Acute pneumonitis associated with low dose methotrexate treatment for rheumatoid arthritis: report of five cases and review of published reports

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Abstract

Background Low dose methotrexate has become established in the treatment of refractory rheumatoid arthritis. Until recently it has been considered that the use of a low dose regimen (<20 mg/week) would avoid the pulmonary toxicity associated with the higher doses prescribed in malignant disease. Although initial experience with low dose methotrexate was encouraging, an increasing number of cases of an acute, life-threatening pneumonitis are being reported in patients with refractory rheumatoid arthritis.

Patients Since 1984 43 patients with refractory rheumatoid arthritis have been established on low dose methotrexate in the Oxford Health District. Five of these patients have subsequently developed acute methotrexate induced pneumonitis. The clinical and radiological features of these cases are described and previous reports reviewed.

Results Five patients having low dose methotrexate treatment developed acute pneumonitis. Presentation was subacute and dominated by constitutional features. Respiratory symptoms developed insidiously but progressed rapidly with increasing dyspnoea associated with severe hypoxia. Chest radiographs were non-specific, showing diffuse interstitial infiltration and alveolar shadowing. Microbiological investigation gave negative results. In all cases methotrexate was discontinued and high dose corticosteroids started, with rapid clinical and radiological improvement. After withdrawal of steroid both clinical and radiological resolution was maintained at follow-up.

Conclusion Acute pneumonitis is an uncommon but serious adverse effect of low dose methotrexate treatment for refractory rheumatoid arthritis. The initial presentation is non-specific and a high index of suspicion is required as respiratory failure may develop rapidly. Management depends on exclusion of infection, withdrawal of methotrexate, and high dose corticosteroid treatment. Full supportive treatment is indicated as the prognosis in such patients is good.

The value of low dose methotrexate in the treatment of refractory rheumatoid arthritis is well established. Although pulmonary toxicity associated with methotrexate is recognised, it has been considered that a low dose regimen may avoid such problems. Initial experience was encouraging but in 1983 the first cases of acute methotrexate induced pneumonitis were reported. Since then over 30 such cases have been published worldwide, reflecting increased enthusiasm for the earlier use of methotrexate in rheumatoid arthritis rather that its occasional use in refractory cases. We describe five further cases and review previous reports.

Case reports

Patient 1 In August 1988 a 47 year old woman with a 13 year history of seropositive rheumatoid arthritis was referred for investigation of dyspnoea. In March 1988, as a result of a sustained inflammatory exacerbation, she had been started on methotrexate 10 mg/week (total dose 100 mg), which in conjunction with fenbufen resulted in a good therapeutic response. Three weeks before admission she developed a non-productive cough associated with general malaise and despite antibiotics became increasingly dyspnoeic.

On examination she was cyanosed and dyspnoeic at rest. Her temperature was 38.5°C, pulse 140/min regular, and blood pressure 120/80 mm Hg. Examination of the chest indicated scattered inspiratory crepitations at the right base.

The haemoglobin concentration was 10.3 g/dl and white blood cells 8.4 × 10⁹/l eosinophilia. The erythrocyte sedimentation rate was 39 mm in the first hour. Arterial blood gas analysis (with an inspired oxygen concentration (FiO₂) of 21%) showed that pH was 7.47, arterial carbon dioxide tension (Paco₂) 4-10 kPa, arterial oxygen tension (Pao₂) 5-26 kPa. The chest radiograph showed diffuse bilateral interstitial infiltrates (fig 1). Cultures of sputum and blood samples were negative and serological tests on convalescent phase serum showed no rise in titres of antibodies to atypical agents. Methotrexate was stopped. Ampicillin and erythromycin were administered intravenously.

Over the following 48 hours her condition deteriorated and a repeat chest radiograph showed increase interstitial shadowing (fig 1B).

