Clotrimazole in rheumatoid arthritis

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SUMMARY Forty-seven patients with active rheumatoid arthritis took part in an 8-week controlled study in which clotrimazole was compared with a standard nonsteroidal anti-inflammatory agent, ketoprofen. Although clotrimazole was shown to be effective in the treatment of the disease and superior to ketoprofen in certain measurements, it was also responsible for a high incidence of adverse effects. Improvement with clotrimazole took place more slowly but was more sustained than with ketoprofen. A significant rise in plasma cortisol and a fall in white cell count was observed in the clotrimazole treated patients.

Clotrimazole is a tritylimidazole derivative which has been used chiefly as a broad spectrum antimycotic agent.¹ On the assumption that a protozoon might play a major role in the pathogenesis of rheumatoid disease the drug was used to treat a number of patients suffering from this condition. The results of an uncontrolled study were extremely encouraging, with active disease reported to disappear within 1 month of starting treatment and spectacular improvement observed in practically every case.² It was because of these findings and the fact that other imidazole derivatives had been shown to improve rheumatoid arthritis³⁻⁵ that the present controlled study was embarked upon.

Patients and methods

Forty-seven patients suffering from definite or classical rheumatoid arthritis as defined by the American Rheumatism Association⁶ were admitted into the study. Patients with peptic ulceration, renal or hepatic insufficiency, diabetes, or any other serious medical disorder were excluded. Patients on corticosteroids or on antirheumatic drugs such as gold and penicillamine and on immunosuppressive therapy were not considered for entry into the study. The trial was double-blind. Two parallel groups of patients were randomly allocated to treatment with either clotrimazole or ketoprofen, a propionic acid derivative of proved efficacy in the treatment of rheumatoid arthritis.^{7,8} The double dummy technique was used. The daily dose of clotrimazole was 40 mg per kg weight during the first week,

increasing to 80 mg per kg given in divided doses. The dose of ketoprofen was 50 mg twice daily for 1 week, increasing to 50 mg 3 times a day. The total treatment period was 8 weeks. Because oral clotrimazole was known to be poorly tolerated^{9,10} all patients were observed in hospital for the first fortnight of treatment but not confined to bed. All antirheumatic therapy was discontinued on admission, and only paracetamol was allowed as the 'rescue' analgesic. Paracetamol consumption was recorded throughout the trial period.

Clinical and laboratory measurements were made at the beginning of the trial and these were repeated at weekly intervals throughout the study. These included proximal interphalangeal joint circumference,¹¹ duration of morning stiffness, grip strength, articular index,¹² visual analogue pain assessment, and the patient's total assessment of treatment, as well as a full blood count, platelet count, ESR, Rose-Waaler test, C-reactive protein, blood urea, standard liver function tests, immunoglobulins, and plasma cortisol. The assessment took place each week at approximately the same time in the morning. The 2 groups were well matched for age, sex, and duration of their disease. Student's *t* test was used to compare differences between changes in the treatment groups.

Results

Seventeen of the 24 patients in the clotrimazole group and 20 of the 23 patients in the ketoprofen group completed the study. All 7 withdrawals from the clotrimazole group were caused by gastrointestinal intolerance, a feature which accounted for only 1

withdrawal from the ketoprofen group. The remaining 2 withdrawals from the latter group were due to flare-up of the disease.

Significant improvement of most clinical measurements was noted in both groups of patients (Table 1), but the rate at which this improvement took place differed between the groups. It was faster in the ketoprofen group, with maximum progress being made during the first 4 weeks, thereafter followed by a steady decline. Improvement in the clotrimazole

group was delayed, with many cases making no progress in the first week. However, by the end of the first month this group was beginning to fare better, and superiority became more marked during the second month.

The majority of measurements showed no statistical difference between the groups, although the trend in favour of the clotrimazole treated patients was a feature during the second month of the study. The exception was the consumption of 'rescue'

Table 1 Changes in clinical measurement

		Grip strength (mm)	Pain (VAS)	Articular index (Ritchie)	Joint swelling (mm)	Duration of morning stiffness (hours)	Patients total assessment (5 point scale - =improved)
Week 1	Clotrimazole	+2.3	-0.9	-1.1	-7.9†	-0.2	-0.1
	Ketoprofen	+7.25	-1.0†	-3.45†	-3.2	-0.1	-0.2
Week 4	Clotrimazole	+58.8†	-2.6†	-9.1†	-18.9*†	-0.65†	-1.0†
	Ketoprofen	+50.0†	-2.0†	-7.0†	-7.05†	-0.2	-0.55†
Week 8	Clotrimazole	+74.4†	-2.6†	-9.2†	-23.1†	-0.5	-0.8†
	Ketoprofen	+23.3	-1.3†	-5.25†	-14.6†	-0.2	-0.3

† Significant improvement from pre-entry, $P < 0.05$. * Significant difference between groups, $P < 0.05$.

