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Observations on Drug Prescribing in Rheumatoid Arthritis

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I firmly believe that if the whole material medica, as now used, could be sunk to the bottom of the sea, it would be all the better for mankind, and all the worse for the fishes.—OLIVER WENDALL HOLME.¹

Summary

A total of 125 patients with rheumatoid arthritis were investigated about their drug therapy before referral to a specialist centre. Most referrals were from general practitioners. Only 47 of the patients had received salicylates as the first drug and 18 had never had them at all. Soluble aspirin was the preparation of salicylates most frequently prescribed (for 63 patients). Only 60 patients had been given an adequate dose and only 62 an adequate course of treatment with salicylates. In 28 patients salicylates had been stopped on account of side effects. About one-third of the patients had been prescribed oral corticosteroids.

The referral letters were poor in giving details of past and present drug therapy, and there were serious omissions in reporting of previous side effects.

Seventy-five general practitioners were asked to rate several currently marketed antirheumatic drugs in terms of effectiveness. Though prednisolone 15 mg daily ranked higher than aspirin 4 g daily the difference was not significant. The study shows the inadequacies of drug prescribing for rheumatoid arthritis in the Glasgow area.

Introduction

Many analgesic and anti-inflammatory drugs may now be prescribed to relieve pain in rheumatoid arthritis. Nevertheless, the "sheet-anchor" of treatment is still salicylates.²⁻⁵ We have found that many patients referred for consultation have not received salicylates as their first drug and many have never received them at all. We have therefore reviewed drug treatment in new referrals to this clinic and the information regarding drug

therapy given in the referral letter. The results indicate an unsatisfactory state of drug prescribing in patients with rheumatoid arthritis in Glasgow.

Patients and Methods

We interviewed 125 patients, who were seen consecutively by one doctor, about their drug therapy before they were referred to the centre. The mean age of the patients was $50.6 \pm \text{S.D. } 15.6$ years (range 13 to 82 years) and 93 were female. All had "definite" or "classical" rheumatoid arthritis according to the diagnostic criteria of the American Rheumatism Association.⁶ The mean duration of arthritis was 7.2 ± 7.3 years, with a range from one month to 31 years.

Each patient was questioned about the first drug prescribed for his arthritis, its dose and duration, and whether it gave rise to side effects; similar details were obtained for all other drugs prescribed. When the patient was uncertain about what drug he had been given he was asked to identify it from a display of available antirheumatic agents. When doubt still existed a letter was sent to the family doctor to verify the preparation given. In 13 patients the first drug prescribed could not be ascertained with certainty, and these patients were not included in the study. We also noted the source of patient referral and whether the accompanying letter mentioned previous and current drug therapy and its dosage—also whether side effects had resulted.

We recorded the preparation of salicylate which had been prescribed and whether salicylates were prescribed in proper dosage—that is, 3 g or more per day—and whether this was given a proper trial—three months or more. We noted also the reason for discontinuing salicylate therapy, any self-medication of analgesics, and the preparation and dosage of corticosteroid therapy.

To determine opinions on the effectiveness of available antirheumatic drugs we sent a questionnaire to 75 general practitioners in Glasgow chosen at random from the General Medical Service list of medical practitioners. They were asked to rate the effectiveness of a number of preparations on a five-point scale: totally ineffective 1, ineffective 2, moderately effective 3, effective 4, and highly effective 5 (see table VI). Forty-nine replies were received. A further 250 general practitioners were asked whether they approved of a proposed standard regimen of drug treatment consisting of initial treatment with salicylates 3.5-4 g daily. Should the patient fail to tolerate or respond to salicylates indomethacin up to 150 mg daily, benorylate 10 ml twice a day, or phenylbutazone 300 mg daily could be given. If this treatment failed then gold or penicillamine, corticosteroids, or immunosuppressive agents might be given but only in consultation with a rheumatologist. As adjuncts to treatment we recommended ibuprofen 1,200 mg

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daily, Distalgesic (dextropropoxyphene 32.5 mg and paracetamol 325 mg), or mefenamic acid 1,500 mg daily.

