INTRODUCTION

Hypereosinophilic syndrome (HES) is a rare disease characterized by persistent eosinophilia (>1,500 eosinophils × 10⁶/L) for at least 6 months that remains unexplained in spite of a comprehensive evaluations and evidence of organ dysfunction primarily attributed to the eosinophilia. Loeffler's endocarditis is a myocardial disease, thought to be secondary to idiopathic hypereosinophilia and often secondary to infection, medication or clonal disorders such as acute leukemia. It is characterized by fibrous thickening of the endocardium of one or both ventricles, leading to apical obliteration and restrictive cardiomyopathy, resulting in heart failure, thromboembolic event and/or atrial fibrillation. Prognosis is believed to be poor due to high mortality from heart failure, sudden death or thromboembolism. The reported 10-year survival rate is less than 50%, which is especially lower in corticosteroid resistant cases with cardiac involvement.

Here, we present a patient who presented with anasarca, peripheral hypereosinophilia, and typical echocardiographic findings consistent with Loeffler's endocarditis, which disappeared with corticosteroid and anticoagulation therapy.

CASE

A 41-year-old male visited our hospital for recently progressed generalized edema and abdominal distension. He had a history of persistent malaise and fatigue for 3 months. Prior to the this first visit, he had been admitted for ascites control at the other hospital for several days, but he did not have a favorable response to conventional diuretic therapy. On his visit, vital signs were stable; blood pressure, 120/80 mmHg; pulse rate, 78 bpm; respiratory rate, 18/min and body temperature, 36.5°C. Physical examination revealed jugular venous distension in the sitting position, decreased breaths sound in the bilateral lower lung fields, severely distended abdomen with shifting dullness on percussion and pretibial pitting edema. Chest X-ray revealed cardiomegaly and blunting of bilateral costophrenic angles. Electrocardiography demonstrated normal sinus rhythm with low voltage in the limb leads. Laboratory studies showed marked eosinophilia (5,060 × 10⁶/L), mild elevated cardiac enzyme (troponin-I 0.57 mg/mL), mild elevated liver enzyme (total bilirubin 2.5 mg/dL, alkaline phosphatase 144 U/L) and elevated C-reactive protein (2.43 mg/dL). But serum electrolyte level and thyroid functions were...
within normal limits. No parasite or ova were seen on stool examination. A bone marrow biopsy showed normocellularularity with eosinophilic hyperproliferation, and a fluorescence in situ hybridization test for FIP1L1-PDGFRa tyrosine kinase also demonstrated no evidence of abnormal myeloid maturation or lymphoproliferative disorder.

Two-dimensional (2D) echocardiography revealed preserved left ventricular ejection fraction (>55%) with reduced systolic tissue Doppler velocity (TDV) of septal mitral annulus; Sm, 4 cm/sec and diastolic TDV of septal mitral annulus; Em, 4 cm/sec, and also showed finding of diastolic dysfunction; E/E’, 15.4, E/A, 0.8. In color Doppler imaging, there was no significant tricuspid regurgitation, but the inferior vena cava was distended independent of respiration. However, a small amount of pericardial effusion and both ventricular apical obliteration by homogeneous echogenic materials suggestive of thrombus extending mid lateral and into the septal area were noted; both atria, particularly the right atrium, were markedly distended in contrast to the normal to small sized ventricles (Fig. 1A and B). Chest computerized tomography was performed to rule out pulmonary thromboembolism and lymphoma, and it revealed no specific mass or lymph node enlargement without an evidence of thromboembolism. However apical obliteration of both ventricles by non-enhanced homogenous materials and presence of same material in right atrium were noted (Fig. 2).

The patient was treated with oral corticosteroid (prednisolone 30 mg) and anticoagulation by intravenous infusion of unfractionated heparin, which was switched to warfarin on the 3rd hospital day. Also intravenous furosemide (20-40 mg/day) was administered to control ascites and edema. On the 2nd day of treatment with corticosteroid, peripheral eosinophilic counts decreased to a normal range (186 × 10^6/L) and peripheral edema was markedly improved. After a week of corticosteroid treatment, we attempted steroid tapering to 10 mg of prednisolone but because of fluctuations in eosinophilic counts, the initial dose was restarted and maintained.