She was transferred to the medical intensive care unit, where she was electively intubated and ventilated before bronchoscopy. Bronchoalveolar lavage fluid showed no pathogens. Prednisolone 60 mg daily was prescribed and over the subsequent 72 hours her condition...
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PATIENT 1
In November 1989, a 58 year old man, who had a 25 year history of seropositive rheumatoid arthritis was referred for assessment of dyspnoea. In November 1989, after he had developed rheumatoid-vasculitis ulceration of his legs, he was started on methotrexate 10 mg/week (total dose 430 mg).

Two months before presentation he developed a non-productive cough associated with a coryza type illness. Despite antibiotics he deteriorated and became increasingly breathless.

On examination he was afebrile and dyspnoeic with minimal activity. His pulse was 80/min regular and blood pressure 120/70 mm Hg. Examination of the chest revealed scattered inspiratory crepitations at both lung bases. The results of arterial blood gas analysis (FiO2 21%) were: pH 7.42, PaO2 4.27 kPa PaO2 11.99 kPa. Pulmonary function testing showed that forced expiratory volume in one second (FEV1, was 2.7 (predicted 2.1–3.8) l), forced vital capacity (FVC) 3.1 (predicted 2.6–4.7) l, alveolar volume (VA) 4.2 (predicted 4.9–7.3) l, transfer coefficient (Kco) 0.79 (predicted 1.00–1.89) mmol min⁻¹ kPa⁻¹ l⁻¹. The chest radiograph showed diffuse interstitial infiltration.

Methotrexate induced pneumonitis was considered likely and he was advised to discontinue methotrexate and to report any subsequent deterioration without delay.

Four days later he presented with increasing dyspnoea. On examination he was cyanosed and dyspnoeic at rest with basal inspiratory crepitations. The haemoglobin concentration was 13.0 g/dl and white blood cells 12.7 x 10⁹/l with no eosinophilia. The results of arterial blood gas analysis (FiO2 21%) were: pH 7.45, PaCO2 4.02 kPa, PaO2 5.18 kPa. A repeat chest radiograph showed increased interstitial shadowing. Cultures of sputum and blood samples were negative and serological tests on convalescent phase serum showed no rise in titres of antibodies to atypical agents.

Ampicillin and hydrocortisone succinate (200 mg six hourly) were administered intravenously with oral prednisolone 60 mg daily.

By the fourth day he had considerably improved and oxygen saturation was consistently above 95% (FiO2, 28%). Amoxicillin and hydrocortisone were discontinued and oral prednisolone continued. He was discharged after 12 days taking prednisolone 30 mg daily, which was tailed off over the subsequent two months.

When reviewed four months later he was symptom free, though troubled by a recurrence of articular symptoms. A chest radiograph and results of repeat pulmonary function tests were normal.

PATIENT 2
In October 1990 a 58 year old man with a 25 year history of seropositive rheumatoid arthritis was referred for assessment of dyspnoea. In November 1989, after he had developed rheumatoid-vasculitis ulceration of his legs, he was started on methotrexate 10 mg/week (total dose 430 mg).

Two months before presentation he developed a non-productive cough associated with a coryza type illness. Despite antibiotics he deteriorated and became increasingly breathless.

On examination he was afebrile and dyspnoeic with minimal activity. His pulse was 80/min regular and blood pressure 120/70 mm Hg. Examination of the chest revealed scattered inspiratory crepitations at both lung bases. The results of arterial blood gas analysis (FiO2 21%) were: pH 7.42, PaO2 4.27 kPa PaO2 11.99 kPa. Pulmonary function testing showed that forced expiratory volume in one second (FEV1, was 2.7 (predicted 2.1–3.8) l), forced vital capacity (FVC) 3.1 (predicted 2.6–4.7) l, alveolar volume (VA) 4.2 (predicted 4.9–7.3) l, transfer coefficient (Kco) 0.79 (predicted 1.00–1.89) mmol min⁻¹ kPa⁻¹ l⁻¹. The chest radiograph showed diffuse interstitial infiltration.

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Ampicillin and hydrocortisone succinate (200 mg six hourly) were administered intravenously with oral prednisolone 60 mg daily.