Results

Most patients were referred by their general practitioner, 69 coming from this source, while 22 were referred by a consultant physician, 14 by a consultant surgeon, and 20 by a medical registrar. Only 47 patients had first received salicylates and 18 patients had not received them at all before they were referred for consultation (table I). The most popular type of aspirin prescribed was soluble aspirin (table II). Only 60 patients out of the 94 who were given salicylates were prescribed what we considered to be an adequate dose of salicylates—3 g per day or more—and only 62 had had an adequate course of treatment—three months or longer. Therapy had had to be discontinued on account of side effects in 28 patients—in 21 because of dyspepsia. Ten patients had discontinued salicylate therapy on advice from their family doctor that the drug was dangerous. Two of these patients, however, had had duodenal ulceration (table III).

The most frequently used oral preparation for systemic corticosteroid therapy was prednisolone (21 patients), and

tetracosactrin depot (six patients) was more commonly prescribed than natural corticotrophin (four patients; table IV). Excessive dosage of oral corticosteroid therapy—that is, more than 10 mg prednisolone equivalent—had been prescribed in 19 patients.

TABLE IV—Systemic Corticosteroid Therapy in 48 Patients with Rheumatoid Arthritis

Systemic Corticosteroid Therapy	No. of Patients
Preparation:	21
Prednisolone	2
Prednisone	3
Triamcinolone	2
Methyl prednisolone	3
Dexamethasone	2
Betamethasone	1
Sintison	7
Unknown	4
Corticotrophin	6
Tetracosactrin depot	3
Two or more preparations	
Dose:	
Excessive (>10 mg prednisone equivalent)	19
Undetermined	3

TABLE I—Initial Treatment and Other Drugs Prescribed for 112 Patients with Rheumatoid Arthritis before Referral to Specialist Centre

	First Drug	Other Drugs
	No.	No.
Salicylates	47	94
Paracetamol	3	44
Indomethacin	22	83
Phenylbutazone	12	43
Oxyphenbutazone	1	4
Ibuprofen	8	40
Flufenamic acid		6
Mefenamic acid		3
Benorylate		2
"Safapryn"		2
Alclofenac		3
"Distalgesic"	2	4
Dihydrocodeine	1	5
Corticosteroids:		
Oral	5	36
Intra-articular	1	2
Corticotrophin		4
Tetracosactrin depot		8
Gold	1	6
No treatment	2	
Total	113*	

*One patient started on both salicylates and indomethacin.

TABLE II—Types of Aspirin Prescribed in 94 Patients with Rheumatoid Arthritis Treated with Salicylates

Aspirin Formulation	No. of Patients
Aspirin B.P.	22
Soluble	63
Enteric coated	21
Aloxiprin (calcium)	1
Unknown	2
Two or more formulations prescribed	14

TABLE III—Dosage, Duration, and Reasons for Stopping Therapy in 94 Rheumatoid Arthritic Patients Treated with Salicylates

Salicylate Therapy	No. of Patients
Dose:	
Adequate (≥ 3 g/day)	60
Inadequate	34
Course:	
Adequate (≥ 3 mnths)	62
Inadequate	32
Reason for stopping:	
Side effects	28
Dyspepsia	21
Gastrointestinal bleeding	2
Other*	5
General practitioner's advice†	10
Ineffective	15

*This comprised two patients with rash, one with dizziness, one with tinnitus, and one with constipation.

†These included two patients with duodenal ulceration.

Of 639 drugs which had been prescribed only 107 were mentioned in the referral letter (table V). Fifty-one patients had suffered between them 85 definite side effects, but only three side effects were recorded in the letter. Four serious side effects were omitted from the letters: haematemesis on aspirin; severe depression on ibuprofen; agranulocytosis on phenylbutazone; and a peptic ulcer on phenylbutazone. Dosage of past drug therapy was mentioned for only 15 patients in 22 of the prescribed drugs. The details of current therapy showed much the same pattern of documentation. Of 160 drugs currently being prescribed only 56 were mentioned in the letter. The dosage of these drugs was recorded only in 20 instances for 17 patients, and only four out of 19 side effects were mentioned.

