Pancytopenia related to azathioprine in rheumatoid arthritis

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SUMMARY Two patients with rheumatoid arthritis developed pancytopenia during treatment with azathioprine 100 mg daily. In one patient this side effect occurred after three weeks, in the other after eight weeks of treatment. Rapid fall of platelets in one patient necessitated platelet transfusion. In the other patient additional treatment with allopurinol was probably responsible for the toxic effect. Haematological side effects of azathioprine are discussed.

Key words: adverse drug reaction, bone marrow suppression.

Azathioprine has been extensively used for more than 20 years in the treatment of patients with rheumatoid arthritis (RA) who are unresponsive or react with side effects to gold salts or D-penicillamine, or both. At a dosage of 1.5–2.5 mg/kg/day the toxic reactions of azathioprine are moderate and limited mainly to gastrointestinal and mild haematological complications.^{1 2}

Pancytopenia in relation to this drug is a rare side effect. So far, 18 cases of pancytopenia during azathioprine treatment used for a variety of diseases, given in different dosages and in combination with different other drugs, have been reported world wide.³ Detailed descriptions of these cases are mostly lacking.

We report two patients with rheumatoid arthritis who developed pancytopenia while receiving azathioprine 100 mg daily. In one patient platelet transfusion was necessary because of severe thrombocytopenia. The pancytopenia disappeared in both patients after azathioprine treatment was stopped.

Case reports

CASE 1

A 36 year old woman was diagnosed as suffering from seropositive RA in November 1974. Besides non-steroidal anti-inflammatory drugs she was treated successively with hydroxychloroquine, gold thioglucose, and D-penicillamine. D-Penicillamine

had to be discontinued twice owing to dermatitis. In August 1985 this drug had to be stopped permanently on account of leucopenia ($2.4 \times 10^9/l$) and thrombocytopenia ($60 \times 10^9/l$). Other abnormal laboratory findings were a positive antinuclear antibody (ANA) test and on immunoblotting weak anti-Sm and antihistone antibodies. Physical examination was normal and there were no signs of active arthritis. Drug induced systemic lupus erythematosus was suspected, and she was treated with prednisone for several months.

Because of active erosive RA involving wrists, knees, and proximal interphalangeal joints of the hands treatment with azathioprine 100 mg daily was started on 12 February 1987. Except for indomethacin 150 mg daily she took no other drug. Three weeks after starting azathioprine RA activity was diminished. Blood tests showed leucopenia of $1.6 \times 10^9/l$ and thrombocytopenia of $78 \times 10^9/l$. Azathioprine was immediately stopped. Five days later the platelet count fell to $27 \times 10^9/l$. Petechiae developed on her legs and she was admitted to hospital. Examination showed little RA activity. Laboratory findings were erythrocyte sedimentation rate (ESR) 60 mm/1st h, haemoglobin 90.2 g/l, platelets $18 \times 10^9/l$, leucocytes $1.8 \times 10^9/l$ with 47% granulocytes. Mean corpuscular volume (MCV) was 81 fmol/l, reticulocytes 0.2%. Except for a positive ANA test and anti-double-stranded DNA (anti-dsDNA) of 57 U/l (normal less than 20 U/l), other laboratory tests, including lactic dehydrogenase, direct Coombs', antibodies to platelets and granulocytes, complement, and immunoblotting, showed no abnormalities.

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HLA analysis was A2 A28 B15 Bw4 Bw6, DR1, DQw1. A bone marrow biopsy showed a hypocellular marrow with a shift of the myeloid series towards immaturity. Megakaryocytes were absent and the erythroid precursors were partly megaloblastic.

The pancytopenia was considered most probably to be related to azathioprine. From the first day in hospital indomethacin was also stopped. Treatment with prednisone 60 mg daily was started and reduced in 10 days to a 5 mg maintenance dose. Two days after admission platelet transfusion was necessary because of a rapid fall of the platelets to $4 \times 10^9/l$. Otherwise there were no other problems. The lowest value of the leucocytes was $1.4 \times 10^9/l$ and of haemoglobin 75.7 g/l (Fig. 1).

