Loeffler endocarditis as a rare cause of heart failure with preserved ejection fraction
A case report and review of literature

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Abstract
Rationale: Hypereosinophilic syndrome (HES) is a rare disease characterized by hypereosinophilia and its ensuing organ damage. Cardiac involvement is divided into 3 chronological stages: an acute necrotic stage; a thrombus formation stage; and a fibrotic stage. Infiltration of the myocardium by eosinophilic cells followed by endomyocardial fibrosis is known as “Loeffler endocarditis.”

Patient concerns: We report a case of a 60-year-old man diagnosed with left-sided restrictive cardiomyopathy.

Diagnosis: The patient experienced heart failure with preserved ejection fraction. The cardiac MRI showed intense, linear, delayed gadolinium enhancement of the endocardium of the lateral wall of the left ventricle, and obliteration of the LV apex. He was ultimately identified as Loeffler endocarditis.

Intervention: A bone marrow smear and biopsy revealed the FIP1L1-PDGFRA fusion gene was positive in 82% of segmented nucleated cells.

Outcome: Our patient responded well to prednisone at 1 mg/kg/d.

Lessons: HES is a rare disease that often affects the heart. Cardiac involvement in hypereosinophilia, especially Loeffler endocarditis, carries a poor prognosis and significant mortality. Early detection and treatment of the disease is therefore essential. Further studies are needed to ascertain therapeutic corticosteroid dosages and develop targeted gene therapies, both important steps to ameliorate the effects of Loeffler endocarditis and improve patient outcomes.

Abbreviations: CS = corticosteroids, ECG = electrocardiography, FISH = fluorescence in situ hybridisation, HES = hypereosinophilic syndrome, LV = left ventricle, MRI = magnetic resonance imaging.

Keywords: heart involvement, HES, hypereosinophilic syndrome, Löffler endocarditis

1. Introduction

Hypereosinophilia is defined by the presence of $\geq 1500$/mm$^3$ eosinophils in the peripheral blood, and may be reactive, neoplastic, or idiopathic.$^{[1,2]}$ A marked and persistent overproduction of eosinophils that subsequently infiltrate and damage multiple organs via a toxic protein is referred to as hypereosinophilic syndrome (HES). The age-adjusted incident rate of HES is approximately $0.036$ per 100,000 person-years.$^{[3]}$ The clinical manifestations of HES are variable and depend on which organ is targeted by the proliferative eosinophils. Cardiac manifestations occur in half of patients with HES and are a major cause of morbidity and mortality among HES patients. Infiltration of the myocardium by eosinophilic cells following endomyocardial fibrosis is known as “Loeffler endocarditis.”$^{[4]}

In the present study, we report a case of a 60-year-old man diagnosed with left-sided restrictive cardiomyopathy with a preserved ejection fraction, which was identified as Loeffler endocarditis.

2. Case report

A 60-year-old male with an unremarkable prior medical history presented to our institution with progressively worsening dyspnea on exertion and paroxysmal nocturnal dyspnea for a duration of 20 days. The physical examination was within normal limits, apart from a finding of bilateral crackles of the lung bases. His complete blood count at presentation showed a normal white blood cell count ($7280$/mL), with $30\%$ neutrophils, $13\%$ lymphocytes, and $5\%$ eosinophils, but demonstrated hypereosinophilia (absolute eosinophil counts $520$/mL), with severely elevated pro-B-type natriuretic peptides of $3710$pg/mL. Echocardiography revealed an apical left ventricular infiltration with a $14 \times 7$ mm thrombus that was associated with leaflet restriction and moderate mitral and tricuspid valve regurgitation (Fig. 1). Both atria were enlarged and a restrictive left ventricle (LV) filling pattern was noted. Both ventricles displayed normal dimensions in the absence of regional

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wall motion abnormalities, and both ventricular systolic functions were preserved (left ventricular ejection fraction 64%). The estimated systolic pulmonary artery pressure was 71 mm Hg. Electrocardiography (ECG) showed a right axis deviation and T waves changes. Cardiac magnetic resonance imaging (MRI) was performed to aid in the diagnosis. The cardiac MRI showed intense, linear, delayed gadolinium enhancement of the endocardium of the lateral wall of the LV, and obliteration of the LV apex (Fig. 2). A bone marrow smear and biopsy revealed a markedly elevated eosinophil count (>36%), but was absent dysplasia and a number of blasts, and there was no evidence of myeloproliferative disorder. The FIP1L1-PDGFRA fusion gene was positive in 82% of segmented nucleated cells, based on fluorescence in situ hybridization (FISH) analysis (Fig. 3). An additional, extensive workup of secondary causes of eosinophilia proved negative and excluded other causes, leading to the final diagnosis of idiopathic HES with heart involvement. Treatment was started immediately with warfarin and a high dose of prednisone (1 mg/kg/d). Over the next few months, the patient began to feel much better, demonstrating marked improvement in his symptoms, with no complaints of shortness of breath or dyspnea on exertion. Eosinophil counts also normalized.

The study protocol was approved by the ethics review board of the First Hospital of Jilin University (No. 2016–263). Informed consent was obtained from the patient for his information used for research.

