Pneumonia and seizures due to hypereosinophilic syndrome—organ damage and eosinophilia without synchronisation: A case report

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Abstract

BACKGROUND

Hypereosinophilic syndrome (HES) is a condition characterized by increased eosinophil proliferation in the bone marrow, as well as tissue eosinophilia, often causing organ damage. The cause of the disease is unknown. Initial symptoms include fatigue, cough, shortness of breath, myalgia, angioedema, fever, and pneumonia. In addition to the respiratory symptoms, damage to the central nervous system can lead to severe seizures. Here, we report a case with pneumonia and complex partial seizures secondary to HES.
CONCLUSION

HES symptoms are variable and atypical, and the level and timing of eosinophilia and organ damage are often discordant.

Key Words: Case report; Hypereosinophilic syndrome; Pneumonia; Seizures; Prednisolone; Levetiracetam

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Core Tip: Although there are diagnostic criteria for hypereosinophilic syndrome, the various degrees of organ damage and hypereosinophilia often make diagnosis difficult in clinical practice. In addition, the organ damage and blood changes do not always occur concurrently. Therefore, clinicians must consider many differential diagnoses, especially when patients present with atypical symptoms and disease course. Early initiation of treatment is no less important than an accurate diagnosis, and the balance between the two should be considered according to the patient's condition as well as the level and quality of medical resources available at the hospital.

INTRODUCTION

Hypereosinophilic syndrome (HES) is a marked blood and tissue eosinophilia of unknown aetiology with a variety of clinical manifestations. Since 1975, the disease has been defined by three criteria: (1) Blood eosinophilia ≥ 1500/mm$^3$ for longer than 6 mo (or death before 6 mo associated with signs and symptoms of hypereosinophilic disease); (2) Lack of evidence for parasitic, allergic, or other known causes of eosinophilia; and (3) Presumptive signs of organ involvement, such as heart failure, gastrointestinal dysfunction, central nervous system abnormalities, fever, or weight loss[1].

However, there are several problems with these criteria. First, according to these criteria, clinicians must wait 6 mo to diagnose a patient with multiple organ involvement. Second, an increase in eosinophils does not necessarily correlate with organ damage. Some patients may have a marked increase in eosinophils but only mild symptoms, while others may have a mild increase in eosinophils but significant organ damage[2]. Therefore, two diagnostic criteria have now been proposed to replace the classic three: (1) Blood eosinophilia of greater than 1500/mm$^3$ on at least two occasions or evidence of prominent tissue eosinophilia associated with symptoms and marked blood eosinophilia; and (2) Exclusion of secondary causes of eosinophilia, such as parasitic or viral infections, allergic diseases, drug-induced or chemical-induced eosinophilia, hypoadrenalism, and neoplasia[3]. However, this revision of the diagnostic criteria does not solve all the problems in the diagnosis and treatment of HES. We anticipate that this case will aid in the diagnosis and treatment of similar cases.

CASE PRESENTATION

Chief complaints

A 94-year-old Asian woman presented to our hospital for dyspnoea and wet cough. She also had abdominal pain and bloody stools. She was admitted to our hospital with a diagnosis of heart failure and sigmoid colon ulceration (day 1).

History of present illness

Her respiratory distress started during the night 1 d before presentation. It improved after 1 h and was still mild over the next morning and evening hours. However, 22 h later, her symptom got worse again during the night.

History of past illness

The patient had no chronic illnesses such as hypertension, diabetes, or asthma, and no history of cancer. She was a non-smoker and did not habitually drink alcohol. She had a well-balanced diet and lived a healthy lifestyle.
Personal and family history
There was no specific history.

Physical examination
The patient was 148 cm tall, weighed 42 kg, and her vital signs were as follows: blood pressure 111/43 mmHg; pulse 90/min; respiratory rate 14/min; and SpO₂ 98% (room air). On chest auscultation, her heart rhythm was regular and no heart murmur was found. However, auscultation of respiratory sounds found wheezing. Her abdomen was flat and soft and abdominal auscultation found neither increased nor decreased intestinal peristalsis. There was no rebound tenderness or abdominal guarding. However, there was intermittent and spontaneous abdominal pain and bloody stools. Her eyelid conjunctiva did not show jaundice or pallor. Her oral and nasal cavities and skin surfaces showed no abnormal findings. Her upper limbs showed no abnormalities. However, her lower extremities showed indurated oedema.

