

Research Article

# Spontaneous pneumomediastinum associated with COVID-19: Rare complication of 2020 pandemic

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## Abstract

Spontaneous pneumomediastinum (SPM) is a rare condition, more commonly seen in patients with history of asthma, chronic obstructive pulmonary disease, infections, or drug users. Today, we face one novel virus that has caused an outbreak of acute respiratory illness, affecting over a million individuals worldwide. New knowledge is being gained of the virus and possible complications have been seen. Following, we present the case of a 71-year-old man with diagnosis of COVID-19 pneumonia complicated with spontaneous pneumomediastinum.

## Background

SPM or spontaneous mediastinal emphysema is a rare condition characterized by free air in the mediastinum not preceded by thoracic trauma, surgery, or any other medical procedure [1-3]. It was first described by Louis Hamman in 1939, which is why it is also known as Hamman's syndrome [1,2,4,5]. SPM typically presents with symptoms of pleuritic chest pain, dyspnea and cough. Upon suspicion of SPM, the goal is to evaluate for potential triggers, exclude differential diagnosis such as pneumothorax or esophageal rupture and finally evaluate for possible complications such as tension pneumomediastinum. Treatment is supportive and includes rest, analgesia and avoidance of maneuvers that may increase the pulmonary pressures. In most cases, the condition self-resolves and rarely, invasive surgical intervention is required and most patients do not develop long-term sequelae or recurrence.

## Introduction

SARS-CoV-2 disease commonly known as COVID-19, is an emerging viral infection which started in December 2019 and has become one of the world's deadliest pandemic

known in history. Everyday new knowledge is acquired, on the various forms of clinical presentations and radiological changes which exist in patients infected with COVID-19. The most known radiological findings include ground glass opacities and consolidations with peripheral distribution. In approximately 1% of the cases, spontaneous pneumothorax have been identified. These findings usually serve as potential markers of COVID infection. Nevertheless, new complications are observed such as spontaneous pneumomediastinum (SPM). SPM is a rare condition characterized by free air in the mediastinum not preceded by thoracic trauma, surgery, or any other medical procedure [1-3]. It was first described by Louis Hamman in 1939, which is why it is also known as Hamman's syndrome [1,2,4,5]. SPM typically presents with symptoms of pleuritic chest pain, dyspnea and cough. It is unknown what is the precise mechanism of spontaneous pneumomediastinum after COVID infection, but possible mechanisms include diffuse alveolar damage secondary to a severe pneumonia. In severe pulmonary infections, repeated cough spells, increase distal airway pressure and result in increased alveolar rupture and gas leakage to the peribronchovascular interstitium that may extend to the mediastinum. Upon suspicion of SPM, the goal is to evaluate for potential triggers, exclude differential

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**Keywords:** Spontaneous pneumomediastinum; COVID-19; Pneumonia



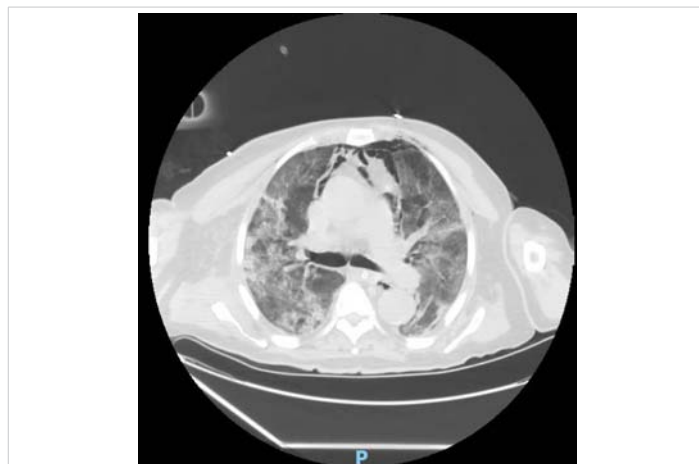
diagnosis such as pneumothorax or esophageal rupture and finally evaluate for possible complications such as tension pneumomediastinum. Reported cases secondary to viral infection are very rare. Further studies of SP and COVID-19 infection are needed in order to determine its role as prognostic factor or disease progression marker. Treatment for SP is mainly supportive and includes rest, analgesia and avoidance of maneuvers that may increase the pulmonary pressures. In most cases, the condition self resolves and rarely, invasive surgical intervention are required and most patient do not develop long term sequela or recurrence.