TABLE V—Past and Current Antirheumatic Drug Therapy in 112 Patients mentioned by the Patients and their Referring Doctors. Figures in Parentheses are Numbers of Patients

	Mention by Patients	Mention by Referring Doctor
Past drug therapy:		
No. of drugs	369	107 (54)
Mean no. of drugs per patient $\pm$ S.D.	3.0 ± 1.7	
No. of dosages		22 (15)
No. of side effects	85 (51)	3 (2)
Present drug therapy:		
No. of drugs	160	56 (46)
Mean no. of drugs per patient $\pm$ S.D.	1.3 ± 0.7	
No. of dosages		20 (17)
No. of side effects	19 (18)	4 (4)

Three patients were never prescribed antirheumatic drugs and seven were receiving no antirheumatic drugs at the time of study.

All the 49 general practitioners replying to our questionnaire considered that all the drugs were superior to placebo (table VI). Surprisingly, however, though prednisone in a daily dose of 15 mg was rated higher than aspirin 4 g per day the difference was not statistically significant.

The results of a questionnaire sent to 250 general practitioners in Glasgow asking about their prescribing habits showed that 25 of them did not consider salicylates as the drug of first choice. The reason was that patients in practice fail to tolerate high-dose salicylate therapy and most patients find low doses of salicylates ineffective. Of these 25 practitioners 20 first prescribed ibuprofen or indomethacin, two prescribed phenylbutazone, and three prescribed oral corticosteroids.

Discussion

This study was designed to obtain information about prescribing practices in Glasgow for patients with rheumatoid

TABLE VI—Effectiveness of Various Antirheumatic Drugs according to Opinions of 49 General Practitioners. Mean Effectiveness calculated on Five-point Scale (see Text)

Drug	Dose/Day	Effectiveness (Mean $\pm$ S.E. of Mean)	Statistical Significance		
			Comparison	t	p
A Prednisone	15 mg	4.2 $\pm$ 0.2	A v. B	1.04	N.S.
B Aspirin (soluble)	4 g	3.9 $\pm$ 0.1	B v. C	0.73	N.S.
C Phenylbutazone	300 mg	3.8 $\pm$ 0.2	B v. D	0.59	N.S.
D Prednisone	10 mg	3.8 $\pm$ 0.2	B v. F	2.67	<0.01
E Tetracosactrin depot	0.5* mg	3.8 $\pm$ 0.2	B v. H	2.51	<0.01
F Aspirin (enteric coated)	4 g	3.6 $\pm$ 0.1	B v. J	3.86	<0.0005
G Corticotrophin	12 tabs	3.6 $\pm$ 0.2	B v. L	9.12	<0.0005
H Indomethacin	100 mg	3.5 $\pm$ 0.1	J v. L	1.20	N.S.
I "Safapryn"	12 tabs	3.4 $\pm$ 0.1	A v. D	3.99	<0.0005
J Ibuprofen	1,200 mg	3.3 $\pm$ 0.1	D v. K	5.50	<0.0005
K Prednisone	5 mg	3.3 $\pm$ 0.2	D v. G	1.67	N.S.
L Aspirin (soluble)	2 g	3.1 $\pm$ 0.1	E v. G	1.46	N.S.
M Oxyphenbutazone	300 mg	3.1 $\pm$ 0.2	G v. A	0.97	N.S.
N Benorylate	20 ml	3.1 $\pm$ 0.2	E v. A	0.34	N.S.
O Alclofenac	3 g	2.9 $\pm$ 0.1	L v. Q	3.00	<0.0025
P Mefenamic acid	1,500 mg	2.8 $\pm$ 0.1	L v. P	2.10	<0.025
Q Paracetamol	4 g	2.7 $\pm$ 0.1			
R Flufenamic acid	600 mg	2.6 $\pm$ 0.2			
S Calcium lactate (placebo)	4 tabs	1.7 $\pm$ 0.1			

N.S. = Not significant.
* = Twice weekly dose

arthritis. The study was retrospective and, as such, suffers from the limitations of this type of investigation. Nevertheless, the results do highlight several disturbing features.