Laboratory examinations
On day 1, her rapid antigen tests using a throat swab were negative for influenza A and B and COVID-19. Her electrocardiogram (ECG) showed no abnormalities on admission and on loss of consciousness on day 108. In her blood analyses, there were no abnormalities in the electrolyte, glucose, lipid, liver function, and renal function parameters. The antinuclear antibody (ANA) titre at 1:40 serum dilution was positive but staining patterns at 1:40 serum dilution was negative for homogeneous, discrete speckled, speckled, nucleolar, and peripheral staining. Anti-neutrophil cytoplasmic antibody (ANCA) and HIV tests were negative. On day 32, the patient’s IgE level was slightly high at 237 IU/mL (normal: 27.54–138.34), but her IgG, IgA, and IgM levels were normal. Blood analyses also showed that eosinophils and brain natriuretic peptide (BNP) levels were high. The highest values for each were 1760/mm³ on day 26 for eosinophils and 738.1 pg/mL on day 1 for BNP (Figures 1 and 2). The blood and sputum taken on day 26 were negative on culture for bacteria or fungi and no parasites were found. However, on day 109, blood culture was positive for methicillin-resistant Staphylococcus epidermidis, and the β-D-glucan level was high (316.0 pg/mL). On day 128, blood tests showed normal levels of ammonia. The patient’s post-consciousness EEG showed spikes and waves in Fp1, F1, C3, P3, and O1 (Figure 3). Her echocardiogram showed no thrombi in the atra or ventricles.

Imaging examinations
Non-contrast computed tomography imaging of the patient’s chest showed bilateral pleural effusions and infiltrative shadows (Figure 4). Her gastrointestinal endoscopy showed multiple ulcers in the sigmoid colon (Figure 5). Histopathological examination showed colonic gland prolapse, vitrification of the stroma, and an infiltration of inflammatory cells, but no evidence of malignant transformation. Non-contrast magnetic resonance imaging of the patient’s head showed extensive periventricular hyperintensity (Figure 6).

MULTIDISCIPLINARY EXPERT CONSULTATION
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The patient’s eosinophils in the blood tests exceeded 1500/mm³ only once, but she had high eosinophils multiple times without meeting this threshold. Additionally, it was clear that the eosinophilic infiltrate was causing serious damage to the nervous, respiratory, cardiovascular, and digestive systems. We should not have delayed the initiation of her treatment any longer to meet the classical diagnostic criteria. Biopsies of organs other than her sigmoid colon were considered for a more precise examination and imatinib initiation was considered for more effective treatment, but both were too invasive for the elderly patient. Given that she and her family did not wish to undergo these tests and treatments, we did not pursue these options. Although Japan is a country with some of the richest medical resources in the world, the hospital where the patient was admitted had no intensive care unit or specialists in collagen diseases and autoimmune diseases, and the same is true for other similar rural hospitals. The use of limited medical resources is an important issue in this country as in the rest of the world.

FINAL DIAGNOSIS
The patient was diagnosed with epileptic seizures and pneumonia caused by HES.
Figure 1 The patient’s serial blood test results showing changes in total white blood cells and eosinophil count. The eosinophil count was automatically calculated from the ratio of eosinophils to total white blood cells. When the ratio was less than 0.1%, it was counted as 0. WBC: White blood cells; PSL: Prednisolone.

Figure 2 Serial changes in the patient’s brain natriuretic peptide values. BNP: Brain natriuretic peptide.

**TREATMENT**

In addition to rehabilitation, the patient was treated with furosemide for heart failure, ceftazidime and vancomycin and fungard and PSL for pneumonia and levetiracetam (LEV) for seizures.

**OUTCOME AND FOLLOW-UP**

Furosemide 20 mg/d for 10 d from the day of admission (day 1) relieved her congestive heart failure symptoms for a period of time. A diet suitable for digestion and absorption made her bloody stools and abdominal pain disappear. Rehabilitation, including gait training and flexibility exercises to prevent loss of range of motion and contractures throughout the body, was also initiated. However, from day 56, her congestive heart failure symptoms recurred, and pneumonia also appeared. Treatment with 20 mg/d furosemide was restarted from that day, but without effect. Ceftazidime 2 g/d was also ineffective in treating the pneumonia. On day 68, the patient was diagnosed with HES and treatment with prednisolone (PSL) 25 mg/d was started. After that, the pneumonia and heart failure symptoms gradually
Ishida T et al. Pneumonia and seizures due to HES

Figure 3 The patient's post loss of consciousness electroencephalogram showing a slow wave indicative of a clinical epileptic seizure.

