Short Report

Placebo controlled pilot trial to study the remyelinating potential of intravenous immunoglobulins in multiple sclerosis

Martin Stangel, Friedrich Boegner, Christel H Klatt, Christoph Hofmeister, Sepp Seyfert

Abstract

Currently there is no treatment available to improve a stable deficit in multiple sclerosis. It was shown in animal models that intravenous immunoglobulins (IVIg) can enhance central nervous remyelination, and the first open trials were promising. We therefore conducted a double blind, placebo controlled pilot study to evaluate the effect of IVIg treatment in patients with multiple sclerosis with a stable clinical deficit. The primary outcome parameter was the change in central motor conduction time as an indirect measure of central myelination. Secondary outcome parameters were neurological examinations including the expanded disability status scale (EDSS), neurological rating scale (NRS), and manual muscle testing (MMT). Ten patients were treated first with placebo and then with IVIg (0.4 g/kg body weight on 5 consecutive days), the two treatments being separated by an interval of 6 weeks. There was no difference in the central motor conduction times measured before and 6 weeks after each treatment. Clinically there was a small improvement after IVIg treatment, but there was no significant difference when compared with placebo. In conclusion, our data do not support a role for IVIg in the remyelination of stable multiple sclerosis lesions as measured by central conduction time. The importance of the small clinical benefit is currently not clear.

Keywords: multiple sclerosis; intravenous immunoglobulins; remyelination

Multiple sclerosis is a disabling disease characterised by demyelination of the CNS. There is currently no therapy to improve a persistent clinical deficit. Spontaneous remyelination occurs in multiple sclerosis lesions, but it is often incomplete.1 2 Augmentation of such remyelination could provide repair of lesions and possibly reverse a neurological impairment. Treatment with intravenous immunoglobulins (IVIg) modulates the immune system and has been used successfully in some autoimmune neurological disorders.3 In multiple sclerosis, administration of IVIg can reduce the relapse rate and the number of gadolinium enhancing lesions on MRI.4 5 In an experimental model of multiple sclerosis—Theiler’s virus encephalomyelitis—it was demonstrated that immunoglobulins can enhance remyelination.6 7 In small open trials, visual acuity and colour vision improved in stable optic neuritis in patients with multiple sclerosis after IVIg treatment,8 and there was clinical improvement of paresis.9 10 We have conducted a pilot trial investigating the remyelinating potential of IVIg using central conduction times as a measure of CNS myelination.

Patients and methods

PATIENTS

Inclusion criteria were clinically definite relapsing-remitting multiple sclerosis11 with an expanded disability status scale (EDSS)7 score between 2.0 and 4.5, and a clinically stable neurological deficit. The last clinical relapse must have occurred more than 3 months ago without an ongoing improvement. Exclusion criteria were treatment with interferon-β or other immunomodulatory therapies, pregnancy, diabetes mellitus, renal insufficiency, liver disease, and heart disease.

All patients gave written informed consent before inclusion, and the study has been approved by the local ethics committee.

STUDY DESIGN

The study was designed as a double blind placebo controlled trial. A 6 week run in period to document the stability of the clinical deficit preceded two identical and consecutive 6 week treatment periods. Based on previous trials9 11 we chose a time span of 6 weeks for treatment evaluations. Each patient was treated first with placebo and then with IVIg to act as his or her own control. This non-randomised design was chosen to avoid a carry over effect due to the long half life of IVIg. A much larger sample size would have been required for the comparison in a two group analysis,12 and a longer wash out period would increase the chance of relapses that would interfere with the study. Patients, evaluating neurologist, electrophysiological
Table 1  Baseline characteristics for the enrolled patients

