Pulsed methylprednisolone for rheumatoid arthritis

Sir: A recently published letter in the Annals suggested that pulse methylprednisolone therapy would have caused the overwhelming infection in a 71 year old patient, and the authors stated that they would be extremely wary of allowing pulse therapy to become an outpatient procedure. This seems to be an overreaction to an unfortunate death in a patient, where there were several possible contributing factors (methylprednisolone pulse therapy, intravenous access, azathioprine treatment, admission to hospital, age).

We have previously reported one patient who developed an infection of his rheumatoid nodule with the same organism that colonised his intravenous cannula. It is likely that the patient described in the letter developed her infection as a result of an indwelling intravenous cannula, although the letter does not state the method of intravenous access for the infusion of methylprednisolone, whether the same intravenous access was maintained for the five days of pulse therapy (three alternate day pulses), and whether the cannula and intravenous access site were subsequently cultured for bacterial organisms which might have been present in the intravenous solution.

Pulse therapy in the 480 patients with rheumatoid arthritis reported in published studies has a low incidence of major and minor side effects, and infections have rarely complicated the treatment in reported series. As a result of our experience with one case of infection as a direct result of an intravenous cannula we no longer use indwelling cannulae to give pulse therapy. Both oral and intravenous infusions of methylprednisolone are given in our hospitals as outpatient procedures with no serious complications (particularly infective ones) being reported. It is difficult to see how giving pulse therapy to the patient at an outpatient clinic would have altered the outcome in the case reported because she developed an overwhelming infection and died despite being in hospital.

We therefore, despite the unfortunate demise of the patient reported I maintain that pulse therapy can be safely given as an outpatient procedure as long as sensible precautions for infection control are used when pulse therapy is given orally. Continuing studies suggest that the efficacy and toxicity of intramuscular gold therapy in patients with rheumatoid arthritis is favourably influenced by the addition of pulse methylprednisolone therapy in the first three months of disease modifying antirheumatic drug treatment.


Effects of eel calcitonin on rheumatoid arthritis

Sir: We have obtained exciting results in the treatment of rheumatoid arthritis (RA) in our clinic by administration of eelcalcitonin, an eelcalcitonin derivative, which is a polypeptide consisting of 32 amino acids and an endocrine substance secreted from the thyroid. The present various treatments for RA are used but few have been found valuable. This study was undertaken to obtain information about treatment of RA with calcitonin.

Two hundred and eighty-nine patients with RA, according to the revised criteria of the American Rheumatism Association, were studied. Their stage and class were determined according to the criteria of Steinbrocker et al. Ten MRC units of calcitonin were given intramuscularly twice a week for six months to 172 patients with RA (age 54 (SEM 1) years, stage 2.8 (SEM 0.06)), who had not been treated with immunomodulators or immunosuppressants (calcitonin only group). Calcitonin and a corticosteroid were given in combination to 63 patients with RA (age 54 (1-4) years, stage 3.1 (0.10)) (combination group), who had been receiving various oral doses of corticosteroids and had also received calcitonin at the same dose and by the same route as the calcitonin only group. Twenty four patients with RA who had received neither calcitonin nor corticosteroids served as the control group (age 58 (2-5) years, stage 3.0 (0-18)); some of this group had been receiving a non-steroidal anti-inflammatory drug.

In the calcitonin only group remarkable improvements in clinical symptoms and laboratory findings were noted after six months of calcitonin treatment. Class, pain, and morning stiffness were significantly (p<0.01) improved (86%) of the calcitonin only group. Additionally, grip strength of both hands, rheumatoid factor, C reactive protein, erythrocyte sedimentation rate, serum iron, haemoglobin, IgG, IgM, and albumin/globulin ratio were all significantly improved after six months of calcitonin treatment (table); all changes were significantly (p<0.05) different compared with the control group.

