Intravenous immunoglobulin treatment in patients with chronic inflammatory demyelinating polyneuropathy

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Abstract

Intravenous immunoglobulin (IVIg) treatment is shown to be effective in a selected group of patients with a chronic inflammatory demyelinating polyneuropathy (CIDP). The proportion of patients that improve after IVIg treatment varies between studies. Because 40% of a group of IVIg treated CIDP patients needed intermittent IVIg infusions to maintain their improved clinical condition, it is expected that IVIg is effective, at least in this subgroup of patients. However, the proportion of patients that improve after IVIg is highly dependent on the selection of patients. Patients with signs and symptoms of an active disease and clear involvement of both arms and legs appear to have the highest chance of improvement after IVIg treatment, but additional or prospective studies are needed to verify these criteria. Results obtained from small numbers of patients treated in open studies suggest that CIDP patients with a monoclonal gammopathy may also improve after IVIg treatment. Further studies are required to evaluate the prognostic factors for improvement after IVIg treatment in CIDP patients and to compare the efficacy and safety of IVIg with other long-term treatment regimes. New studies should also focus on the mechanism of IVIg treatment in patients with CIDP.

Intravenous immunoglobulin (IVIg) treatment is shown to be effective in a selected group of patients with a chronic inflammatory demyelinating polyneuropathy (CIDP), as demonstrated in randomised clinical trials. The chronic course of CIDP in principle offers the possibility of using various kinds of treatment or different dose schedules in single patients. However, the course of CIDP is not completely predictable as these patients can either have spontaneous relapses or a rapid spontaneous clinical improvement which has been observed in some patients even after a course of severe progressive weakness during several months, and indicates the need for properly designed controlled studies.

IVIg treatment in CIDP

Long term treatment with corticosteroids can have serious side-effects. PE is a time consuming procedure, needing specialised equipment and good vascular access. Complications of PE in patients with CIDP are generally not severe. In the early 1980s, when the results of the corticosteroid and PE trials in patients with CIDP were not available, fresh-frozen plasma (FFP) and later on IVIg was administered to 17 severely disabled CIDP patients. Improvement was observed in 13 patients (76%). Most of these patients needed intermittent treatment and none of them became refractory to IVIg treatment. Since then, several uncontrolled studies claimed that 20–100% of the CIDP patients may improve after IVIg treatment. The results of the studies concerning more than two patients with CIDP treated with IVIg are shown in the table.

These observations were supported by the results of a double-blind placebo controlled cross-over trial in patients with CIDP, who were judged to have responded to IVIg treatment and who needed repeated IVIg infusions to maintain their improved condition. In this cross-over study all the patients deteriorated after discontinuation of regular IVIg treatment. Thereafter, the patients were randomised to IVIg (0.4 g/kg) or placebo (albumin) for five consecutive days of treatment. All the patients responded after IVIg and none responded after placebo treatment. Improvement after IVIg was always observed within one week after the start of treatment. The time-lapse from the end of trial treatment to deterioration was significantly longer after treatment with IVIg (mean 6–4 weeks) than after placebo treatment (mean 1–3 weeks).

In a retrospective study involving 52 patients with CIDP, it was found that 32
patients (62%) improved after IVlg. Twenty-one patients (40%) needed intermittent IVlg infusions to maintain clinical improvement suggesting that improvement, at least in these patients, was caused by IVlg treatment. In two of the 32 patients, there was a short-lasting improvement, and subsequent IVlg infusions had no effect. Eighteen of 30 patients (60%) who needed one or more (intermittent) IVlg infusions to maintain the improved condition, have reached a spontaneous or therapeutically (IVlg) induced complete remission after a mean follow up period of 6-5 years. The following variables were significantly associated with improvement after IVlg treatment: disease duration of less than one year, progression of weakness until treatment, absence of discrepancy in weakness between arms and legs, areflexia of the arms, and reduced nerve conduction velocity of the motor median nerve. It is surprising that it was not possible to demonstrate the efficacy of IVlg treatment in a double-blind, placebo controlled study in 28 patients with a clinical diagnosis of CIDP who were not treated with IVlg previously. The reason for this may be the selection of CIDP patients: as many as 10 of 13 patients in the placebo group and only six of 15 patients in the IVlg treatment group fulfilled the five criteria that were associated with improvement after IVlg treatment. These observations indicate that only a subgroup of patients with a CIDP seem to respond to IVlg treatment. These prognostic factors, however, need to be evaluated in a properly designed prospective study. Another surprising finding was that three patients with a slowly progressive deterioration before the start of trial treatment had a rapid and dramatic clinical improvement within days after the start of placebo treatment. The previously performed cross-over trial, however, showed that CIDP patients responded to IVlg and not to placebo. It is therefore concluded that we require better criteria to select CIDP patients for treatment with IVlg, and that the follow up must be carefully carried out to prevent continuous prescription of therapy in the case of spontaneous or therapeutically induced clinical remission.