<table>
<thead>
<tr>
<th>Patient No</th>
<th>Age</th>
<th>Sex</th>
<th>Duration of MS (y)</th>
<th>Last relapse (months)</th>
<th>Baseline EDSS</th>
<th>Baseline NRS</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>38</td>
<td>M</td>
<td>10</td>
<td>24</td>
<td>3.5</td>
<td>71</td>
</tr>
<tr>
<td>2</td>
<td>46</td>
<td>M</td>
<td>25</td>
<td>4</td>
<td>4.0</td>
<td>71</td>
</tr>
<tr>
<td>3</td>
<td>49</td>
<td>M</td>
<td>10</td>
<td>4</td>
<td>3.0</td>
<td>92</td>
</tr>
<tr>
<td>4</td>
<td>54</td>
<td>F</td>
<td>13</td>
<td>4</td>
<td>4.0</td>
<td>78</td>
</tr>
<tr>
<td>5</td>
<td>53</td>
<td>F</td>
<td>27</td>
<td>11</td>
<td>3.5</td>
<td>84</td>
</tr>
<tr>
<td>6</td>
<td>28</td>
<td>M</td>
<td>8</td>
<td>4</td>
<td>4.0</td>
<td>60</td>
</tr>
<tr>
<td>7</td>
<td>38</td>
<td>M</td>
<td>12</td>
<td>4</td>
<td>4.0</td>
<td>60</td>
</tr>
<tr>
<td>8</td>
<td>38</td>
<td>M</td>
<td>6</td>
<td>4</td>
<td>3.5</td>
<td>75</td>
</tr>
<tr>
<td>9</td>
<td>31</td>
<td>F</td>
<td>6</td>
<td>14</td>
<td>3.5</td>
<td>67</td>
</tr>
<tr>
<td>10</td>
<td>55</td>
<td>F</td>
<td>12</td>
<td>11</td>
<td>2.0</td>
<td>81</td>
</tr>
<tr>
<td>Mean</td>
<td>43</td>
<td>M/F</td>
<td>6.2</td>
<td>4.2</td>
<td>2.0–4.0</td>
<td>60–92</td>
</tr>
<tr>
<td>Range</td>
<td>28–55</td>
<td>M/F</td>
<td>4–27</td>
<td>2.0–4.0</td>
<td>60–92</td>
<td></td>
</tr>
</tbody>
</table>

Table 2  Electrophysiological and clinical measurements

<table>
<thead>
<tr>
<th></th>
<th>At inclusion</th>
<th>Before placebo</th>
<th>After placebo/ before IVIg</th>
<th>After IVIg</th>
</tr>
</thead>
<tbody>
<tr>
<td>MEP (ms):</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>To left leg</td>
<td>22.78 (8.6)</td>
<td>23.14 (10.5)</td>
<td>22.77 (9.0)</td>
<td>21.71 (8.6)</td>
</tr>
<tr>
<td>To right leg</td>
<td>23.34 (10.4)</td>
<td>24.68 (11.7)</td>
<td>23.95 (11.2)</td>
<td>23.6 (11.3)</td>
</tr>
<tr>
<td>Amplitude (mV):</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>To left leg</td>
<td>1.51 (0.9)</td>
<td>1.45 (1.3)</td>
<td>1.46 (1.1)</td>
<td>1.44 (1.2)</td>
</tr>
<tr>
<td>To right leg</td>
<td>1.14 (0.6)</td>
<td>1.36 (0.7)</td>
<td>0.89 (0.5)</td>
<td>1.01 (0.8)</td>
</tr>
<tr>
<td>EDSS</td>
<td>3.5 (0.6)</td>
<td>3.5 (0.6)</td>
<td>3.45 (0.9)</td>
<td>3.4 (0.6)</td>
</tr>
<tr>
<td>NRS</td>
<td>73.9 (10.3)</td>
<td>71.7 (9.1)</td>
<td>74.8 (8.4)</td>
<td>78.0 (6.8)</td>
</tr>
<tr>
<td>MMT</td>
<td>223.8 (9.9)</td>
<td>227.7 (11.9)</td>
<td>228.7 (9.0)</td>
<td>232.1 (6.0)</td>
</tr>
<tr>
<td>Walk 20 m (s)</td>
<td>13.9 (3.1)</td>
<td>13.8 (2.6)</td>
<td>14.4 (3.4)</td>
<td>13.5 (2.3)</td>
</tr>
<tr>
<td>Climb 12 steps (s)</td>
<td>7.3 (1.3)</td>
<td>7.3 (1.3)</td>
<td>7.3 (1.3)</td>
<td>7.2 (2.1)</td>
</tr>
</tbody>
</table>

Values in parentheses are SD.