3. Discussion

Eosinophils are granulocytes that play a role in the immune response to inflammation and infection. In healthy subjects, the
upper normal limit of eosinophils in the peripheral blood is 3–5% (absolute eosinophil count of 350–500/mm³). Extensive infiltration of eosinophils in tissue results in organ damage through associated tissue fibrosis via the secretion of TGF-β and IL-1β. A chronic state of eosinophil proliferation and infiltration can lead to a diagnosis of HES.

HES is defined by 1 or more of the following:

- an elevated eosinophil count (>1500/mm³) on 2 separate tests (≥1 mo);
- tissue hypereosinophilia based on >20% eosinophils in a bone marrow section;
- marked disposition of eosinophilic granule proteins in tissue plus organ damage directly due to hypereosinophilia.

Our patient presented with symptoms of heart failure, and demonstrated a moderately elevated eosinophil count. The diagnosis of HES with cardiac disorder was made by cardiac MRI, the current non-invasive gold standard for diagnosis. The underlying causes of HES are various and were classified in 2011 at the Working Conference on Eosinophil Disorders and Syndromes into following terms: hereditary (familial) HE variant; primary (clonal/neoplastic) HE variant; secondary (reactive) HE; and HE of undetermined significance/idopathic variant.

Our patient received a complete workup to find the underlying etiologies, and the negative results led to the diagnosis of idiopathic HES.

HES is a potentially fatal disease with cardiac involvement in about 40% to 50% of all cases. Cardiac manifestations, particularly eosinophilic myocarditis and endomyocardial fibrosis, are typical causes of mortality in HES. Eosinophilia cardiac disease was first documented in 1936 by Loffler, who described 2 cases of “endocarditis parietalis fibrosistica,” a condition consistent with the endomyocardial fibrosis seen in HES at later stages of cardiac disease. Cardiac involvement usually follows 3 stages. The first stage, frequently asymptomatic, begins with acute necrosis and eosinophilic infiltration and toxic degranulation. The second (thrombotic) stage is characterized by formation of mural thrombi and thrombus. Both ventricles can be involved, but the thrombus most often occurs at the apex of the LV. The last (fibrotic) stage occurs when the granulation tissue is replaced by fibrosis and restrictive cardiomyopathy ensues. Severe valvular disease is a common complication during this stage, and patients present with clinical restrictive features but preserved systolic function. It is important to note that the stages may overlap; the degree of cardiac dysfunction does not necessarily correlate with the degree of eosinophilia.

Histological evidence is necessary for the diagnosis of HES; however, the invasive endocardial biopsy procedure necessary to obtain sample is difficult to implement clinically. It may be more valuable in diagnosing early cardiac involvement, but the acquisition of an adequate biopsy sample is difficult, especially in the presence of endocardial fibrosis and superimposed thrombi. By contrast, echocardiography is a very useful first-line method to detect impairment of cardiac and valvular function, ant to visualize thrombus and pericardial effusions and their dynamics. Echocardiography should be used in cases of moderate (1500/mm³) to severe eosinophilia (5000/mm³), whereas the initial necrotic stage may be undetectable by echocardiography. Cardiac MRI is currently the gold standard for non-invasive diagnosis, monitoring, and prognostic stratification of myocarditis. It can easily detect endocardial fibrosis with subendocardial late gadolinium enhancement before functional consequences, and reduce the need for invasive biopsy in HES patients. Genetic testing for Fip1-like1-platelet-derived growth factor receptor (alpha) fusion gene (FIP1L1-PDGFRα) has recently been reported in a few cases of HES and has been added to the diagnostic algorithm of HES. The FP fusion gene can activate tyrosine kinase activity, stimulate proliferation, and mediate survival of eosinophils in HES. The prevalence of FIP1L1-PDGFRα fusion is 10% or less in idiopathic HES. Presence of the mutation appears to correlate with more common cardiac involvement. The tyrosine kinase inhibitor imatinib is a specific inhibitor of the PDGFR alpha receptor and has been a clear target in the development of treatments for the FLP1L1-PDGFRα mutation. Imatinib can achieve complete hematological and molecular remission within weeks to months, but cannot eliminate the FLP1L1-PDGFRα clone in most patients.

Therapeutic interventions for HES typically proceed in a stepwise fashion, beginning with immunosuppressive treatment corticosteroids (CS) to reduce eosinophil count and counteract inflammation. There is little consensus regarding the initial dosage of CS or the treatment duration. In a multicenter study of 188 subjects with HES, 179 of whom had the FLP1L1-PDGFRα mutation and were treated with CS, 65% were non-responders. Among the responders, there are no clear evidence-based recommendations that can be given. It seems reasonable to adjust the dosages of CS and the treatment duration with respect to the severity of EM manifestation. Recommended starting doses are typically 0.5 to 1mg prednisone/kg body weight. The prednisone dose required to maintain disease control is highly variable from 1 patient to another overtime. Our patient, who tested positive for the FLP1L1-PDGFRα mutation responded well to prednisone at 1mg/kg/d.

4. Conclusion

HES is a rare disease that often afflicts the heart. Cardiac involvement in hypereosinophilia, especially Loffler endocarditis, carries a poor prognosis and significant mortality. Early detection and treatment of the disease is therefore essential. Further studies are needed to ascertain therapeutic corticosteroid dosages and develop targeted gene therapies, both important steps to ameliorate the effects of Loffler endocarditis and improve patient outcomes.

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