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A Very Rare Complication of Covid-19: Penumothorax and Penumomediastinum

Aynur Yurtseven*

Department of Emergency Medicine, Ankara diskipi yildrim beyazit tezinin and research hospital, Turkey

*Corresponding author: Aynur Yurtseven, Department of Emergency Medicine, Ankara diskipi yildrim beyazit tezinin and research hospital, Turkey, E-mail: aynuryurt7@gmail.com

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Abstract

Background: Pneumothorax and pneumomediastinum are defined as free air within pleural ve mediastinal spaces, respectively. Spontaneous pneumothorax and pneumomediastinum combination is a very rare complication of Covid-19 pneumonia. With this case report we want to underline etiology and possible pathophysiology of these rare complications of Covid-19. At the same time, with this case, we wanted to contribute to the literature in terms of complications that may be seen due to Covid-19 infection.

Case Report: 24 years old male patient visited emergency service due to dyspnea and pleuritic chest pain since 3 days. High resolution thorax CT scan revealed spontaneous pneumothorax, pneumomediastinum and Covid-19 pneumonia. Due to absence of predisposing factors, we considered pneumomediastinum and pneumothorax as rare complications of Covid-19 and administered patient to ICU unit.

Conclusion: Spontaneous pneumothorax and pneumomediastinum are rare but significant complications of Covid-19 and should be kept in mind as a differential diagnosis of chest pain and dyspnea at these patient group. Further researches are needed for etiology of pneumomediastinum and pneumothorax due to Covid-19.

Keywords: COVID-19 pneumonia; Spontaneous pneumothorax; Spontaneous pneumomediastinum; Chest pain.

Introduction

Since 2019, outbreak of Covid-19, there has been an increasing trend at numbers of Covid-19 patients and numbers of complication associated with Covid-19. In this case we want to underline a rare complication; spontaneous pneumothorax and spontaneous pneumomediastinum, due to Covid-19 pneumonia at a patient with no history of positive pressure ventilation or trauma.

Case

24 years old male patient administered to emergency service with pleuritic chest pain and dyspnea of 3 days. Patient had cough with sputum and headache, without loss of taste and loss of smell. He received 2 doses of vaccination with Biontech[®], last dose received 2 months ago. Physical examination revealed diffuse rhonchus bilaterally on auscultation, blood pressure 130/80 mmHg, heart rate of 134, body temperature of 36.5°C, plasma glucose 139 mg/dL, respiratory rate of 24 and saturation of 84 at room air. ECG had no specific features with sinus tachycardia. Patient had WBC 18.0000/uL, PLT 229.000/uL, serum lactate 2, D-Dimer 620 ng/dl and troponin within normal range. Covid-19 RT-PCR test was positive. High resolution computed tomography showed bilateral minimum pneumothorax and pneumomediastinum with perihilar and peribronchovascular focal ground glass opacities compatible with Covid-19 (Figure 1). During emergency service follow-up patient developed hemoptysis and severe dyspnea. Thoracic surgery and cardiovascular surgery consultations are done and no surgical interventions planned. For further follow-up ICU admission was planned.

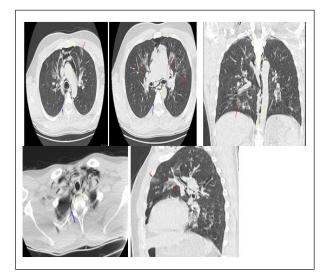


Figure 1: Images of pneumomediastinum and plastering pneumothorax in a patient who presented with Covid-19 pneumonia. Chest computed tomography scan showing

Vol.8 No.2:210

pneumomediastinum shown with yellow arrows, ground glass image compatible with Covid-19 pneumonia shown with red arrows and plastering pneumothorax shown with blue arrows.

Discussion

Covid-19 virus passes through cell membrane *via* Angiotensin Converting Enzyme 2 receptor (ACE 2) and induces wide tissue damage by stimulating immune system and causing overreaction. Interleukin 6 (IL 6) is thought to be the main trigger of this overraction. Another important factor is IL 1B known as main causative agent of widespread lung inflammation and fibrosis [1]. Most frequent radiological findings are early phase reticular changes and mid-late phase ground glass opacities. Distribution of inflammation through lungs is usually bilateral, peripherally located and consolidation increases sequentally [2].

During Severe Acute Respiratory Syndrome (SARS) outbreak at 2002-2003, a research from Hong Kong, showed increased incidence of spontaneous pneumomediastinum (11,6%) [3]. For Covid-19 patients incidence of spontaneous pneumothorax and spontaneous pneumomediastinum is not well known yet. Spontaneous pneumothorax incidence of SARS-CoV-2 infected patients is 1%-2%, and spontaneous pneumomediastinum incidence is very low [4-6].

We found 15 cases of spontaneous pneumomediastinum of Covid-19 patients from literature review. 7 of these 15 patients had no history of invasive or non-invasive mechanical ventilation and no concomitant spontaneous pneumothorax was diagnosed. 6 patients has concomitant spontaneous pneumothorax. According to literature review, some of these patients developed pneumothorax and pneumomediastinum during follow-up while some had these complications at first contact with healthcare system [7]. For our case it was at first medical contact. Researchers declare that the main underlying mechanism for pneumomediastinum and pneumothorax is rather alveolar rupture associated with intense alveolar damage and air leakage throughout bronchoalveolar sheath and not the viral load. Main determinant is thought to be immunological damage. Also, literature review of these patients who mainly had no significant medical history or smoking habit support immune reaction theory. Our case, likewise, had no medical history or smoking habit.

Since spontaneous pneumomediastinum is mostly a selflimiting clinical condition, a conservative approach is preferred first. However, in order to prevent possible serious complications in the course of the disease, it is very important to detect predisposing factors and take precautions accordingly. Thorax CT scan has a very important role in determining the cause of respiratory distress and pulmonary complications associated with COVID-19 in the early period.

Results

Spontaneous pneumomediastinum and spontaneous pneumothorax should be considered in the differential diagnosis of patients with Covid-19 whose general clinical condition worsens. More research is needed to elucidate the underlying pathophysiological process and to determine its true incidence.

Acknowledgments

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Conflict of Interest

The authors have no conflict of interests to declare.

Ethics Approval

Informed consent was obtained from family of the patient prior writing this case report.

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