Successful Treatment of Progressive Rheumatoid Interstitial Lung Disease With Cyclosporine: A Case Report

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INTRODUCTION

Interstitial lung disease (ILD) in rheumatoid arthritis (RA) has been controversial. Although there have been several anecdotal reports on the efficacies of corticosteroids or cytotoxic agents such as methotrexate, cyclophosphamide, azathioprine, and D-penicillamine for the treatment of ILD associated with RA, no controlled studies have been performed. To date, corticosteroids have been a central agent for the treatment of this disease, but their effects are partial and temporary in most cases. In addition, the adverse effects of these agents are considerable. On the other hand, limited information is available on the cyclosporine use in ILD associated with RA. We describe a 49-yr old female patient with RA and ILD that had initially responded to high dose prednisolone and cyclophosphamide intravenous pulse therapy, and the lung disease was aggravated with the tapering of prednisolone. After 10 months of follow-up loss, the patient was successfully treated with low dose cyclosporine without high dose corticosteroids.

Key Words: Lung Disease, Interstitial; Arthritis, Rheumatoid; Cyclosporine

CASE REPORT

In March 1999, a 49-yr-old woman presented with a 1-yr history of polyarthritis and shortness of breath on exertion. She did not smoke and had no past history of environmental toxin exposure. There was no medical history of photosensitivity, oral ulcer, alopecia, and Raynaud phenomenon. On physical examination, inspiratory crackles were heard on the both lower lung fields, and heart sounds were normal. The joint swelling and tenderness were noted on the small joints of both hands, right elbow, right first metatarsophalangeal joint, and left third metatarsophalangeal joint. Laboratory studies at this time showed a hemoglobin level of 11.3 g/dL and a white blood cell count of 5,200/μL. The erythrocyte sedimentation rate was 50 mm/hr. The C-reactive protein was not increased. The rheumatoid factor was 340 IU/mL and the antinuclear antibody titer was 1:640 with a speckled pattern. The anti-extractable nuclear antigen and anti-double-stranded DNA antibodies were absent. The complement level was normal. There were no specific abnormalities on other chemistry profiles.

A chest radiograph revealed bibasilar interstitial infiltrates (Fig. 1A). High resolution computerized tomography (HRCT) of the chest showed interstitial fibrosis, and ground glass
opacities suggesting active inflammation on both lower lung fields. A pulmonary function test revealed a mild restrictive pattern with forced vital capacity (FVC) of 78% of predicted. The carbon monoxide diffusing capacity was 46% of predicted. Although juxta-articular osteoporosis in the plain radiographs was noted around proximal interphalangeal joints of both hands, right wrist, and right elbow, bony erosions were absent.

A diagnosis of RA with active ILD was established. Her initial medications included prednisolone 1 mg/kg daily, hydroxychloroquine 200 mg b.i.d. and nabumetone 500 mg b.i.d. Monthly cyclophosphamide intravenous (IV) pulse therapy with a dose of 600 mg/m² was started. Three weeks later, dyspnea and polyarthralgia improved. The pulmonary function test was also improved with FVC of 85% of predicted and diffusing capacity of 56% of predicted. The dosage of prednisolone was gradually tapered to 7.5 mg over the next 4 months. Monthly cyclophosphamide IV pulse therapy was continued.

In September 1999, she complained of a shortness of breath of a greater severity. At that time, pulmonary function was aggravated with FVC 72% of predicted and diffusing capacity of 50%. HRCT showed progressive basal interstitial infiltrates and increased ground glass opacities compared with
the findings in March 1999. Cyclosporine 3.6 mg/kg (200 mg) daily was prescribed, and prednisolone was increased to 20 mg daily, but she did not take this medication and was lost from follow-up. During this period she had not taken any medical treatment except herb medications.

In July 2000, she was admitted again with complaints of shortness of breath on minimal exertion and severe polyarthralgia. Physical examination revealed inspiratory crackles and coarse breath sounds on both lower lung fields. There were swelling and tenderness on the multiple joints such as small joints of both hands, both elbows, both knee joints, and right ankle. On laboratory studies, a white blood cell count was 10,000/μL and a hemoglobin level was 11 g/dL. Arterial blood gas analysis without O₂ supplementation showed a PaO₂ of 54.1 mmHg, PaCO₂ of 27.6 mmHg, and O₂ saturation of 90.1%. The lung infiltrates on the chest radiograph (Fig. 1B) and HRCT (Fig. 2) were more aggravated with an extension to left upper lobe. The pulmonary function was also aggravated with FVC of 54% of predicted and diffusing capacity of 37% of predicted. A transbronchial lung biopsy showed interstitial fibrosis and mononuclear cell infiltrates, which were compatible with interstitial pneumonitis. Bacterial cultures grew normal flora of the upper respiratory tract. Cultures for fungus and mycobacteria were negative.