After three weeks in hospital the patient was dismissed with a haemoglobin of 90.2 g/l, reticulocytes 4.5%, leucocytes $2.7 \times 10^9/l$, and platelets $173 \times 10^9/l$. Later a marrow biopsy specimen was completely normal and anti-dsDNA was repeatedly

negative. At this time she felt well. Her only treatment being prednisone 5 mg daily.

CASE 2

A 68 year old man was diagnosed as suffering from seropositive RA in January 1976. He was then treated successively with hydroxychloroquine and gold thioglucose. Because of a recurrent duodenal ulcer he received a maintenance dosage of cimetidine 400 mg daily. In January 1985 hyperuricaemia and nephrolithiasis from urate calculi were detected, and treatment with allopurinol 100 mg daily was started. One year later he was admitted to hospital because of active RA involving wrists, knees, proximal interphalangeal and metacarpophalangeal joints of the hand. Subcutaneous nodules were present. Abnormal laboratory findings were ESR 81 mm/1st h, haemoglobin 114.3 g/l, creatinine 118 $\mu\text{mol/l}$, and a creatinine clearance of 70 ml/min. Treatment with azathioprine 100 mg daily was started. In addition to allopurinol and cimetidine, the patient had taken indomethacin 125 mg daily for several years. At the end of February 1986 a declining haemoglobin (82.1 g/l) occurred without melena or positive benzidine reaction. Values of leucocytes and platelets were normal. Gastroscopy showed superficial erosions in the bulbus. Two weeks later the patient was readmitted with pancytopenia. Physical examination was unchanged. Laboratory results were as follows: ESR 122 mm/1st h, haemoglobin 69.2 g/l, MCV 90 fmol/l, reticulocytes 0.1%, platelets $50 \times 10^9/l$, leucocytes $2.8 \times 10^9/l$ with normal differential cell count. Renal function was unchanged. Other laboratory values were normal, including iron, folic acid, and vitamin B₁₂. Bone marrow biopsy showed only a decreased number of megakaryocytes. After discontinuation of azathioprine, allopurinol, and indomethacin the counts of leucocytes and platelets returned to normal in two weeks. Two units of packed red blood cells were administered. Because of active RA azathioprine was restarted three weeks after admission at a dosage of 50 mg but now without allopurinol. Daily blood tests were performed. The dosage was increased after one week to 100 mg, but the drug had to be discontinued owing to isolated thrombocytopenia of $107 \times 10^9/l$. As the RA activity persisted, prednisone 10 mg daily was started and the platelet count became normal within one week.

Fig. 1 Haematological changes during azathioprine and indomethacin treatment in patient 1. (▼ = one unit of platelet transfusion.)

Discussion

These two patients with RA developed pancytopenia induced by azathioprine. Bacon *et al* have reported two patients with this serious side effect. They emphasised in one patient the possible addi-

tional role of allopurinol. In our second patient also allopurinol probably played a part in the development of pancytopenia.

Azathioprine is an imidazole derivative of 6-mercaptopurine with immunosuppressive and anti-inflammatory effects. Its mode of action is based on incorporation of a false purine, leading to alteration and inhibition of the purine, DNA, and RNA synthesis. Haematological side effects of azathioprine are dependent on the dosage. When the usual dosage (1.5–2.5 mg/kg/day) is given the interval before maximum bone marrow suppression is about two weeks.⁵ We found the same interval in patient 1.

Leucopenia is the most frequent haematological complication of azathioprine treatment. In published reviews about this adverse drug reaction in patients with RA the incidence of leucopenia ranges from 11 to 14.5%.^{1,2} Agranulocytosis within one week of azathioprine administration is very rare and possibly an idiosyncratic reaction.⁵ Other haematological changes are thrombocytopenia, anaemia, and macrocytosis. Thrombocytopenia is an uncommon side effect reported in 0.8–4.8% of cases.^{1,2} It rarely develops without leucopenia. In our second patient isolated thrombocytopenia developed two weeks after restarting azathioprine. Pure red cell aplasia induced by azathioprine is a very rare adverse reaction. It reflects a selective toxic effect of azathioprine on erythropoiesis.^{6,7} Macrocytosis has been reported in transplant recipients and ascribed to interference with DNA synthesis.^{7–9} It is unrelated to the metabolism of folic acid or vitamin B₁₂, or both. In our patients we could not find a significant alteration in the MCV. In the second patient a declining haemoglobin concentration occurred two weeks before the leucopenia and thrombocytopenia. When the negative result of the gastroscopy and the laboratory values are considered the anaemia appears to be related to azathioprine.