By the fourth day he had considerably improved and oxygen saturation was consistently above 95% (FiO2, 28%). Amoxicillin and hydrocortisone were discontinued and oral prednisolone continued. He was discharged after 12 days taking prednisolone 30 mg daily, which was tailed off over the subsequent two months.

When reviewed four months later he was symptom free, though troubled by a recurrence of articular symptoms. A chest radiograph and results of repeat pulmonary function tests were normal.

PATIENT 3
In November 1988 a 67 year old woman with a 15 year history of seropositive rheumatoid
arthritis was admitted with a short history of dyspnoea.

Intramuscular gold injections and chloroquine had been of limited value and she continued to have symptoms despite oral prednisolone. In August 1988 she was started on methotrexate 10 mg/week (total dose 130 mg), with moderate symptomatic improvement. In the two weeks before admission she became increasingly dyspnoeic. A chest radiograph showed diffuse interstitial and alveolar infiltration.

On examination she was cyanosed and dyspnoeic at rest. Her temperature was 39·0°C, pulse 60/min, and blood pressure 105/70 mm Hg. Examination of her chest indicated harsh inspiratory crepitations throughout both lung fields.

The haemoglobin concentration was 8·9 g/dl and white blood cells 6·6 × 10⁹/l, with 12% eosinophils. The erythrocyte sedimentation rate was 110 mm in the first hour. The results of arterial blood gas analysis (FiO₂ 40%) were: pH 7·42, PaCO₂ 7·7 kPa, PaO₂ 7·21 kPa. Cultures of sputum and blood samples were negative and serological tests on acute and convalescent phase serum showed no rise in titres of antibodies to atypical agents.

Methotrexate was stopped and broad spectrum antibiotics were administered intravenously with oral prednisolone 100 mg daily. Her condition thereafter rapidly improved with both clinical and radiological resolution. She returned to the referring hospital after 10 days taking prednisolone 30 mg daily, which was tailed off over two months.

When reviewed two months later she was symptom free and her chest radiograph normal.

PATIENT 4
In December 1990 a 75 year old woman with an 11 year history of seropositive rheumatoid arthritis was admitted with dyspnoea. In August 1988 she had been started on methotrexate 5 mg/week (total dose 600 mg), with an excellent clinical response. Two months before admission she developed a non-productive cough, fever, and increasing dyspnoea, which had progressed despite antibiotics.

On admission she was cyanosed and dyspnoeic at rest. She was afebrile. Her pulse was 120/min and blood pressure 150/80 mm Hg. Examination of the chest indicated late inspiratory crepitations throughout both lung fields.

The haemoglobin concentration was 10·0 g/dl and white blood cells 24·5 × 10⁹/l, with eosinophilia. The erythrocyte sedimentation rate was 87 mm in the first hour. The results of arterial blood gas analysis (FiO₂ 21%) were: pH 7·5, PaCO₂ 5·3 kPa, PaO₂ 7·1 kPa. The chest radiograph showed bilateral interstitial and alveolar infiltrates, affecting predominantly the upper lobes (fig 2A). Cultures of sputum and blood samples were negative and serological tests on convalescent phase serum showed no rise in titres of antibodies to atypical agents.

Methotrexate was stopped and she was started on prednisolone 40 mg daily. There was considerable clinical and radiological resolution over the next two weeks and she was discharged on a reducing course of corticosteroids. By three months she was symptom free. A chest radiograph confirmed radiological resolution (fig 2B).

PATIENT 5
In March 1991 a 76 year old man with an 11 year history of seropositive rheumatoid arthritis was admitted with cough and increasing dyspnoea. In November 1990 he was started on methotrexate 10 mg/week (total 180 mg). A marginal response was observed.

Two months before admission he had noted sweats, fever, and general malaise associated with anorexia and weight loss. Ten days before admission he had developed a non-productive cough with increasing dyspnoea.

Other medications included prednisolone 5 mg daily, tetracosactrin 2 mg/week by intramuscular injection, azapropazone 600 mg twice daily, glibenclamide 5 mg daily, and atenolol 50 mg daily.
On examination he was dyspnoeic at rest. He was afebrile. His pulse rate was 80/min regular and blood pressure 130/90 mm Hg. Examination of the chest indicated scattered inspiratory crepitations at both bases.

