Markedly high peripheral blood eosinophilia: A rare manifestation of Eosinophilic granulomatosis with polyangiitis

Abstract

Eosinophilic granulomatosis with polyangiitis (EGPA) is a systemic necrotizing multi-system vasculitis. It is rare for EGPA to present with severe eosinophilia. Here we report a case of EGPA, presenting with markedly high eosinophilia with tissue infiltrates. It was successfully treated with immunosuppressive medications.