Follow up 2D echocardiography on the 15th hospital day showed marked resolution of thrombus at both ventricular apices (Fig. 1C).

However, tissue Doppler velocity (TDV) of septal mitral annulus had no interval change compared to the initial tissue Doppler study. Additional follow up echocardiography at 2 months later did not show further resolution of the diastolic dysfunction. The patient was continued on corticosteroid and oral anticoagulation therapy.

**DISCUSSION**

In this report, we present a case consistent with Loeffler’s endocarditis secondary to idiopathic HES. In these patient group, heart is the most frequently involved organ (50-60% in cases). Loeffler’s endocarditis is thought to be a late cardiac manifestation that is characterized by fibrous thickening of the endocardium and extensive eosinophilic
infiltration of the ventricular endocardium, with subsequent
degranulation and necrosis with resultant endomyocardial
fibrosis, leading to apical thrombotic obliteration. In
addition, the intense inflammatory changes result in
superimposed thrombus formation, which subsequently
occupies much of the ventricular cavity. The presence of
asymptomatic endomyocardial lesions may precede overt
cardiac disease by a substantial period of time. Valvular
involvement, as well as endocardial involvement, includes
tricuspid, mitral, and aortic valve regurgitation caused by
endocardial inflammation, and fibrosis affecting papillary
muscles and chordae tendineae producing papillary muscle
dysfunction.

For the treatment of symptomatic patients with HES,
such as in the case of Loeffler’s endocarditis, the first line
drug of choice is corticosteroid, followed by cytotoxic
agents such as hydroxyurea or immunomodulatory agents
such as interferon-alpha. Gene rearrangements that
activate tyrosine kinase may cause a myeloid disorder,
which can be treated with imatinib mesylate. Glucocorticoid
treatment has resulted in clinical and biopsy-proven
improvement of eosinophilic infiltration and myocardial
damage as well as normalization of peripheral hypereosino-
philic. Thus, normalizing the peripheral eosinophilic count
has been accepted as one of the initial therapeutic targets.
Treatment with glucocorticoid (prednisolone 40-60 mg/day)
has been used, and along with conventional treatment of
heart failure and maintenance therapy with a lower dose of
10 mg/day, appears to prevent progression of cardiac
disease.

The vast majorities of the literature emphasize the
importance of oral anticoagulation because infiltration and
degranulation of eosinophils in the endocardium induce an
inflammatory and prothrombotic state that frequently
results in the development of a ventricular mural thrombus.
Consequently, anticoagulation therapy should be also
considered before the ventricular obliteration develops.
These findings were supported by those of Lofiego et al.
who reported five patients with Loeffler’s endocarditis,
including three patients with long-term echocardiographic
follow-up. This report showed that appropriate anticoag-
ulation therapy with a corticosteroid or a cytotoxic agent
(in the case of hypereosinophilic syndrome) not only prevented
pulmonary or systemic embolic events and thrombus
growth but also limited the obliteration of the ventricular
cavity, thereby inducing a favorable long-term left
ventricular remodeling. Endocardial biopsy is
recommended as an invasive diagnostic tool. However,
this case showed definite peripheral hypereosinophilia and
typical echocardiographic findings, therefore another
invasive diagnostic tool would not be needed to make a
diagnosis. To facilitate the choice of appropriate treatment
regimens, studies to determine possible secondary causes of
hypereosinophilia are also important. For instance, parasitic
or other infections, and clonal disorders such as acute
leukemia and lymphoma are one of the causes of secondary
hypereosinophilia. In this case, possible causes of
secondary HES were ruled out, and then we finally could
confirm Loeffler’s endocarditis due to idiopathic HES.
Remnant thrombus in both apices, rigidity of mitral and
tricuspid valve and diastolic dysfunction persisted despite
the initial favorable response to medical therapy. The
patient remains on drug therapy and serial follow up
echocardiography is also planned. In the future, more
studies will be needed to evaluate the efficacy of medical
therapies that would not only result in hematologic
remission of HES but also to bring out full recovery of end
organ damage without remnant sequelae.

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