The haemoglobin concentration was 11.3 g/dl and white blood cells 9.1 × 10⁹/l, with no eosinophilia. The results of arterial blood gas analysis (Fio₂ 21%) were: pH 7.46, Paco₂ 4.61, Pao₂ 8.23. Pulmonary function testing showed that FEV₁ was 1.2 (predicted 2.2–3.99), FVC 2.6 (predicted 3.9–5.1), Kco 0.8 (predicted 0.77–1.67) mmol min⁻¹ kPa⁻¹ l⁻¹. The chest radiograph showed diffuse interstitial and alveolar infiltrates. Cultures of sputum and blood samples were negative and serological tests on convalescent phase serum showed no rise in titres of antibodies to atypical organisms.

Methotrexate was stopped and erythromycin 500 mg four times daily was administered by mouth. Serial blood gas determinations and the chest radiograph appearances suggested continued deterioration and on the fourth hospital day prednisolone 40 mg daily was prescribed. Within 72 hours there was considerable clinical and radiological improvement. He was discharged after 30 days taking prednisolone 10 mg daily.

Discussion

It seems likely that the five patients described had methotrexate induced pneumonitis. This diagnosis, however, can only be made with reasonable confidence retrospectively in the light of the clinical course and negative results of microbiological investigations. At the time of presentation the main differential diagnoses included infection, rheumatoid related alveolitis, and drug induced pneumonitis. The fact that four out of the five patients had constitutional symptoms or fever led to the assumption that their illness could be infective and thus could be treated with antibiotics. The subacute onset of symptoms and the diffuse pattern of radiological infiltrates, however, were unusual for a bacterial infection. Atypical and opportunistic organisms were considered but the clinical response coincided with the introduction of corticosteroids. Sputum and blood cultures were negative in all patients and the serological tests for atypical organisms showed no rise in antibody titres. The decision to undertake bronchoalveolar lavage may be necessary if there is clinical deterioration and diagnostic uncertainty persists. Its potential usefulness must be balanced against the fact that these patients were severely hypoxic and in the one patient who had bronchoscopy elective intubation and assisted ventilation were necessary. Samples obtained at lavage were negative for Pneumocystis carinii and fungi.

Patients with rheumatoid arthritis may develop diffuse interstitial lung disease unrelated to any medication. The onset is usually insidious with breathlessness as the main presenting feature. It would be unusual for the disease to evolve as rapidly as that seen in our patients. The most convincing evidence, however, that their alveolitis was not related to rheumatoid arthritis is the gratifying response to corticosteroids and the fact that the patients remained well from a respiratory viewpoint after the steroids had been stopped.

The clinical presentation of methotrexate induced pneumonitis is usually subacute and dominated by constitutional symptoms, including headache, malaise, and fever. These may persist for weeks or months before the onset of respiratory symptoms heralded by dry cough and dyspnoea. The early symptoms are often indeterminate, though in some instances may advance rapidly. For example, in patient 2 extreme dyspnoea and hypoxia were observed over 72 hours. Similarly, in patient 1 relentless deterioration was observed after admission and ventilatory support was required. The findings from clinical examination are generally nonspecific. A rash has been reported in up to 16% of cases of pneumonitis associated with high dose methotrexate for malignant disease. Fever, dyspnoea, and inspiratory crepitations have been consistently reported. Mild neutrophil leucocytosis has been observed in about 30% of cases. Although eosinophilia has been reported, it appears to be unusual with the low dose regimen. Radiological appearances are non-specific and usually consist of bilateral interstitial infiltrates, often with alveolar shadowing. Transient hilar lymphadenopathy has been reported. Although usually diffuse, the radiological abnormalities may be confined initially to the lower lobes or even to one lung. The radiological appearances in patient 4 were somewhat unusual as the infiltrates were predominantly in the upper lobes.

