LETTERS TO THE EDITOR

Worsening of Felty's syndrome with methotrexate

Sir: Drs Hughes and Abdulla recently reported a favourable response to low dose oral methotrexate treatment in a patient with Felty's syndrome,1 confirming earlier reports.2 3 In contrast, we found such treatment failed, with worsening of neutropenia.

A 55 year old man had had rheumatoid arthritis since 1980. He had received second line drug treatment, including gold compounds and D-penicillamine, which were discontinued because of proteinuria and inefficacy respectively. Haematological signs of toxicity were not noted with these treatments. A diagnosis of Felty's syndrome was established in 1988 when he developed a splenomegaly associated with polyarthritis and was found to have neutropenia (white blood cells 2.41×109/l; neutrophils 0.72×10⁹/l). Prednisone treatment 30 mg daily was started, then decreased to 20 mg/daily. In December 1989 he presented with acute polyarthritis and low peripheral neutrophils at 1.76×10⁹/l. The erythrocyte sedimentation rate was 45 mm/h. Methotrexate 7.5 mg weekly was introduced, together with 15 mg prednisone daily. After one month no marked amelioration of clinical condition was seen, the spleen was unchanged, white blood cells control showed a drop of neutrophils to 1.036×10^9 /l (WBC 2.8×10^9 /l, platelets 231×10⁹/l). Methotrexate was discontinued. Further follow up showed that after one month neutrophils had increased and stabilised. Treatment with corticosteroids was continued alone.

In this patient methotrexate treatment failed to improve Felty's syndrome, and, on the contrary, induced a transient worsening of neutropenia. His condition was well defined,4 and no previous history of toxic neutropenia was found. He was not receiving any other neutropenic drug. His condition did not apparently differ from those of other patients who improved with methotrexate treatment. 1-3 Our data suggest individual sensitivity to methotrexate treatment in Felty's syndrome, and further studies are needed to define the characteristics which would predict a positive response.

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Lymphoedema complicating rheumatoid

Sir: I read with interest the report of Dacre,





Wrist radiographs taken in two planes.

Scott, and Huskisson on lymphoedema as an extra-articular feature of rheumatoid arthritis. 1 The phenomenon of lymphoedema of a limb complicating rheumatoid arthritis is well recognised, and I have successfully treated two patients with unilateral hand and forearm swelling with daily compression methods (Flowtron apparatus).

The article also reminded me of another interesting patient seen a few years ago with an identical appearance of one hand and wrist to that seen in the illustration accompanying the article. This patient had a wrist arthrogram performed, and the films show the resulting radiographs taken in two planes, clearly

showing the dye penetrating the marrow cavity of the radius. This patient's oedema was rather refractory to treatment. I am unaware of this marrow penetration phenomenon being previously reported.

I thank Dr J D Jessop, University Hospital of Wales, Cardiff, for permission to reproduce the radiographs.

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1 Dacre J E, Scott D L, Huskisson E C. Lymphoedema of the limbs as an extra-articular feature of rheumatoid arthritis. Ann Rheum Dis 1990; 49: 722-4.

Smoking and back pain

Sir: The recent study by Symmons and coworkers¹ confirms previous work showing an association between smoking and back pain. The authors also (for the first time) show a similar association between back pain and oral contraceptives. These two findings might have a common mechanism as both smoking and oral contraceptives lead to disturbances in the flow properties of blood. Blood viscosity is a suitable ex vivo parameter for measuring this. It is roughly 50% higher in non-smokers than in heavy smokers (23.3 (SD 8.0) v 31.2 (9.7) mPa.s). When oral contraceptives (levonorgestrel) are taken by young healthy women for three cycles blood viscosity increases on average by 3 mPa.s.³
One might therefore speculate⁴ that this alteration in blood rheology leads to a malnutrition of the highly bradytrophic intervertebral disc, rendering it more vulnerable to injury. This would be an attractive explanation of the fact that these cardiovascular risk factors are also related to back problems. It seems tempting to test this speculation in more detail—for example, by reanalysing some of the numerous epidemiological studies on cardiovascular disease in terms of back problems.

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1 Symmons D P M, van Hemert A M, Vanden-broucke J P, Valkenburg H A. A longitudinal study of back pain and radiological changes in the lumbar spines of middle aged women. I. Clinical findings. Ann Rheum Dis 1991; 50: 158 61

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2 Ernst E, Matrai A, Schmölzl Ch, Magyarosy I.
Dose-effect relationship between smoking and blood rheology. Br J Haematol 1987; 65:

3 Ernst E, Schmölzl Ch, Matrai A, Schramm W.

Hemorheological effects of oral contraception.

Contraception 1989; 40: 571-80.

4 Ernst E. Risk factor for back trouble. Lancet 1989; i: 1305-6.

Rheumatology in Dar-es-Salaam, Tanzania

Sir: May I commend and support the views expressed by Dr Adebajo. 1 During a spell teaching at Muhimbili Hospital, Dar-es-Salaam, Tanzania, I found a dearth of rheumatological experience amidst a wealth of clinical material. The table shows those disorders encountered during two weeks, mainly by scouting through clinics and wards. Good-