Methylprednisolone in multiple sclerosis: a comparison of oral with intravenous therapy at equivalent high dose

S M Alam, T Kyriakides, M Lawden, P K Newman

Summary
A randomised double-blind placebo-controlled trial of intravenous methylprednisolone versus oral methylprednisolone at equivalent high dose was carried out on 35 patients with an acute relapse of multiple sclerosis (MS). After baseline evaluation each was randomly allocated to oral treatment and intravenous placebo or intravenous treatment and oral placebo, receiving 500 mg of methylprednisolone for five consecutive days and with reassessment at days five and twenty-eight. There was no significant difference in response when disability or functional scores were compared in the two groups. Adverse effects were minor and equally distributed. In this study oral treatment with methylprednisolone was as effective as intravenous treatment in acute relapse of MS.

(J Neurol Neurosurg Psychiatry 1993;56:1219–1220)

The influence of short courses of corticosteroid therapy in relapsing multiple sclerosis (MS) has been studied in placebo-controlled trials using ACTH and intravenous methylprednisolone and in comparative studies. There is some suggestion that high doses of steroids is may be more effective than lower doses but this has not been convincingly demonstrated. Many clinicians, influenced by the published material and their own experience, now routinely use high dose intravenous methylprednisolone for patients in relapse. This form of treatment is inherently inconvenient. We have examined the hypothesis that an equivalent dose of oral methylprednisolone is no less effective.

Patients and methods
Thirty-five patients were randomised to either intravenous treatment and oral placebo (N = 20) or oral treatment and intravenous placebo (N = 15). There were no significant differences between the two groups in age, sex distribution, duration of disease or disability score at entry. There was no significant over representation in either group of any particular clinical manifestation. The Kurtzke disability score had improved at day 28 in 16 of the 20 in the intravenous group, mean score of the whole group falling from 4·85 to 3·50, and in 10 of the 15 in the oral group, mean score falling from 4·80 to 3·67. Statistical analysis using Student's t test and the Mann–Whitney rank sum test, indicated a clear improvement in both groups over the course of the study (p < 0·01) but no significant differences between the two groups at days 5 and 28.

Table
Clinical details of trial patients. Values are mean (SD)

<table>
<thead>
<tr>
<th></th>
<th>Injection (20)</th>
<th>Tablets (15)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male: Female</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Age (years)</td>
<td>41·6 (12·8)</td>
<td>41·3 (13·6)</td>
</tr>
<tr>
<td>Duration of disease (years)</td>
<td>6·5 (7·4)</td>
<td>3·8 (3·5)</td>
</tr>
<tr>
<td>Disability score</td>
<td>4·85 (1·9)</td>
<td>4·8 (1·9)</td>
</tr>
</tbody>
</table>

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There were no major side effects in either group. Minor side effects including headache, indigestion, acne, dizziness, flushed feeling, phlebitis, ankle oedema and mild depression were equally distributed in both groups. A metallic taste after injection was only present in the intravenous group. There was no increase in gastrointestinal symptoms in the group that received methylprednisolone orally.

Discussion
Most neurologists use steroid therapy in one form or another to treat patients with MS in relapse, although some hold the view that any benefit is at best only marginal.1 Intravenous methylprednisolone has been used successfully for several years but this form of therapy may be inconvenient for the patient and wasteful of hospital resources. This study suggests that oral methylprednisolone at an equivalent high dose is as effective and safe as intravenous treatment. The treatment response was the same in the orally treated patients as in the intravenous group. There were no serious adverse effects in either group; the frequency of minor side effects was equal and in particular there was no excess of upper gastrointestinal reactions in the oral group.

In an earlier controlled trial of intravenous methylprednisolone a significant improvement was seen compared with placebo when 22 patients in relapse were evaluated.1 Our study had the power to detect a 25% difference in disability grade improvement between the oral and intravenous treatments but would have required 98 randomisations to detect a 10% difference. Thus a large study is desirable which could include another arm of low dose prednisolone and should be ideally placebo controlled.

This experience in patients with MS parallels similar findings when high dose oral methylprednisolone has been evaluated in patients with rheumatoid arthritis10 and in children with idiopathic thrombocytopaenic purpura.11 Oral medication has obvious logistic advantages over intravenous therapy, and our study suggests that in MS it is as safe and no less effective.

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