She was considered to have as advanced ILD with active inflammation in association with RA. Daily cyclosporine 3.6 mg/kg (200 mg), prednisolone 5 mg, and celecoxib 200 mg were started. Within 4 weeks, her breathlessness and joint symptoms markedly improved. The pulmonary function test revealed FVC of 69% of predicted and diffusing capacity of 39% of predicted. Serum trough level of cyclosporine was 119 ng/mL. In December 2000, her shortness of breath further improved and stabilized. Chest radiograph (Fig. 1C) and HRCT at this time showed markedly reduced pulmonary infiltrates. The dosage of cyclosporine could be reduced to 2.7 mg/kg (150 mg) daily. Mild hypertension which was the only side effect of cyclosporine, was controlled with amlodipine. In January 2001, she was free of joint symptoms, and only had mild dyspnea on exertion.

**DISCUSSION**

The clinical course of ILD associated with RA is variable. Whereas some patients take gradually progressive clinical course (7), other patients run a rapidly progressive course that can be fatal within several months (2). Spontaneous remission for this disease has been rarely reported (8). The clinical presentation and the pathologic findings of ILD in RA are known to be very similar to those of idiopathic pulmonary fibrosis, and the therapeutic modalities for these two diseases are not much different. Although some RA patients with progressive ILD may respond to aggressive immunosuppressive therapy, treatment for this disease still remains controversial.

To date, the corticosteroids have been the central agent for the treatment of progressive ILD in RA. However, the favorable response rate to these agents has been reported less than 40%, and documentation of sustained effect is even rare (2, 9, 10). Although there have been several anecdotal reports for the efficacies of cytotoxic agents such as methotrexate, azathioprine, D-penicillamine, and cyclophosphamide in patients with RA and progressive ILD, few controlled studies have demonstrated the sustained effects of these agents (2).

The efficacy of cyclosporine for the treatment of RA has been proven (11-13). Even if there have been a few reports for the successful cyclosporine use in progressive ILD or acute interstitial pneumonitis in association with RA, the available...
information is still limited (14-17). Alegre et al. first described the successful treatment of aggressive pulmonary fibrosis in RA with cyclosporine (14). In contrast to our case, that patient had symptoms for less than one month before treatment, and had not been treated with any other immunosuppressive agents. In addition they used high dose prednisone and higher dose cyclosporine than in our case. Puttick et al. reported the effect of cyclosporine in a patient with long-standing RA and progressive ILD that had been refractory to corticosteroids and cytotoxic agents such as chlorambucil and oral cyclophosphamide. They used similar dose of cyclosporine and higher dose of prednisolone, compared with our case (15).

Our patient had initially responded to high dose prednisolone and cyclophosphamide IV pulse therapy. However, her lung disease was aggravated with the tapering of prednisolone for 6 months. After 10 months of follow-up loss, ILD was much more aggravated. At that time, she was treated with low doses of cyclosporine and prednisolone (5 mg/day). Despite subjective symptoms, her pulmonary function such as forced vital capacity, and radiographic pulmonary infiltrates have much improved with these agents, the diffusing capacity did not change. This may reflect the ventilation-perfusion mismatch in irreversibly damaged lung lesions. Most patients in the literature who showed the effectiveness of cyclosporine, including our case, the improvement exhibited within one month. Because other patients than ours had used cyclosporine together with higher dose of corticosteroids, it is not certain whether the therapeutic improvement was due to cyclosporine or to corticosteroids. On the other hand, as the spontaneous remission is extremely rare in progressive ILD associated with RA, the improvement of the lung disease in the present case was mostly considered to be the effect of cyclosporine.

Cyclosporine, a complex fungal decapetide, has highly selective immunosuppressive properties. This agent appears to reduce interleukin-2 synthesis by activated T-cell (18) and to disrupt cytokine-dependent T lymphocyte-macrophage interaction, and thus prevents fibroblast-mediated fibrosis that is thought to be important in the pathogenesis of ILD (2). Recently, Sekigawa et al. described 4 cases of steroid-resistant autoimmune diseases which were effectively treated with an extremely low dose of cyclosporine (1 mg/kg/day) (19). Due to the limited information of cyclosporine in the treatment of RA with ILD, the controlled studies are needed to clarify whether cyclosporine is effective agent for the treatment of ILD associated with RA, whether high dose corticosteroids are required together with cyclosporine, and which dosage of cyclosporine is optimal.

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