## Case presentation

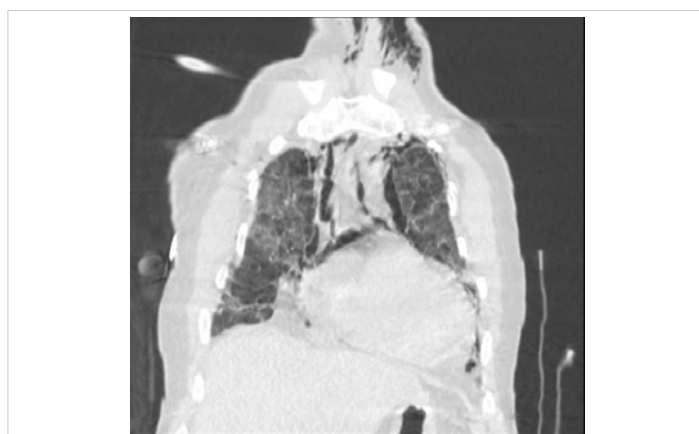
We present the case of a 71-year-old man with past medical history of arterial hypertension and prostate cancer treated with 44 sessions of radiotherapy presented to the emergency room with complains of dyspnea on exertion of five (5) days of evolution. Review of systems included anorexia, nausea and diarrhea. Patient reported recent travel to the Dominican Republic. He denies fever, chills, unintentional weight loss, chest pain, or sick contacts. Medications included Nifedipine, Atenolol, Tamsulosin and Hydralazine. Vital signs remarkable for tachypnea of 30 bpm with peripheral oxygenation of 79% at room air. Physical examination revealed notable respiratory distress and bilateral bibasilar rales.

Laboratories showed leukocytosis with lymphopenia with slight toxic granulation and normochromic normocytic anemia. Laboratory remarkable for elevated Ferritin 2617 ng/mL, elevated D-Dimer 24.3 mg/L, elevated fibrinogen 687 mg/dL, central bicarbonate of 18mEq/l, hyponatremia 130mEq/l, hypochloremia of 97mEq/l, prerenal azotemia with BUN/Cr >20, lactate dehydrogenase of 247 IU/L, and erythrocyte sedimentation rate of 64. CXR shows parenchymal increased interstitial markings and bilateral pulmonary opacities more confluent towards the lung bases with an elongated aorta with atherosclerotic changes. Nasopharyngeal PCR swab test was positive for COVID-19 infection. Patient was treated with mechanical ventilation and lung protective ventilation, dexamethasone therapy, and IV antibiotics. After 5 days of admission, patient developed ARDS as complication of COVID-19. Chest CT (Figure 1) demonstrated bilateral diffuse areas of ground-glass opacity and consolidations, features compatible with COVID-19 pneumonia. On MICU day 28, patient developed a respiratory distress for which chest CT (Figure 2) was performed showing a significant pneumomediastinum with associated subcutaneous emphysema extending to the thoracic wall and lower neck, anterior apical pneumothorax, and cardiomegaly.

Given spontaneous pneumomediastinum was highly likely a complication of SARS-CoV 2 infection, management consisted of supportive care.



**Figure 1:** Lung parenchymal shows increased interstitial markings as well as peribronchovascular cuffing, diffuse bilateral patchy ground glass and confluent densities noted. Small pleural effusion.



**Figure 2:** Diffuse pneumomediastinum and subcutaneous emphysema extending to the thoracic wall and lower neck.

## Conclusion

Spontaneous pneumomediastinum is caused by a pressure gradient between the alveoli and pulmonary interstitium leading to alveolar breakdown. In cases of infection by SARS-CoV, pneumomediastinum may be related to damage and rupture of alveolar membrane caused by the virus. Although pneumomediastinum is usually considered a self-limiting condition, with an unknown precise pathological mechanism; there is limited literature resources and reports on potential complications associated with COVID-19 pneumonia, overall incidence has not yet fully well-established. It is important to highlight this possible complication of COVID-19 pneumonia, which serves as an uncommon marker of worsening disease. In the era of SARS-CoV 2 pandemic, physicians need to be aware of rare and uncommon complications of this novel virus.

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