A Rare Presentation of Eosinophilic Granulomatosis with Polyangiitis with Diffuse Lymphadenopathy

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Abstract
A 56-year-old man, who recently returned from a trip to Guyana, presented to hospital with a 10-week history of neuropathic pain, paresthesias, sinus pain, weight loss, and diffuse lymphadenopathy. Bloodwork was remarkable for significant eosinophilia (6.57 ×10^9 cells/L), erythrocyte sedimentation rate of 119 mm/hour and C-reactive protein of 44.3 mg/L. An extensive work up for an underlying lymphoproliferative disorder or infectious process was negative. Anti-MPO ANCA was initially borderline positive at 23 CU but on repeat testing 3 months later was strongly positive at 409 CU. He was diagnosed with eosinophilic granulomatosis with polyangiitis (EGPA) and was started on corticosteroids with an excellent clinical response. We review the epidemiology, typical clinical manifestations, diagnostic criteria and management of EGPA.

Résumé
Un homme de 56 ans, de retour récemment d’un voyage en Guyane, se présente à l'hôpital avec des problèmes de douleur neuropathique, de paresthésie, de douleur aux sinus, de perte de poids et d’une adénopathie diffuse qui durent depuis dix semaines. Les analyses sanguines révèlent une éosinophilie importante (6,57 ×10^9 cellules/L), une vitesse de sédimentation érythrocytaire de 119 mm après une heure et un taux de protéine C réactive de 44,3 mg/L. Une recherche poussée relative à la présence d’un trouble lymphoprolifératif sous-jacent ou d’un processus infectieux s’est avérée négative. L’ANCA anti-MPO était tout d’abord légèrement positif à 23 CU, mais une nouvelle analyse trois mois plus tard s’est révélée fortement positive, à 409 CU. On a alors diagnostiqué une granulomatose éosinophile et une inflammation simultanée de plusieurs vaisseaux sanguins ou lymphatiques (EGPA, pour eosinophilic granulomatosis with polyangiitis). Un traitement aux corticostéroïdes a été entrepris, avec une excellente réponse clinique. Nous passons en revue l’épidémiologie, les manifestations cliniques, les critères diagnostiques et la prise en charge de l’EGPA.
**Case Presentation**

A 56-year-old male presented with a 10-week history of pain and paresthesias in his extremities with sinus pain, weight loss, diffuse lymphadenopathy and generalized malaise. Past medical history included a recent left leg deep venous thrombosis diagnosed one month prior and chronic obstructive pulmonary disease (COPD) diagnosed 8 years prior. He had infrequent COPD exacerbations and was symptom free between episodes. Medications included rivaroxaban, salbutamol, tiotropium, and fluticasone/salmeterol. He emigrated from Guyana several years ago, but recently vacationed there.

Physical examination demonstrated cachexia, bilateral cervical and inguinal lymphadenopathy, and hepatomegaly. There were no rashes, swollen joints, or ulcers. There was decreased sensation in the median nerve distribution on the left forearm, the ulnar nerve distribution on the right forearm, and the sural nerve distribution of the left leg. Bloodwork revealed a leukocytosis of 19.9 × 10^9 cells/L with eosinophilia (6.57 × 10^9 cells/L). Erythrocyte sedimentation rate was 119 mm/hr and C-reactive protein (CRP) 44.3 mg/L. c-ANCA was positive, p-ANCA was negative. Interestingly, his anti-PR3 ANCA titre was within normal limits at 3 CU but his anti-MPO ANCA was borderline positive at 23 CU. Chest computed tomography (CT)-scan showed subpleural groundglass opacification in the left upper lobe. Abdominal CT-scan showed thrombosis of the hepatic vein and ischemic changes in the liver parenchyma consistent with Budd-Chiari syndrome. There was bilateral inguinal adenopathy. The patient was admitted to hospital for further investigations for a possible lymphoproliferative disorder, infection, or vasculitis.

