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Development of Anti-CCP-positive Rheumatoid Arthritis Following Pegylated Interferon-α2a Treatment for Chronic Hepatitis C Infection

To the Editor:

Interferon-α (IFN-α) is a group of cytokines with antiviral and antiproliferative effects, used for the treatment of chronic hepatitis C infection and various malignancies. Immunomodulatory effects of IFN may lead to the induction or exacerbation of autoimmune diseases such as psoriasis, systemic lupus erythematosus, and rarely, rheumatoid arthritis (RA). Covalent attachment of a polyethylene glycol (PEG) moiety (pegylation) to IFN-α results in a significantly higher sustained virological response rate in patients with chronic hepatitis C compared to conventional IFN-α2a. Pegylation also reduces the immunogenicity of IFN-α2a. We describe the first case of a patient who developed anticyclic citrullinated peptide antibody (anti-CCP)-positive RA following treatment of chronic hepatitis C infection with pegylated IFN-α2a.

A 54-year-old Chinese man with chronic hepatitis C (genotype 2a) infection was placed on a 24-week course of PEG-IFN-α2a 180 µg weekly and ribavirin 400 mg BID. His hepatitis C virus (HCV) RNA became undetectable at Week 12 and remained undetectable throughout his treatment. With no history of arthritis, he developed a diffuse pain syndrome 18 weeks into antiviral treatment. Initially this was attributed to myalgias or possible myositis, but creatine phosphokinase (CPK) and aldolase levels were normal. Treatment with ibuprofen 200 mg every 6 h, celecoxib 200 g/day, and tramadol 100 mg every 6 h provided only minimal relief. Upon completion of a 24-week course of antiviral therapy he developed a sustained virological response as evidenced by persistent undetectable HCV RNA and normal aminotransferase activities. However, by the end of his 24 weeks of antiviral therapy, he developed bilateral pain in his shoulders, elbows, hands, knees, and feet. The musculoskeletal examination was notable for active synovitis of the proximal phalangeal joints, metacarpophalangeal joints, wrists, elbows, shoulders, knees, ankles, and feet. Distal interphalangeal joints were spared. Handgrip was 50% of normal. He had no musculoskeletal symptoms prior to antiviral therapy. Review of systems was otherwise unremarkable. Laboratory results revealed anti-CCP 41.27 IU/ml (negative < 5 IU/ml), erythrocyte sedimentation rate 163 mm/h, C-reactive protein > 192 mg/l (normal < 6 mg/l), CPK 17 U/l (normal 24–170 U/l), negative antinuclear antibodies, undetectable cryoglobulins, negative rheumatoid factor (RF; normal < 8 IU/ml), hemoglobin 11.3 g/dl, and thyroid-stimulating hormone 3.18 IU/ml. Radiographs showed juxtaarticular osteopenia and early erosions of wrist joints. A positive tuberculin skin test precluded the initiation of an anti-tumor necrosis factor agent. Instead he was treated with hydroxychloroquine 400 mg and sulfasalazine 2 g/day, which resulted in dramatic improvement of symptoms.

IFN-α has rarely been reported to induce or exacerbate RA in patients with chronic hepatitis C infection. Okanoue, et al reported that out of 677 patients treated with high-dose pegylated IFN for chronic hepatitis C infection, 2 patients developed RA. Most case reports of patients developing RA after treatment with pegylated IFN for hepatitis C show persistent arthritis unresponsive to nonsteroidal antiinflammatory drugs despite discontinuation of IFN therapy.

The possibility of pegylated IFN inducing or exacerbating RA in previously predisposed individuals should be considered in patients with hepatitis C who develop arthritis. Screening for RA and anti-CCP may be considered before treating with IFN. Detection of anti-CCP, the presence of nodules, and erosive disease help distinguish RA from HCV-related arthritis, but these characteristics are not always found in patients who develop RA with IFN treatment. The prognosis of patients with PEG-IFN-associated RA remains unclear because of the small number of cases reported. If the symptoms of RA do not resolve after discontinuing PEG-IFN, these patients may require treatment with disease-modifying antirheumatic drugs.

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