Abstract

Eosinophilic Granulomatosis with Polyangiitis (EGPA) was earlier named as Churg Strauss syndrome or allergic granulomatosis with angiitis. It is a multisystem disease which is categorized as an Anti-Neutrophil Cytoplasmic Antibody Positive (ANCA) small vessel vasculitis. But it can affect medium sized vessels as well. The exact pathogenesis of EGPA is not known yet. Vasculitis features may not appear in the early phase of the disease and patients can present with several non-specific complaints. Although neuropathy is a commonly observed manifestation of EGPA, finding of isolated foot drop is a rare representation. As EGPA is a potentially life-threatening condition, knowledge regarding the unusual manifestations of this rare disease is important for the timely diagnosis and initiation of therapy. This case describes a 68-year-old known Asthmatic patient, presented with fever for 3 weeks and later developed unilateral foot drop who was ultimately diagnosed to be having EGPA.