In the combination group no significant improvements were seen in grip strength (left), C reactive protein, erythrocyte sedimentation rate, serum iron, and haemoglobin, IgG, IgA, and albumin/globulin ratio. Improvements were found in class (p<0.01), grip strength (right) (p<0.05), rheumatoid factor (p<0.01), and IgM (p<0.05), but the improvements in grip strength and rheumatoid factor were significantly (p<0.01) less than those in the calcitonin only group. There were no significant improvements in clinical signs and symptoms or laboratory data in the control group.

Thus this study has shown that the clinical signs and symptoms of RA as well as laboratory data are dramatically improved in the group treated with calcitonin alone. It has been reported that calcitonin has an anti-inflammatory effect in rats with adjuvant arthritis. In the calcitonin only group decreases in C reactive protein and erythrocyte sedimentation rate were noted after calcitonin treatment, indicating that the inflammatory response was suppressed by calcitonin. In addition, however, reductions in IgG, IgA, and IgM also occurred after calcitonin treatment, resulting in an increase in albumin/globulin ratio and a decrease in rheumatoid factor. These results suggest that calcitonin has an effect on the immunological mechanism of RA. It is reasonable to suggest, therefore, that the therapeutic effect of calcitonin was subsequently brought about by an immunomodulating effect of calcitonin as the reduction of immunoglobulins cannot be attributed to suppression of the inflammatory response.

The current paper seems to be an addition to the findings that both serum iron and haemoglobin significantly increased, because anaemia associated with RA is thought to be induced by the deposition of iron in the synovial tissue after repeated small haemorrhages caused by the immunological reaction in the joints.

Thus it seems that calcitonin may be considered for use as a remission inducing drug in RA. Corticosteroid treatment, however, appears to have beneficial effects of calcitonin treatment.

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Improvements in symptoms and laboratory findings in patients with rheumatoid arthritis after calcitonin treatment.

<table>
<thead>
<tr>
<th>Value</th>
<th>Before</th>
<th>After</th>
</tr>
</thead>
<tbody>
<tr>
<td>Grip strength (mmHg)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Left</td>
<td>56 (4-3)</td>
<td>89 (5-1)**</td>
</tr>
<tr>
<td>Right</td>
<td>75 (5-6)</td>
<td>99 (5-3)**</td>
</tr>
<tr>
<td>Class 2</td>
<td>2-5 (0-6)</td>
<td>1-5 (0-6)**</td>
</tr>
<tr>
<td>Rheumatoid factor</td>
<td>8-2 (0-6)**</td>
<td>2-1 (0-6)**</td>
</tr>
<tr>
<td>C reactive protein (mg%)</td>
<td>33 (0-3)</td>
<td>13 (0-3)**</td>
</tr>
<tr>
<td>Erythrocyte sedimentation rate (mm/h)</td>
<td>51 (2-8)</td>
<td>39 (2-7)**</td>
</tr>
<tr>
<td>Serum iron (mmol/l)</td>
<td>11.6 (0-3)</td>
<td>12.7 (5-0)**</td>
</tr>
<tr>
<td>Haemoglobin (g/l)</td>
<td>132 (1-5)</td>
<td>129 (1-5)**</td>
</tr>
<tr>
<td>IgG (g/l)</td>
<td>20 (1-0)</td>
<td>19-5 (4-6)</td>
</tr>
<tr>
<td>IgA (mg/l)</td>
<td>121 (0-1)</td>
<td>124 (0-1)**</td>
</tr>
<tr>
<td>IgM (mg/l)</td>
<td>2750 (81)</td>
<td>2580 (75)**</td>
</tr>
<tr>
<td>Albumin/globulin ratio</td>
<td>1-31 (0-26)</td>
<td>1-50 (0-05)**</td>
</tr>
</tbody>
</table>

*p<0.05; **p<0.01 for comparison of values at the start of the study and after six months' treatment. Results analysed by Student's t test or Wilcoxon's signed rank test.
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M D Smith

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