Cultures of blood and stool were negative. Interferon-gamma release assay for tuberculosis was negative. Serology for strongyloides, filariasis, malaria, human immunodeficiency virus, and human T-cell lymphotropic virus I/II was negative. Electromyography showed a length-dependent neuropathy suggestive of mononeuritis multiplex. Molecular genetic testing for mutations associated with clonal eosinophilic disorders was negative. Bone marrow biopsy demonstrated reactive changes with no malignant cells but significant eosinophilia (20% of all bone marrow cells). An excisional lymph node biopsy revealed a necrotic and hemorrhagic node with vague non-caseating granulomas in the surrounding lymph node capsule (Figure 1). ANCA serology was repeated and anti-MPO ANCA was now strongly positive at 409 CU. A sural nerve biopsy was planned, but the patient's symptoms progressed to the point where he was debilitated. He was started on corticosteroids for a presumed diagnosis of eosinophilic granulomatosis with polyangiitis (EGPA). His symptoms improved and his eosinophilia resolved on steroid therapy. Azathioprine was started as his steroid dose was tapered.

![Figure 1. Histopathology from lymph node biopsy.](image)

**Discussion**

EGPA, formerly known as Churg-Strauss syndrome, is a rare, life-threatening ANCA-associated systemic vasculitis. The annual incidence is estimated at 2.4–6.8 cases per million people. The disorder was first described by Jacob Churg and Lotte Strauss in 1951, when they published a series of 13 patients who presented with a necrotizing systemic vasculitis, with pathologic evidence of extravascular granulomas and eosinophilia on a background history of asthma and atopy. The pathophysiology is poorly understood and the role of ANCA remains controversial. Patients have a prodrome of asthma with progressive development of peripheral eosinophilia, culminating in a life-threatening necrotizing vasculitis affecting small to medium sized vessels. Airways disease can precede the vasculitic manifestations for several years.

The diagnosis of EGPA is challenging to make. The classic features of eosinophilia, extravascular granulomas and necrotizing vasculitis described by Churg and Strauss rarely co-exist in the same patient at a given time. Biopsy of an affected organ is recommended but can be falsely negative.

The American College of Rheumatology (ACR) proposed six diagnostic criteria for EGPA: presence of asthma, eosinophilia >10%, mononeuropathy or polyneuropathy, transient pulmonary infiltrates, paranasal sinus abnormalities, and pathologic confirmation of extravascular eosinophils. The ACR found that satisfying at least 4 of these 6 criteria yielded a sensitivity of 85% and specificity of 99.7% for diagnosing EGPA. Our patient satisfied at least 4 criteria (eosinophilia >10%, mononeuropatitis multiplex, paranasal sinus abnormalities, pulmonary infiltrates). Though our patient was diagnosed with COPD remotely, we suspect he actually had asthma given the absence of symptoms between exacerbations. The timeline of his initial diagnosis with...
airways disease and the development of the vasculitis fits with the natural history of EGPA.

The largest study of patients with EGPA is the French Vasculitis Study Group Cohort. The study enrolled 383 patients diagnosed with EGPA from 1957–2009. The mean age of subjects was 50.3 years and 98.2% were Caucasian. Only 31.0% were ANCA positive. Common disease manifestations included peripheral neuropathy, weight loss, otolaryngological symptoms, skin lesions, and pulmonary infiltrates. This case of EGPA is unique because the patient's lymphadenopathy was very pronounced, and he had a history of 2 recently diagnosed deep venous thromboses (left leg DVT and Budd-Chiari syndrome). Venous thromboembolic disease was unusual among the French Vasculitis Study Group patients. The prevalence of lymphadenopathy in the group was not described by the authors. Lymph node involvement in EGPA has previously been described as exceedingly rare.

EGPA was frequently fatal prior to the development of immunosuppressive therapy. The prognosis has improved significantly with the advent of glucocorticoids. Guidelines issued jointly by the British Society for Rheumatology (BSR) and British Health Professionals in Rheumatology (BHPR) for the management of adults with ANCA-associated vasculitis recommends induction therapy with high-dose glucocorticoids and cyclophosphamide or rituximab. Methotrexate or mycophenolate mofetil are alternative induction agents in non life- or organ-threatening disease. Some sources suggest that glucocorticoid therapy alone is sufficient to induce remission in EGPA in up to 40% of cases. Our patient has had an excellent response to glucocorticoids.

References