

Comparative trial of Co-trimoxazole and Chloramphenicol in Typhoid Fever

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Summary

A comparative trial of co-trimoxazole and chloramphenicol was conducted in two groups of 50 patients each to try to resolve conflicting opinions on the relative merits of the two drugs in the treatment of typhoid fever. We conclude that in our part of India co-trimoxazole is superior to chloramphenicol and that differences in our findings to those of others may perhaps be accounted for by differences in strains of *Salmonella typhi*, ethnic differences, and possibly differences in herd immunity to typhoid.

Introduction

Comparative clinical studies of co-trimoxazole (trimethoprim-sulphamethoxazole) and chloramphenicol in enteric fever in different parts of the world have given conflicting results, some indicating that co-trimoxazole is superior to chloramphenicol (Kamat, 1970) and others the opposite (Scragg and Rubidge, 1971). To try to resolve the issue we undertook a comparative clinical trial of the two drugs.

Patients and Methods

The trial was conducted in the Sassoon Hospitals, Poona, and was restricted to inpatients over 12 years of age for better control and standardization of dosage and follow-up.

The following investigations were carried out in patients suspected to be suffering from enteric fever: routine urine and blood examinations, blood-clot culture for salmonella, and Widal reaction. No specific therapy was given for 96 hours, by which time the result of the blood-clot culture was available. Only patients whose culture was positive for *Salmonella typhi* were admitted to the trial. They were allotted to one of two treatment groups on a randomized basis, the control group receiving chloramphenicol and the trial group receiving co-trimoxazole. Antipyretics were not given since they might have affected fever, one of the criteria for assessment. Intravenous fluids were given when indicated. The patients were examined daily.

The response to specific therapy was assessed on the following criteria: (1) the duration of fever; (2) the duration of toxic symptoms such as severe headache, mental confusion, disorientation, delirium, and involuntary movements; (3) exacerbation of toxæmia after starting specific therapy (toxic crises); (4) relapse of fever within three weeks of defervescence; and (5) occurrence of complications of typhoid fever.

Patients in the trial group were given four tablets of co-trimoxazole (each containing trimethoprim 80 mg and sulphamethoxazole 400 mg) twice daily until defervescence, and then two tablets twice daily for seven days. Patients in the control group were given chloramphenicol two capsules six-hourly until defervescence and then one capsule six-hourly for 14 days, each capsule containing 250 mg.

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There were 50 patients in each group, the trial group consisting of 40 males and 10 females and the control group of 37 males and 13 females. The mean age of patients in the trial group was 23.44 years (range 13-45) and in the control group 24.6 years (range 11-75). Their distribution among age groups is shown in table I.

TABLE I—Age-group Distribution of Patients

Age	No. in Trial Group	No. in Control Group
12-20	20	21
21-30	21	19
31-40	6	7
41-50	3	1
Over 50	—	2

Most patients had an axillary temperature on admission of from 100° to 102°F (37.8°-38.9°C) but in 11 in the trial group and in 13 in the control group it exceeded 102°F. The duration of fever before admission is shown in table II. The mean

TABLE II—Duration of Fever before Admission

Duration (Days)	No. of Patients	
	Trial Group	Control Group
1-3	4	7
4-6	16	14
7-9	16	18
Over 9	14	11

duration of fever was 6.98 days in the control and 7.53 days in the trial group. Thirty-two of the patients in the trial group and 36 of those in the control group were toxæmic at the time of starting specific therapy.

Results

All patients in both groups responded to treatment. There was no significant difference in the response of fever to treatment in either group. The mean period of fever before defervescence was 4.55 days in the control group and 4.98 days in the trial group. Toxæmia responded to treatment earlier in those treated with co-trimoxazole (trial group) than in those treated with chloramphenicol (control group) (table III).

TABLE III—Response of Toxæmia to Treatment

Duration of Treatment before Response (Days)	No. of Patients	
	Trial Group (n = 32)	Control Group (n = 36)
1-3	25	18
4-6	6	12
Over 6	1	6

A toxic crisis—drug-induced exacerbation of toxæmia—was seen in two patients in the control group but in none in the trial group. Two patients in the control group relapsed. One developed fever with a positive blood culture 17 days after completing a course of chloramphenicol treatment. He responded to co-trimoxazole in four days and had no further relapse. The other patient who relapsed developed fever and a positive

blood culture 16 days after completing a course of chloramphenicol. He also responded to co-trimoxazole in four days and had no further relapse. Both these patients may have been reinfecting, but far more probably they were genuine cases of relapse.