Figure 4 Non-contrast computed tomography image of the patient's chest showing bilateral pleural effusions and infiltrative shadows.

improved, so the PSL was gradually reduced to 15 mg/d. Her rehabilitation was also resumed. However, on day 108, her eyes suddenly rolled upwards, and she showed eyelid twiching and loss of consciousness. Her breathing stopped and her $\text{SpO}_2$ dropped to 70% (room air). Oxygen therapy (reservoir mask: 10 L/min) was started immediately. Diazepam 10 mg intramuscular injection was also given and after 5 minutes her $\text{SpO}_2$ improved to 90% (reservoir mask: 10 L/min) and her seizures stopped. LEV 1000 mg/d was started and subsequently, the seizures did not recur. However, her state of consciousness continued to be unfavourable on the Glasgow Coma Scale, E2 V2 M4.

Her respiratory status also improved reaching $\text{SpO}_2$ 90% (nasal cannula: $\text{O}_2$ 1L), but not higher. On day 109, fungard 50 mg/d, vancomycin 1 g/d and ceftazidime 2 g/d were started and her PSL was increased to 30 mg/d. However, her pneumonia continued to worsen, and she died of pneumonia on day 135.

DISCUSSION

First, we will discuss diagnosis of HES. As mentioned at the beginning of this report, the diagnostic requirement for HES is a blood eosinophil count $\geq 1500/\text{mm}^3$ for at least 6 mo according to the classical diagnostic criteria or measured at least twice according to the diagnostic criteria proposed by Simon et al [3]. In this case, neither of these criteria were met at the time the diagnosis was made. In our case, the patient’s blood eosinophil count exceeded 1500/\text{mm}^3 only once and this did not necessarily coincide with the most severe period of organ damage. Some clinicians suggested that we should have waited.
until her eosinophil count had risen again and her symptoms had worsened before deciding on a diagnosis. However, as stated in the Multidisciplinary Expert Consultation, we needed to start her treatment as soon as possible. This was considered a higher priority than meeting the strict diagnostic criteria. In addition, the disease course in this case meets the criterion of “or death before 6 mo associated with signs and symptoms of hypereosinophilic disease” listed in the supplementary information in the classical criteria\[3\]. Heart failure, sigmoid colon ulceration, pneumonia, and epilepsy signified damage to each of these organs in our patient. Therefore, the diagnosis of HES in this case was appropriate. With regard to treatment, the focus is on PSL. As is well known, PSL is a drug that needs to be tapered off, not stopped abruptly. Furthermore, a longer PSL treatment course is commonly required for eosinophilic pneumonia. However, in this case, given the early decline in the eosinophil count after initiation of PSL therapy, perhaps it should have been tapered or stopped earlier.

Next, we would like to discuss the diagnosis and treatment of epilepsy. Common causes of loss of consciousness in older people include heart disease, autonomic disorders, other conditions such as anaemia, ischaemia and varicose veins, or anticholinergic drugs\[5\]. Our patient had congestive heart failure at the time of admission, but her condition improved. She did not have any blood clots identified that could have caused cerebral embolism. Her vital signs, including blood pressure and ECG, were normal. Therefore, it was considered unlikely that the loss of consciousness was due to circulatory problems. Her post-consciousness electroencephalogram (EEG) suggested that there were epileptic discharges around the left anterior frontal lobe, motor cortex, central sulcus, sensory cortex, and visual cortex. The rather unremarkable EEG findings compared with the clinical findings may be because this EEG was done after the LEV treatment was started. The epileptic discharges may have been more intense before the administration of antiepileptic drugs. We were not able to do rapid or continuous monitoring EEG at our hospital. In diagnosing and treating her epilepsy, we were forced to prioritise life-saving treatment over rigorous diagnosis.
Regarding the seizure type, her epilepsy was most likely a complex partial seizure. Complex partial seizures are more likely to occur in older adults and cause disorientation, but they do not cause tonic or clonic seizures[6]. The fact that LEV treatment was effective in preventing the seizure symptoms was also one of the reasons for the diagnosis. However, the presence of eyelid twitching also suggests that her seizure type may have been absence seizures. Absence seizures are generally more common in young children and have a shorter duration of disorientation. However, if the seizure is an atypical absence seizure, the disturbance of consciousness may be prolonged. In Japan, LEV may be administered at doses of up to 3000 mg/d. However, in this case, the patient was elderly, and it was feared that increasing the dose of LEV might cause side effects such as malignant syndrome. The LEV dose was not increased because there were no obvious epileptic seizures after the start of LEV at a dose of 1000 mg/d. In this case, the patient’s loss of consciousness was prolonged even after an intramuscular injection of 10 mg diazepam and was, therefore, considered to be benzodiazepine-refractory. LEV, fosphenytoin, and valproic acid are suitable for the treatment of epilepsy which does not improve with benzodiazepines. In elderly patients, there is no difference in the safety or therapeutic efficacy between these three drugs[7]. LEV was chosen in this case because it can be started intravenously and can be seamlessly transferred to the oral route if oral medication becomes available later. We considered hepatic encephalopathy as a possible cause of our patient’s impaired consciousness, but this was ruled out by the normal blood ammonia levels and the fact that the EEG showed no triphasic waves.

