Mortality due to COVID-19 in a Patient with Severe Asthma Receiving Omalizumab Treatment: A Case Report

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ABSTRACT

Coronavirus disease 2019 (COVID-19) originated in Wuhan, China, and caused a pandemic in the world. SARS-CoV-2 infections have a great health risk, especially in patients with chronic diseases. GINA (Global Initiative for Asthma) recommends that asthmatic patients continue their controller medications including inhaled corticosteroids, during the COVID-19 outbreak. GINA recommends adding biological agents to the treatment of selected patients with severe asthma to reduce the need for oral corticosteroids (3).

There are concerns that COVID-19 may be more severe, especially in patients with severe asthma who use biological agents. However, the efficacy and safety of treatment with biological agents during SARS-CoV-2 infections are currently unknown. In this case report, we present a severe asthmatic patient who had been using omalizumab for five years and died due to COVID-19.

Keywords: COVID-19, asthma, omalizumab, severe allergic asthma, anti-IgE

INTRODUCTION

Coronavirus disease 2019 (COVID-19) originated in Wuhan, China, and caused a pandemic in the world (1). SARS-CoV-2 infections have a great health risk, especially in patients with chronic diseases (2). GINA (Global Initiative for Asthma) recommends that asthmatic patients continue their controller medications including inhaled corticosteroids, during the COVID-19 outbreak. GINA recommends adding biological agents to the treatment of selected patients with severe asthma to reduce the need for oral corticosteroids (3).

There are concerns that COVID-19 may be more severe, especially in patients with severe asthma who use biological agents. However, the efficacy and safety of treatment with biological agents during SARS-CoV-2 infections are currently unknown. In this case report, we present a severe asthmatic patient who had been using omalizumab for five years and died due to COVID-19.
shortness of breath increased and she presented again to the emergency department. On the physical examination performed in the emergency department, fever was 38.2 °C, pulse 108/min, respiratory rate 29/min, blood pressure 90/58 mmHg, and saturation 88% while taking 4 L/min supplemental nasal oxygen. On the respiratory system physical examination, there were rales and rhonchi in both lung fields. Other physical examination findings were normal. The laboratory tests showed: ferritin 86.9 µg/ml (normal: 11-307), C reactive protein (CRP) 20.4 mg/dL (normal: 0-0.5), lactate dehydrogenase (LDH) 251 mg/dL (normal <247), lymphocyte count 900 µL (normal: 1200-3600), sedimentation rate 39 mm/hour (normal: 0-25), procalcitonin 0.023 ng/mL (normal: 0-0.1), troponin-1 2.7 ng/L (normal: 8.4-18.3), and D dimer 0.47 mg/L (normal: 0-0.55). The thoracic computed tomography of the patient revealed opacities and ground-glass densities in the peripheral, lower, and posterior zones in both lungs (Figures 1, 2). Favipiravir was started at a 2x1600 mg first-day loading dose followed by 2x600 mg maintenance dose for five days. Dexamethasone 8 mg/day was added to the treatment and the patient was transferred to the intensive care unit (ICU). As her hypoxemia worsened during ICU follow-ups, she was intubated on the third day of her hospitalization in the intensive care unit and followed up under mechanical ventilation and in the prone position. The patient, who developed septic shock and multi-organ failure during follow-up, died after 14 days in the ICU.

DISCUSSION

In this case report, we presented a patient who had been using omalizumab for five years with the diagnosis of severe asthma and died due to COVID-19 infection. To the best of our knowledge, this is the first Turkish case receiving omalizumab treatment due to severe asthma who died of COVID-19. It is thought that treatment with omalizumab, an anti-IgE antibody, may protect against severe COVID-19 infection. Omalizumab has been shown to reduce the high-affinity IgE receptors on plasmacytoid dendritic cells and increase the antiviral immune response (4). In the PROSE (Preventative Omalizumab or Step-up Therapy for Severe Fall Exacerbations) study, it was shown that omalizumab caused a decrease in the frequency of viral infections due to decreased expression of high-affinity IgE receptors in plasmacytoid dendritic cells in asthmatic children (5,6). There is limited data in the literature on COVID-19 patients receiving omalizumab. In a study by Dominguez-Ortega J et al., seven of 71 patients with severe asthma who took biological agents were diagnosed with COVID-19. Five of these seven patients were receiving omalizumab treatment. Pneumonia requiring hospitalization was diagnosed in only one of the seven patients receiving omalizumab, and the clinical findings were very mild in the other patients. During the COVID-19 infection, no asthma exacerbation or loss of asthma control was observed in these seven patients (7). In this case we presented, asthma control was impaired during COVID-19 while asthma was under control, and the infection was mortal.

Coronavirus infections other than COVID-19 limited to the upper respiratory tract have been associated with asthma exacerbations (8). The effect of more virulent coronaviruses such as SARS-CoV-2 on asthma control
is not well known. According to the current literature, asthma is not among the top 10 comorbid diseases in patients who die from COVID-19. Comorbid diseases such as diabetes, chronic heart disease, and obesity have been found to be more closely associated with COVID-19-related deaths (9). However, the Center for Disease Control and Prevention (CDC) reported that the most common comorbid diseases in young patients hospitalized with the diagnosis of COVID-19 are obesity, asthma, and diabetes (10). In this case, accompanying comorbid diseases were asthma, diabetes, FMF, and obesity. Concomitant diabetes and obesity may have contributed to the mortality of the case. The impact of asthma on COVID-19 severity remains uncertain. This uncertainty also causes anxiety in healthcare providers and patients.

The anxiety that asthma medications may adversely affect the clinical course of COVID-19 infection causes concern among healthcare providers. Inhaled steroids are the mainstay of asthma treatment and oral steroids are frequently used in asthma exacerbations. In *vitro* studies have shown that corticosteroids inhibit viral cytokines, but not interferons, which are an important antiviral defense mechanism (11). According to the current literature data, it is not possible for us to make a clear judgment about the effects of corticosteroid therapy on the prognosis of COVID-19 in patients with asthma. In addition, there is no data indicating that biological drugs approved for asthma treatment increase the risk of COVID-19. The GINA guidelines also recommend that asthmatic patients continue their current controller medication during the COVID-19 outbreak (3).

In conclusion, the effect of biological agents used in asthma treatment on the clinical course of COVID-19 is unknown. However, biological agents should be preferred over corticosteroids in severe asthma patients requiring regular use of systemic corticosteroids. We think that other comorbid diseases (obesity, diabetes) may have contributed to mortality in this patient with severe asthma who received omalizumab treatment.

**REFERENCES**