Despite the acceptance of salicylates as the drug of choice in rheumatoid arthritis,²⁻¹⁰ only 47 patients were first treated with this drug. In addition 18 patients had never received salicylate therapy at any time during their illness. Of those who had had salicylate therapy about one-third had not received an adequate course of treatment, in terms either of dosage or of duration of therapy. Their family doctors had told 10% of patients to discontinue treatment with salicylates because they considered them dangerous. In a poll of over 5,000 doctors in the United States only a quarter prescribed salicylates as the drug of first choice in rheumatoid arthritis.¹¹

The question arises Why do some family doctors not prescribe salicylates to their patients with rheumatoid arthritis? The answer may be that they consider that salicylates in high doses cause side effects whereas in low doses they are inadequate to relieve pain. In the 250 replies to our questionnaire 25 family doctors gave this as their reason for not prescribing salicylates as the drug of first choice. It is difficult to fault this answer since 28 patients in the present study had to discontinue salicylates because of side effects, mostly dyspepsia. We do not, however, agree with the two family doctors who considered phenylbutazone or the three doctors who preferred oral corticosteroids as the drug of first choice. Clearly several family doctors in Glasgow prescribe phenylbutazone and oral corticosteroids as the drugs of first choice in treating patients with rheumatoid arthritis (see table I). It would be interesting to know their views on the potential toxicity of these particular antirheumatic agents.

The high percentage of patients in this study who had been treated with systemic corticosteroids is surprising and runs contrary to present recommendations to use corticosteroid therapy as infrequently as possible. Many of the patients had severe rheumatoid arthritis, and it could be argued that corticosteroid therapy was justified but for the fact that most patients had not had a careful and unhurried trial of non-steroidal

analgesic drugs. A particular criticism is that 19 of those given oral corticosteroids received doses over 10 mg prednisolone per day or equivalent. In view of the popularity of corticosteroid therapy it is interesting that the doctors assessed the mean analgesic effect of 15 mg prednisolone as not being more effective than 4 g of soluble aspirin.

Only 18 patients in the present study admitted to self-medication, mostly as salicylates; only two patients said they were taking phenacetin-containing mixtures. It is impossible to know how accurate these data are, and they are probably an underestimate. Murray¹² found that 128 of 740 Glasgow citizens admitted to taking analgesics at least weekly, and most of these did so without reference to a doctor. Furthermore, the incidence of analgesic nephropathy in the West of Scotland is four times greater than that found in England and Wales.¹³

The referring doctor's recording of past and present drug therapy can hardly be described as adequate. The letters, however, were reasonably adequate about the patient's symptoms, past and family histories, physical findings, and results of laboratory data. Thus, for example, one doctor's letter which ran to two pages of typescript gave every detail of the patient's illness except what drugs had been and were being prescribed. This patient had had a haematemesis from a duodenal ulcer while receiving phenylbutazone. Hence the importance of drug therapy in the management of patients still lags far behind in most doctors' thoughts. Interestingly, there was no difference between details of drug therapy in letters from hospital doctors and family doctors, though letters from surgical colleagues lagged far behind in this matter. Clearly more education is needed in this important aspect of medical care.

Regional differences in drug prescribing have been noted both in general¹⁴ and in rheumatic disease.¹⁵ In our region there is, for example, little gold prescribed for rheumatoid arthritis, though in several areas in the United States it is extremely popular. It remains to be seen whether the inadequacies of drug prescribing which we have observed in Glasgow are similar in other parts of the world.

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