There were no serious complications such as haemorrhage, perforation, or circulatory collapse in any patient in either group. Nor were there any severe side effects. Minor side effects were equally common in both groups, the only notable difference being a significantly greater incidence of weakness in the control group (19 patients) compared with the trial group (5 patients).

Discussion

All patients in both groups in our series responded to specific therapy, thus confirming the results of Akinkugbe *et al.* (1968), Farid *et al.* (1970), and Kamat (1970). In contrast, two out of 23 patients in Geddes *et al.* (1971) series and 8 out of 103 in Scragg and Rubidge's (1971) series treated with co-trimoxazole failed to respond. Farid's (1971) observations on the dosage in Scragg and Rubidge's series that underweight and undernourished children who were given co-trimoxazole on a body-weight basis might have received less than a therapeutic dose is very pertinent and may explain Scragg and Rubidge's poor results with the drug. This also confirms our earlier experience (Sardesai, 1971) when we did not encounter a single failure in 40 patients treated with co-trimoxazole.

In this trial the response of toxæmia to treatment occurred significantly earlier in the trial (co-trimoxazole) group compared with the control (chloramphenicol) group. Before the end of the third day 78.1% of toxæmic patients in the trial group became non-toxæmic compared with only 50% in the control group. At the end of six days only 3.1% of patients in the trial group remained toxæmic as compared to 16.6% in the control group. These results are similar to almost all previous workers (Kamat, 1969; Kamat, 1970; Farid *et al.*, 1970; Geddes *et al.*, 1971; Scragg and Rubidge, 1971).

Two patients in our control group relapsed but none in the trial group did so. Kamat (1970), and Farid *et al.* (1970) did not encounter any relapse in patients treated with co-trimoxazole. In Geddes *et al.* (1971) series there were two relapses out of 23 patients and in Scragg and Rubidge's series a staggering relapse rate of 12.6% in patients treated with co-trimoxazole.

Our overall results are similar to those in the very large series of Kamat (1970) in that co-trimoxazole seemed superior to chloramphenicol in relieving toxæmia but equal to it when judged by the duration of fever. In our series, however, co-trimoxazole also seemed to be superior to chloramphenicol when judged by the incidence of relapse. The comparatively poorer results with co-trimoxazole in the series of Geddes *et al.* (1971) and Scragg and Rubidge (1971) are difficult to explain. It seems that various factors such as differences in strains of *Salm. typhi*, ethnic differences, and possibly differences in herd immunity to typhoid may be responsible for the apparent contradictions. Clearly, however, in our part of India co-trimoxazole is superior to chloramphenicol in the treatment of typhoid fever.

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Association between Previous Tuberculous Infection and Cerebral Glioma

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Summary

An increased incidence of previous infection with tuberculosis has been found in a series of patients with cerebral gliomas, and it is suggested that such an association may be due to defective immunity acting as a common aetiological factor.

Introduction

When it was noticed that three patients with cerebral gliomas, seen consecutively, had associated disease a survey of further patients with gliomas was conducted (Finn *et al.*, 1972). This showed that about a quarter had a history or significant radio-

logical evidence of previous tuberculous infection compared with a tenth of a series of 56 controls. Consequently, the larger study was carried out.

Patients and Methods

For the purpose of this investigation previous tuberculous infection was defined as either a previous history of clinical tuberculosis or the presence on chest radiography of healed apical fibrotic lesions, with or without extensive calcification, often involving the hilar regions.

Altogether, 100 case records of patients, each with a histologically-proved glioma, were studied. Eight had to be rejected because of inadequate information. The ages of the remaining 92 patients ranged from the mid-20s to over 80 years.

As controls we examined in detail the previous histories and chest radiographs of 100 patients admitted consecutively to two general medical units. In view of the decline in tuberculous infection over recent years we limited selection to patients aged 50 years or over, with the exception of three under the age of 50 who were known to have had tuberculosis in the past.

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