In the case of patient 1 the general practitioner had not brought to our notice the haematological changes (leucocytes $3.3 \times 10^9/l$, platelets $123 \times 10^9/l$) a week before she visited us. At that time we could have discontinued or diminished the dosage. An additional toxic effect of indomethacin on the bone marrow was unlikely because she had taken that drug for many years. In retrospect, owing to the absence of megakaryocytes from the bone marrow and the absence of antibodies to granulocytes and platelets, the indications for administering high dosage prednisone were debatable. As she had developed leucopenia and thrombocytopenia while taking D-penicillamine in the past she may have been predisposed to bone marrow suppression due to different drugs.

There are several reports of pancytopenia due to the combination of azathioprine and allopurinol,^{10–12} as occurred in patient 2. Azathioprine is rapidly converted to 6-mercaptopurine. The major metabolite of 6-mercaptopurine, 6-thiouric acid, is excreted in the urine. The enzyme xanthine oxidase is responsible for this oxidation. Allopurinol inhibits this enzyme and so administration of azathioprine together with allopurinol gives rise to an increased 6-mercaptopurine blood concentration, which can induce bone marrow toxicity. If it is necessary to administer both drugs it is advisable to reduce the usual dose of azathioprine by about 75%.^{3,5} In patient 2 this rule was unfortunately not applied.

Finally, when azathioprine is prescribed, close clinical and laboratory controls are necessary. Increased toxicity due to additional drugs has to be considered. Fortunately, both these patients recovered completely from this rare but dangerous adverse reaction to azathioprine.

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References

- Whisnant J K, Pelkey J. Rheumatoid arthritis: treatment with azathioprine (Imuran(R)). Clinical side-effects and laboratory abnormalities. *Ann Rheum Dis* 1982; **41** (suppl): 44–7.
- Bunch Th W, O'Duffy D M. Disease modifying drugs for progressive rheumatoid arthritis. Series on pharmacology in practice. *Mayo Clin Proc* 1980; **55**: 161–79.
- Lawson D H, Lovatt G E, Gurton G S, Hennings R C. Adverse effects of azathioprine. *Adverse Drug React Acute Poisoning Rev* 1984; **3**: 161–71.
- Bacon B R, Treuhaf W H, Goodman A M. Azathioprine-induced pancytopenia. *Arch Intern Med* 1981; **141**: 223–6.
- Decker J L, Steenberg A D. Immunoregulatory drugs. In: McCarty D J, ed. *Arthritis and allied conditions*. Philadelphia: Lee and Febiger, 1985: 529–30.
- Old C W, Flannery E P, Grogan Th M, Stone W H, San Antonio R P. Azathioprine induced pure red blood cell aplasia. *JAMA* 1978; **240**: 552–4.
- Mcgrath B P, Ibels L S, Raik E. Erythroid toxicity of azathioprine. *Q J Med* 1975; **173**: 57–63.
- Nicholls A J, Davidson R J L. Development of macrocytosis during azathioprine therapy after renal transplantation. *Transplantation* 1978; **27**: 220–1.
- Wickramasinghe S N, Dodsworth H, Rault R M J. Observations on the incidence and cause of macrocytosis in patients on azathioprine following renal transplantation. *Transplantation* 1974; **18**: 443–6.
- Reinicke H, Weber F W, Hausamen T U. Myelosuppression mit Makrozytärer Anämie und Leukopenie unter Behandlung mit Azathioprine bei Myasthenia gravis. *Dtsch Med Wochenschr* 1982; **107**: 1592–4.
- Zazgornik J, Kopsa H, Schmidt P, Pils P, Kusch K, Deutsch E. Increased danger of bone marrow damage in simultaneous azathioprine-allopurinol therapy. *Int J Clin Pharmacol Ther Toxicol* 1981; **19**: 96–7.
- Glogner P, Henri N. Pancytopenie nach Kombinationsbehandlung mit Allopurinol und Azathioprine. *Medizinische Welt* 1976; **277**: 1545–6.