presented to the emergency department with a 4-day history of spastic cough due to a cold. He had no known allergies and no history of pulmonary or cardiovascular disease. He had a consultation at a private practice at the beginning of his cold and was given oral antibiotics (macrolide) and aspirin. One hour after taking his medicine, he developed chest pain, dyspnea and swelling of the face, neck and upper chest.

At his admission, the patient presented with respiratory distress syndrome so he was immediately transferred to the intensive care unit of the emergency department and put under oxygen mask. On physical examination, crackling-feel was noted in the neck, the clavicular areas, and upper chest. Pulmonary and cardiac auscultations were normal.

The chest X-ray on admission showed a large subcutaneous emphysema and pneumomediastinum (Figure-1). The computed tomography revealed a pneumomediastinum, subcutaneous emphysema and bilateral pneumothorax (Figure-2). The lungs were otherwise clear.

The patient was given nasal oxygen and codeine for the pain. He was kept for close observation in the hospital during the next 2 days. The subcutaneous emphysema improved clinically and the chest X-ray after 2 days showed significant resolution of the emphysema and pneumomediastinum (Figure-3).

Case 2

29-year-old patient admitted to emergency department for vomiting secondary to diabetic ketoacidosis complicating his type 1 diabetes. The physical examination revealed subcutaneous emphysema and epigastic tenderness, urine test strips shows three glucose crosses and two acetone crosses, the blood bicarbonates level was 12 mmol/l. The chest radiograph showed a large subcutaneous emphysema and pneumomediastinum. The CT scan of chest revealed a right pneumothorax, a pneumomediastinum with subcutaneous emphysema extending to the lower neck (Figure-4). Intravenous fluid perfusion and insulin were administered in order to treat the acidocetosis; as well as nasal oxygen which was given to help resolve the pneumothorax. After glycemia control and conservative treatment of the pneumothorax and pneumomediastinum, the patient was discharged. The
pneumomediastinum and pneumothorax resolved spontaneously according to the follow up imagery.

Fig-1: Chest X ray showing pneumomediastinum, and subcutaneous emphysema

Fig-2: Section 1-computed tomography (thorax) revealing apneumomediastinum with subcutaneous emphysema and bilateral pneumothorax

Fig-3: Chest X-ray after 2 days showing a significant resolution of the emphysema and pneumomediastium
DISCUSSION

The mediastinal cavity is defined as the area demarcated by the pleural cavities laterally, the cervicothoracic inlet as an upper limit and the diaphragm as an inferior one [3]. Pneumomediastinum (PM) also referred to as mediastinal emphysema was originally described in 1819 by Laënnec [4, 5] and is defined as the presence of free air in the mediastinal cavity. The discovery of free air in the mediastinal cavity is often thought of as a phenomenon secondary to a more serious medical condition. PM is broken down into two categories: the first one is as a result of trauma or caused iatrogenically due to endoscopic or other therapeutic procedures. Such cases are referred to as secondary PM and have a clear causative factor. The second one is the presence of free air in the mediastinal cavity without a clear etiology, which is referred to as spontaneous pneumomediastinum (SPM). It is a rare condition, reported with incidence of less than 1/44000, and often has a benign outcome [6].

Cough is an explosive expiration which is a physiological protective mechanism allowing the clearing of the trachea and bronchi from secretions and foreign material. However, the sudden increase in intrathoracic pressure during coughing is a risk factor of alveolar over distension and rupture. The forceful contraction of expiratory muscles limits the expansion of lungs to their total lung capacity by splinting the chest wall and thus usually preventing any volutrauma that may occur and causes alveolar volume disruption of alveolar walls [7]. Rarely, the transient increase in intrapulmonary pressure during coughing may also cause alveolar rupture. Coughing is usually uneventful, but occasionally results in microscopic alveolar ruptures [6].

SPM usually follows vomiting, intense exertion, and Valsalva maneuvers, all of which cause sudden increase in intra-alveolar pressure without specific cause such as trauma. Studies have also reported its association with cocaine and marijuana use and asthma attacks [8, 9]. Cough has also been suggested as a possible cause of pneumomediastinum [8, 10]. In this reported case, cough was the only etiology found justifying the occurrence of bilateral spontaneous pneumothorax, pneumomediastinum and subcutaneous emphysema in a otherwise healthy young man with no known respiratory condition. No obvious preceding events or factors are detected in nearly 20% of all mediastinal emphysema patients [8].

Chest and neck pain, dyspnea, subcutaneous emphysema, hypotension, dysphagia, and cough are common symptoms of SPM. Chest pain is characteristically sudden and of pleuritic type, and may radiate to the back and shoulders. Subcutaneous emphysema has a prevalence ranging from 40% to 100% [7] and occurs when intrathoracic air leaks into the soft tissues after pneumothorax or pneumomediastinum. Hamman’s sign is another clinical finding of SPM and was described by Hamman as the crunching, rasping sound, synchronous with the heartbeat during auscultation.

There is no consensus on the method of diagnosis of SPM [11]. While some authors report that simple chest X-ray alone is enough for the diagnostic [12, 13], others state that chest X-ray is sufficient for the diagnosis of SPM if a patient did not have preceding trauma, foreign body inhalation or vomiting, and that further investigations are not required if the patient is stable [14]. On the other hand, some authors suggested that all patients should undergo chest CT to diagnose SPM in order to investigate any other underlying conditions [9]. It is also recommended that chest X-ray is the initial method of diagnosis, but chest CT and other investigations should be performed if SPM and other conditions are suspected [15, 16].

Primary SPM is retained after all other causes of mediastinal emphysema are excluded and can be treated easily. However, this condition may lead to severe complications such as tension pneumomediastinum that can be fatal and occurs in case of increased mediastinal pressure. Once the diagnosis of SPM is established, the patient has to be hospitalized for a minimum of 24 hours observation to prevent potential complications which may require a surgical intervention [8]. Treatment includes bed rest, analgesic and oxygen administration, mediastinal emphysema usually resolves within several days.
CONCLUSION

In this article we described two cases of spontaneous pneumomediastinum, subcutaneous emphysema and pneumothorax that were induced by coughing or vomiting. The occurrence of the swelling right after the intake of antibiotics and aspirin as well as the respiratory distress could have induced a false diagnosis of an allergy. Also the incidence of chest pain complaints is lower in cases of pneumomediastinum accompanied by DKA than in cases of generalized pneumomediastinum, and dyspnea can be regarded as a symptom of DKA; this increases the chances of missing the correct diagnosis, however, life-threatening complications (pneumothorax, pneumopericardium, and mediastinitis) might accompany this condition. Therefore, clinicians need to include this complication in the differential diagnosis. The diagnosis of SPM was achieved thanks to a thorough physical examination and the immediately performed chest X-ray. The follow up was favorable and the patients recovered spontaneously without any complications.

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