While all ANA staining patterns were negative, the ANA titre was positive at 1:40 dilution. This result indicates that our patient may have had a collagen disease. Therefore, we needed to differentiate the collagen diseases with eosinophilia granulomatosis with polyangiitis (EPGA) and polyarteritis nodosa (PAN), from HES. EPGA, formerly known as Churg–Strauss syndrome, is characterised by asthma, eosinophilia of the blood and tissues, and small vessel vasculitis. The clinical symptoms are variable but can be divided into two main subtypes: The "vasculitis" type, which is positive for ANCA and shows glomerulonephritis, purpura, and inflammation of multiple peripheral nerves. The other is the "eosinophilic" type, which is negative for ANCA but is characterised by a markedly high level of eosinophils and damage in the lungs and myocardium[8]. This case is similar to the "eosinophilic" type, but EPGA is less likely to cause central nervous system damage such as seizures.

PAN is a vasculitis that targets medium-sized arteries and leads to multi-organ failure. The target organs include the heart and the gastrointestinal tract. We note that damage to the central nervous system has also been reported[9]. However, damage to the lungs has rarely been reported[10]. The present case is not typical but is similar to both EPGA and PAN. However, the pathological findings in the sigmoid colon did not show the features of either of these diseases.

In terms of brain imaging findings, the differential diagnosis in this case also includes cerebral amyloid angiopathy (CAA). The Modified Boston Criteria for CAA are based on imaging and pathological findings. The disease can be classified as: (1) Definite CAA; (2) Probable CAA with supporting pathology; (3) Probable CAA; and (4) Possible CAA. The novelty of this criteria is that (3) and (4) do not require a pathological examination[11]. In this case, at the request of the patient and her family, no pathological examination of the brain tissue was performed after her death. Generally, the hallmark of CAA on brain imaging is multiple sporadic lesions confined to the cerebral cortex, cortico-subcortical junction grey matter, and subcortical white matter. However, some subtypes of CAA, such as cerebral amyloid angiopathy-related inflammation (CAA-ri), present with mass-like lesions as in this case[12]. CAA-ri is known to respond well to treatment with steroids and immunosuppressive drugs. In this case, treatment was given with the expectation that the patient might be diagnosed with CAA-ri. There is a report that CAA-ri can cause seizures[13], which seems to be consistent with the symptoms in this case. However, there are no previous studies on its frequency. Similar cases need to be studied in the future.

CONCLUSION

HES can cause damage to many organs and has an undulating disease course. Therefore, HES must be differentiated from other diseases such as EPGA, PAN, and CAA. In this case, the diagnosis was more difficult because of the time lag between the various clinical symptoms and the eosinophilia. The focus of treatment was to determine the appropriate dose and duration of PSL and LEV therapy. We conclude that this case report can be used as a reference for the diagnosis and treatment of similar cases.

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FOOTNOTES

Author contributions: Ishida T was the patient’s primary care physician and contributed to the literature review and manuscript preparation; Murayama T and Kobayashi S reviewed the literature and contributed to the preparation of the manuscript; and All authors gave their approval for the final version to be submitted.

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