Fig. 1 'Rescue' drug count.

Fig. 2 Plasma cortisol (mean ± SEM).

drug, which became significantly lower in the clotrimazole treated patients during the second month of the study and, in contrast to the ketoprofen group, was still decreasing at the conclusion of the study (Fig. 1). Adverse effects were substantially commoner in the clotrimazole treated patients, with gastrointestinal symptoms predominating (Table 2). Lethargy, drowsiness, and pain on micturition were also noted by some patients in this group.

Little change was observed in most of the laboratory values (Table 3). In the clotrimazole treated group, however, a marked rise in plasma cortisol (Fig. 2) and a significant fall in white cell count

(Fig. 3) involving predominantly the polymorphs was seen. In no case, however, was leucopenia observed.

Discussion

The results of this controlled study fail to demonstrate the dramatic improvement which has previously been described in patients with rheumatoid arthritis taking clotrimazole.² In the present study the improvement in the clotrimazole treated group was slower but more sustained than that in patients who received treatment with ketoprofen. The extent of improvement leaves us in no doubt as to the efficacy of clotrimazole in the treatment of rheumatoid arthritis. Previously reported work suggested that a 2-month treatment period was sufficient to observe optimum therapeutic effect, and the design of the current study was drawn up with this in mind. However, the fact that after 2 months' treatment improvement was still continuing suggests that longer periods of treatment may be necessary before maximum benefit is observed. Our observations of cases treated for up to 8 months in an uncontrolled study tend to substantiate this, and

Table 2 Main adverse effects

Complaint	Number of patients		
	Clotrimazole	Ketoprofen	
Nausea/vomiting	16	6	P<0.02
Diarrhoea	14	5	P<0.025
Anorexia	10	0	P<0.002
Anorexia	10	0	P<0.002

Table 3 Changes in laboratory values

		C-reactive protein (Dilution)	ESR (mm/h)	Urea (mmol/h)	Alkphos (U/L)	SGOT (U/L)	LDH (U/L)	IgG (IU/ml)	IgM (IU/ml)	IgA (IU/ml)
Week 1	Clotrimazole	—	+6.3	-0.5*	+4.4	+3.2	-1.4	—	—	—
	Ketoprofen	—	-1.7	+0.8*	-1.4	+0.9	-0.3	—	—	—
Week 4	Clotrimazole	-0.07	+9.8	-0.7*	+7.9†	+3.4	+21.5	—	—	—
	Ketoprofen	-0.18	-0.1	+0.7*	+7.9†	-0.2	+5.5	—	—	—
Week 8	Clotrimazole	-0.64*	+11.5	-1.5*†	+7.3	+1.8	+14.0	-24.4†	+8.8	-31.4
	Ketoprofen	+0.28*	+9.9	+0.4*	+5.2	+0.5	+15.9	-5.8	+45.3†	-163.6

† Significant change from pre-entry, P<0.05. * Significant difference between groups, P<0.05.

Fig. 3 White cell count (mean ± SEM).

SIGNIFICANT CHANGE IN CLOTTRIMAZOLE GROUP P<0.01 FROM WEEK 5 ONWARD
SIGNIFICANT DIFFERENCE BETWEEN GROUPS P<0.05 FROM WEEK 3 ONWARD

suggest that maximum clinical improvement occurs between 3 and 6 months and is accompanied by a fall in the ESR. Certainly experience with imidazole derivatives, such as levamisole, indicates that full benefit may not take place until 6 months after treatment is begun.

The mode of action of clotrimazole in rheumatoid arthritis is open to debate. It has been suggested on the basis of animal studies that this drug might exert an anti-inflammatory effect by stimulating the adrenal glands.¹³ The raised levels of plasma cortisol that were so consistently observed in the patients on clotrimazole would certainly support this suggestion.

