COID-19 Pneumonia or Hypereosinophilic Syndrome?

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Abstract

Hypereosinophilic syndromes (HESs) are a group of disorders characterized by pathological proliferation of eosinophils. Diagnostic criteria include eosinophil count of 1,500/mm3 or higher, presence of organ involvement and exclusion of other causes of eosinophilia for at least 6 months. A 69-year-old male patient was referred to the pandemic clinic with a preliminary diagnosis of coronavirus disease 2019 (COVID-19) with fever and dyspnea. Computed tomography (CT) was compatible with COVID-19, nasopharyngeal swab polymerase chain reaction (PCR) was negative for two times. He had 20,000/mm3 eosinophilia and retrospective examinations showed that he have had eosinophilia for more than 1 year. Platelet-derived growth factor receptor alpha (PDGFRα) resulted positively. After starting parenteral methylprednisolone with a dose of 1 mg/kg, he was discharged with oral maintenance therapy. In outpatient follow-up, it was observed that eosinophilic pneumonia completely regressed. HES may occur with multiple system and organ involvement and findings. In the differential diagnosis of patients presenting with heart failure, pulmonary involvement and eosinophilia, HES must definitely be considered.

Keywords: COVID-19; Eosinophilia; Hypereosinophilic syndrome; Pulmonary involvement

Introduction

Hypereosinophilic syndromes (HESs) are a group of disorders characterized by excessive proliferation of eosinophils, with eosinophilic infiltration and mediator release. Although it has been considered idiopathic for a long time, etiologies have been described for some forms of HES. Percentage of eosinophilia has increased white blood cell count and eosinophils, and is between 350 - 500/mm3 [1]. Eosinophilia, which expresses the increase of eosinophils in peripheral blood, is examined by grading mild (500 - 1,500/mm3), moderate (1,500 - 5,000/mm3) and heavy (> 5,000/mm3) [1, 2]. The term “hypereosinophilia” is generally used for values above 1,500/µL [3].

In the data of “The Surveillance, Epidemiology and End Results (SEER)” between 2001 and 2005, the incidence of HES was determined as 0.036 per 100,000 [4]. Fatigue, cough, shortness of breath, myalgia, angioedema, rhinitis, rash and fever are the most common symptoms of hypereosinophilia. The most common clinical findings in the follow-up of patients with hypereosinophilia are due to dermatological involvement (69%). There then are pulmonary (44%) and gastrointestinal (38%) findings [5]. Eosinophilia-induced cardiomyopathy has been reported at 20% [6]. Progressive heart failure is the most typical example of eosinophilic organ damage and develops as a result of a multi-step pathophysiological process.

During the coronavirus disease 2019 (COVID-19) pandemic, which affects the whole world, the patient was referred to emergency clinic with cough, high fever, and pulmonary infiltration was detected in lung imaging. In our case, a patient who was referred to the pandemic service with suspicion of COVID-19 based on clinical and imaging results, but was diagnosed with HES through an advanced examination, is mentioned.

Case Report

A 69-year-old male patient with New York Heart Association stage IV (NYHA IV) heart failure and chronic obstructive pulmonary disease (COPD) was admitted to our hospital’s emergency clinic with fever, shortness of breath and cough. Polymerase chain reaction (PCR) was planned with thorax imaging and nasopharyngeal swab, considering COVID-19 in the foreground. In thorax computed tomography, patched ground glass consolidation areas were observed at the level of the middle and lower lobe segments of the lung (Fig. 1). In his examinations, hemoglobin was 12.3 g/dL, white blood cell count 56,410/mm3, neutrophil was 31,720/mm3, eosinophil was 20,000/mm3, platelet was 402,000/mm3, urea was 34 mg/dL, and creatinine 0.66 mg/dL. Lactate dehydrogenase (LDH) was 312 mg/dL and C-reactive protein (CRP) was 35 mg/dL. Retrospective examinations showed that he have had eosinophilia for more than 1 year. The peripheral smear evaluated for remarkable eosinophilia has increased white blood cell count and eosinophils, and toxic granulation is observed in neutrophils. In stool microscopy examined for secondary reasons, trophozoite, parasitic cyst and egg were not observed. Malignancy and additional pathology were not detected in the neck, thorax, and abdominal
computed tomography. The COVID-19 PCR sample was found to be negative twice. Platelet-derived growth factor receptor alpha (PDGFRα) mutation was found positive in genetic mutation screening. Bronchoalveolar lavage (BAL) was planned for pulmonary involvement of HES, but it could not be performed due to the risk of transmission in pandemic situation.