The role of open lung biopsy in the management of methotrexate induced pneumonitis is uncertain as the histological changes are non-specific, though consistent with drug induced pneumonitis. Findings include an interstitial mononuclear cell infiltrate associated with a variable degree of fibrosis. Multinucleated giant cells, non-caseating granulomas, and eosinophilic infiltrates have also been reported. Whereas bronchoscopy and bronchoalveolar lavage may help in excluding infection, cytological examination of lavage fluid is generally unhelpful. There may be an absolute or relative increase in the lymphocyte count but this information is of limited diagnostic value.

Treatment requires the withdrawal of methotrexate, which of itself may effect clinical and radiological resolution. Nevertheless, as was the case in at least two of our patients, the disease may progress and clinical improvement coincide with the introduction of corticosteroid treatment. Such observations have prompted treatment with corticosteroids when the diagnosis seems at all likely. No large scale trials have examined the efficacy of corticosteroids in methotrexate induced pneumonitis but reports throughout the literature attest to their value. It has been suggested that a limited controlled trial is warranted, though we believe that this would be inappropriate in a life threatening condition. Full supportive measures, including assisted ventilation, is indicated in patients who are critically ill because the ultimate prognosis is good.
The mechanism of methotrexate induced pneumonitis is uncertain. Some observations have suggested that it may represent an acute hypersensitivity reaction. The presentation with fever and eosinophilia is consistent with the hypersensitivity reaction and is supported by the finding in alveolar lavage fluid of lymphocytic alveolitis. The exact subpopulation of lymphocytes affected is not clear. One study reported an increase in OKT4 (helper) cells similar to that seen in sarcoidosis, whereas another reported an increase in OKT8 (suppressor) cells consistent with hypersensitivity pneumonitis—for example, in response to gold or organic antigen. Further evidence of an immunological mechanism is provided by the reported release of a lymphokine, leucocyte inhibitory factor, from peripheral blood lymphocytes of patients with methotrexate induced pneumonitis. Finally, the often dramatic response to corticosteroids supports an immune mediative phenomenon. The spontaneous remission during treatment, however, and reports that pulmonary toxicity may not recur after rechallenge remain to be explained.

As methotrexate has been shown to accumulate preferentially in the lung and clinical resolution may follow dose reduction, it has been suggested that a direct cytotoxic action may be important in some instances. Acute pulmonary toxicity appears, however, to be unrelated to either the dose or the duration of treatment. One important variable appears to be the frequency of administration since patients receiving daily or even weekly doses have a higher risk of toxicity than those receiving the drug less frequently.

Two recent prospective studies have addressed the incidence of pulmonary toxicity during low dose methotrexate treatment for active rheumatoid arthritis. Hanrahan et al described one such case from a cohort of 128 patients followed for up to four years, and Furst et al reported no instances of acute pneumonitis from 45 patients during a treatment period extending over three years. Retrospective analysis based on several series, however, would suggest an incidence of 3–5%. The five cases described here are derived from a cohort of 43 patients prescribed low dose methotrexate over the previous three years, representing an overall incidence of almost 12%. The Committee on Safety of Medicines (personal communication) has received eight reports of pneumonitis associated with methotrexate.

Although low dose methotrexate has been used for over 30 years in the management of psoriasis there have been relatively few reports of acute pulmonary toxicity. Likewise, the incidence of pulmonary toxicity in patients with psoriasis remains uncertain, though Nifors reports an incidence of 2.4% over an average follow up of three years, lending support to suggestions that rheumatoid patients may be at particular risk of methotrexate lung disease.

Methotrexate represents an important advance in the management of refractory rheumatoid arthritis, where a gratifying clinical response may be observed in selected patients. Accordingly, it is disappointing that, despite low dose regimens, methotrexate may be associated with acute life threatening pneumonitis. We would suggest that before starting treatment patients should have a full clinical, radiological, and physiological assessment for detection of occult lung disease. Further prospective studies are required to examine both the incidence of acute pneumonitis during low dose treatment and potential risk factors for its development.
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