An in-vitro study¹⁴ has shown that the addition of clotrimazole to normal lymphocytes causes both an inhibitory and an enhancing effect on mitogenic stimulation. This was dependent on the concentration used, but a predominantly immunosuppressive effect was observed at concentrations equivalent to therapeutic serum levels. A subgroup of rheumatoid patients in the present study also showed a significant reduction in mitogenic lymphocyte responsiveness while on clotrimazole, whereas no change occurred in patients taking ketoprofen.¹⁴ Although it is possible that this finding related to the increased cortisol production in the clotrimazole group, the of correlation with cortisol concentration and the existence of suppression despite the absence of autologous serum suggest an inherent immunosuppressive action of clotrimazole. This contrasts with the immunopotentiating effect of levamisole.

On the basis of our results we would suggest that clotrimazole affects rheumatoid arthritis in 2 ways. The initial effect, though not immediate, appears to take place sooner than is normally seen in patients who respond to immunosuppressive drug therapy for rheumatoid arthritis and may be the result of increased cortisol release by adrenal stimulation. The fact that plasma cortisol increased gradually during the first 3 weeks before reaching a steady level would lend support to this suggestion. The continued improvement during the latter part of the study and associated with the gradual fall in white cell count could be attributed to the immunosuppressive action of the drug.

What role, if any, is clotrimazole likely to play in the management of rheumatoid arthritis? The positive features of the present study are more than counterbalanced by the unacceptably large

number of side effects. If, as has been suggested,² treatment with a substantially lower dose of clotrimazole were to prove equally effective without the disadvantage of a high degree of intolerance, then clotrimazole could yet play a role as an alternative to some of the more specific drugs currently used in the treatment of the more severe and resistant cases of rheumatoid arthritis. On the other hand we suggest that investigation of other similar but potentially less toxic oral antimycotic agents for the antirheumatic properties which clotrimazole has been shown to possess might prove a more useful area of research.

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References

- ¹ *Br Med J* 1969; **4**: 444-5.
- ² Wyburn-Mason R. Clotrimazole in rheumatoid arthritis. *Lancet* 1976; **1**: 489.
- ³ Abd-Rabbo H, Abaza H, Hillal G, Moghazy M, Asser L. Nitro-imidazole in rheumatoid arthritis. *Am J Trop Med Hyg* 1972; **75**: 64-6.
- ⁴ Schuermans Y. Levamisole in rheumatoid arthritis. *Lancet* 1975; **1**: 111.
- ⁵ Huskisson E C, Dieppe P A, Scott J, *et al.* Immuno-stimulant therapy with levamisole for rheumatoid arthritis. *Lancet* 1976; **1**: 393-5.
- ⁶ Ropes M W, Bennett G A, Cobb S, Jacox R, Jessar R. Diagnostic criteria for rheumatoid arthritis, 1958 Revision. *Ann Rheum Dis* 1959; **18**: 49-53.
- ⁷ Gyory A N, Bloch M, Burry H C, Grahame R. Orudis in management of rheumatoid arthritis and osteoarthritis of the hip; comparison with indomethacin. *Br Med J* 1972; **4**: 398-400.
- ⁸ Zutchi D W, Marr S, Bloch M, Mason R M. Double-blind cross-over study of orudis and placebo in 50 patients with rheumatoid arthritis. *Rheumatol Rehabil* 1973; **12**: 62-7.
- ⁹ Tetterborn D. Toxicity of clotrimazole. *Postgrad Med J* 1974; Suppl. 17-20.
- ¹⁰ Bayer A G. Clotrimazole. *Investigators Manual* 1974; 43-59.
- ¹¹ Boardman P L, Hart F D. Clinical measurement of the anti-inflammatory effects of salicylates in rheumatoid arthritis. *Br Med J* 1967; **4**: 264-8.
- ¹² Ritchie D M, Boyle J A, McInnes J M, *et al.* Clinical studies with an articular index for the assessment of joint tenderness in patients with rheumatoid arthritis. *Q J Med* 1968; **37**: 393-406.
- ¹³ Otterness I G, Niblack J R. Clotrimazole and rheumatoid arthritis. *Lancet* 1976; **1**: 148.
- ¹⁴ Gow P J, Corrigan V, Panayi G S. The effects of clotrimazole on human lymphocyte responsiveness to plant mitogenes. *Agents Actions* in press.