He was taking furosemide 80 mg/day and digoxin tablet 0.5 mg/day orally for congestive heart failure, and had also an implantable cardioverter defibrillator (ICD) for 2 years. Methylprednisolone with a dose of 1 mg/kg was added to the treatment of the patient who was evaluated as HES lung involvement. The hemogram after 5 days was as follows: hemoglobin: 9.5 g/dL, white blood cell: 7,430/mm³, neutrophil: 3,830/mm³, eosinophil: 180/mm³, platelet: 403,000/mm³. The patient was discharged with oral maintenance methylprednisolone treatment. In the outpatient follow-up, 3 weeks after beginning methylprednisolone, it was observed through computed tomography that the signs of eosinophilic pneumonia completely regressed (Fig. 2), and his complaints entirely disappeared. While the steroid dose was planned to be reduced, it was planned to continue with imatinib mesylate 100 mg daily.

Discusion

Lung involvement can be seen in 40-60% of cases in HES. The most common respiratory symptom is the chronic and persistent cough. Lung involvement results in sputum, wheezing, and shortness of breath, which increases the suspicion of bronchial hyperreactivity. As with our patient, patients may be mistaken for asthma or COPD. However, pulmonary function tests typically reveal that there is no obstructive pulmonary disease [8]. In these patients, BAL can be performed and the presence of a large number of eosinophils can be demonstrated. BAL sampling could not be done in our patient due to the risks in the pandemic situation.

When cardiac involvement occurs, the most common form is endomyocardial fibrosis. Progressive scarring, especially in the posterior mitral valve, mostly limits the movement of the valve. Although cardiac involvement is best diagnosed by endomyocardial biopsy, this procedure carries high morbidity and mortality risks. Instead, a noninvasive procedure, such as transesophageal echocardiography, is valuable for assessment and detection of possible thrombus formation. Chronic heart failure and severe arrhythmia are frequently found in the late stages of the HES, as in our case [7, 9, 10]. Neuropsychiatric symptoms are common in HES due to the involvement of the central nervous system [9]. It often presents with loss of consciousness, depressed mood, poor coordination, and peripheral neuropathy. There was no evidence of neuropsychiatric involvement in the hospital application and anamnesis in our patient.

Laboratory evaluation of primary eosinophilia should be initiated by screening factor interacting with PAPOLA and CPSF1 (FIP1L1)-PDGFRα gene fusion in peripheral blood. The presence of a deletion in 4q12 can be detected, leading to
the formation of FIP1L1-PDGFRA fusion [9].

Treatment should begin in HES, which includes organ damage by definition. The main treatment is corticosteroids; it is usually started at a dose of 1 mg/kg, after 1 - 2 weeks of use at this dose, it is gradually reduced over a period of 2 - 3 months [3]. Imatinib is the first drug to be considered in the presence of translocations with other genes and changes in FIP1L1-PDGFRA or PDGFRβ. Tyrosine kinase inhibitor therapy results in complete clinical and molecular remission in a high proportion of patients and complications associated with the disease, including transformation into leukemia, are prevented [3]. However, a small number of patients may require treatment changes and additional treatment.

Conclusions

HES can occur with nonspecific multiple cutaneous, immunological, pulmonary, cardiac, ocular, rheumatological and gastrointestinal manifestations. Physicians should definitely consider HES in the differential diagnosis of patients presenting with heart failure, pulmonary involvement and eosinophilia. As in our case, although other comorbidities, such as COPD, may prevent a correct diagnosis of HES with pulmonary involvement, early diagnosis usually provides the most appropriate treatment.

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Conflict of Interest

None to declare.

Informed Consent

An informed consent was obtained from our patients to publish this study.

Author Contributions

All authors contributed to the editing of the manuscript. IS wrote the manuscript and created samples from computed tomography images.

Data Availability

The authors declare that data supporting the findings of this study are available within the article.

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