**RESULTS**

Analysis of the primary and secondary outcome measures with repeated measures analysis of variance (ANOVA) were performed. Multivariate tests were used if assumptions for univariate testing was possibly violated. Due to the small sample size results up to p=0.1 were considered for post hoc analysis by t test for single measure comparisons. Treatment periods were compared using two way ANOVA with repeated measurements on both factors (before and after treatment and placebo and IVIg effect). Differences in the response to the stimulus between placebo and treatment groups were tested by the interaction effect of this two factorial design.

**EFFECT OF IVIg ON NEUROLOGICAL FUNCTION**

There was a significant improvement in the NRS 6 weeks after IVIg treatment (p=0.029; table 2). However, there was also an improvement (non-significant, p=0.081) after placebo treatment, and comparison of the two treatment periods showed no significant difference. A similar small improvement was seen in the MMT with both treatments. Thus these improvements are likely to represent a placebo effect. No significant changes were found for EDSS, time to walk 20 m, and to climb 12 steps.

**EFFECT OF IVIg ON THE PRIMARY OUTCOME MEASURE**

Measurement of MEPs before and 6 weeks after each of the two treatments showed only minor changes that were not significant. After IVIg treatment, the central conduction time decreased by 0.71 (SD 2.3) ms compared with 0.55 (SD 2.7) ms after placebo (fig). Thus, no evidence was discovered for IVIg affecting central myelination as measured by MEP. P Wave latencies as indicators of peripheral conduction time were likewise unaffected. Similarly, no significant change in amplitudes was found (table 2).
ADVERSE EFFECTS

Treatment with IVIg was well tolerated. The only adverse effect occurring more often during IVIg treatment was headache (six patients during IVIg treatment, three patients during placebo treatment). Laboratory tests showed the expected rise of total serum protein and IgG concentration from 11.0 (SD 3.1) g/l to 42.3 (SD 9.1) g/l. Changes for all other laboratory tests (full blood count, liver enzymes, renal variables, and blood glucose) were only minor.

BLINDING

After completion of the trial, patients and evaluating neurologist were asked to assign which of the treatments they considered as placebo or IVIg. Four patients guessed correctly, one because of the experienced side effects (headache) during active treatment. One patient guessed incorrectly and five were not able to assign any treatment to IVIg or placebo. The evaluating physician guessed correctly in four patients and incorrectly in four patients and was not able to decide this question in two patients. This suggests that blinding was valid for both patients and evaluating neurologist.

Discussion

To evaluate the potential of IVIg to induce remyelination in patients with multiple sclerosis with stable motor deficit, this is the first study using as primary outcome central motor conduction as the best available objective measure of CNS myelination. No difference was seen between placebo and active treatment. This is in line with data from another IVIg treatment study in multiple sclerosis with MRI as the main evaluating instrument, in which only a non-significant improvement in...
the central motor conduction was seen during IVIg treatment.9

The initial results from an open study with a similar patient population as ours showed some clinical benefit after IVIg, but this study did not include electrophysiological measurements and had no placebo control.10 Subsequent controlled trials on stable motor deficit and optic neuritis in multiple sclerosis failed to confirm the positive results.19 20

There are several possible explanations why remyelination by IVIg could not be demonstrated in our and in other studies: (1) It is not clear how long and what dose of IVIg should be administered to promote remyelination, and possibly our treatment protocol was too short. However, other trials with longer treatments similarly did not show a reversal of a permanent deficit.19 20 (2) The time of treatment, when there is a fixed deficit, may be too late for a remyelinating therapy, for which there is possibly only a certain yet unknown time window. Another important question is whether IVIg can enter the brain parenchyma to target the remyelinating cells. (3) Studies in the Théier’s virus model have shown that the immunoglobulins most effectively promoting remyelination were IgMs that recognise antigens on oligodendroglial cells.21 The IVIg preparations used in multiple sclerosis studies so far contain no or only traces of IgM. (4) Finally, although the central conduction time is currently the best available measure of CNS myelination, the degree of clinical deficit in multiple sclerosis and the change in evoked potential latency may correlate poorly. In this respect, the importance of the small clinical benefit seen in our study is currently not clear, but the study was not designed to evaluate this aspect.

Many studies have shown that IVIg is a potent immunomodulator22 23 and there is compelling evidence that IVIg has a beneficial effect on relapse rate in multiple sclerosis. Despite promising results from experimental work and clinical pilot trials, this and other studies were thus far unable to demonstrate an effect of IVIg on myelin repair in permanent defects in multiple sclerosis.

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