Comparison between treatment with steroids, PE and IVlg
To date, no controlled study that compares IVlg with corticosteroids or PE treatment has been published. A cross-over study organised by Dyck et al comparing PE with IVlg treatment in 20 CIDP patients has recently been completed in the USA. If the results of this study favour IVlg, this option may be the treatment of choice. However, if the results of the PE treated group are better, this will have to be weighed against the disadvantages of PE treatment: the limited availability for long term treatment in many centres, minor side effects and the requirement of good vascular access in general. Moreover, the costs of treatment are important. IVlg and PE are both expensive treatments. To compare the costs between PE and IVlg, the costs of PE personnel, equipment and other logistics have to be taken into account. The cost for five PE sessions or a IVlg course with in total 2 g/kg seems to be more or less equal, not only in our country but also in the United States. Corticosteroids on the other hand are cheap, but complications can be severe and expensive as well.

Several uncontrolled studies have included CIDP patients who were treated with PE, steroids or other immunosuppressive treatments before IVlg treatment, or alternatively, after a failure of IVlg treatment. Cornblath et al treated 15 CIDP patients with IVlg. Fourteen patients were treated with PE, steroids or other immunosuppressive treatment before IVlg was started. The indications for IVlg were: failure after one or more therapies (four patients), desire to avoid possible side effects (four patients), and unsatisfactory response to current therapy (seven patients). Six patients were on other therapies at the onset of IVlg treatment. The three patients showing objective improvement after IVlg ini-
tially had incomplete improvement after prednisone treatment. Some of the patients who did not improve after IVlg or PE, subsequently improved after prednisone treatment. Faed et al reported that prednisone and azathioprine did not alter the pattern of relapses that occurred in a patient needing intermittent IVlg. In our own series, prednisone treatment resulted in improvement in five of 16 patients (31%) who did not respond to IVlg. These observations indicate that some CIDP patients may improve after immunosuppressive treatment, PE or IVlg, while other patients improve only after a particular treatment. This means that it is generally worthwhile to try another treatment modality in an individual patient if either prednisone, PE or IVlg fails.

What treatment first?
Without a controlled study comparing prednisone, PE and IVlg this is a difficult question to answer. The major drawback of IVlg treatment is its high cost, and frequently this is the reason to choose prednisone as the initial treatment. However, because of the potentially serious side effects of prednisone, especially in children, we start treatment with IVlg as soon as the patient has reached a degree of muscular weakness that significantly interferes with lifestyle or prevents totally independent existence. PE, being roughly as expensive as IVlg, is logistically often difficult to perform if the patient needs long term treatment.

What is the best IVlg dose schedule?
The best IVlg dose schedule in CIDP patients is not known. Initially, most patients were treated with 0.4 g IVlg/kg/day for five consecutive days. With this dose no major complications were observed in CIDP patients. A CIDP patient may be treated with 1-0 g IVlg/kg/day for two days or even the double dose in a single day. In Kawasaki disease such a rapidly administered total dose appeared to be even more effective. However, especially in older patients, one needs to be wary of fluid dysharmony and increased plasma viscosity due to an overload of IVlg, resulting in vascular problems. We usually treat CIDP patients with 0.4 g IVlg/kg/day for five days and we have observed no major complications in the more than 65 patients treated of whom many needed repeated treatment. If a patient improves after IVlg treatment, we wait and see. When secondary deterioration follows, a 0.4 g IVlg/kg course for one day is recommended. We then put the patient on a regular treatment scheme, that comprises one treatment day every three to four weeks, depending on the severity and rate of deterioration. If this is insufficient, treatment frequency is increased to one day every other week. If the patient reaches a stable condition we gradually reduce the dose instead of increasing the interval between the administered IVlg doses.

Patients with CIDP and a monoclonal protein of undetermined significance
Initially it was suggested that the diagnosis of CIDP can only be made in the absence of a serum monoclonal protein. However, some patients with the clinical signs and symptoms of a CIDP also have a monoclonal protein of undetermined significance (MGUS). Whether these patients may be diagnosed as having CIDP is presently still a matter of debate. There are several arguments that favour doing so, especially in CIDP patients with an IgG MGUS, older than 45-50 years of age, since a monoclonal gammopathy is more frequently found in the older population. Patients with a neuropathy and an IgM MGUS, are more difficult to diagnose as CIDP. This is partly due to the fact that roughly half of these patients have serum antibodies against myelin associated glycoprotein (MAG). These antibodies, which are related to the uncompacting of myelin, are not normally found in patients with idiopathic CIDP. Additionally, local antibody deposition can be found in nerve biopsies from patients with an IgM MGUS polynuropathy. This is normally not found in CIDP.

Patients with an IgM MGUS and anti-MAG activity often have a slowly progressive, mainly sensory polyneuropathy which is infrequently found in patients with a CIDP. Patients with a chronic neuropathy and an IgM MGUS more frequently have sensory loss and nerve conduction abnormalities compared with patients with an IgG MGUS. These findings support the view that IgM MGUS neuropathies are different from IgG MGUS neuropathies.

Two recent studies are particularly interesting as 30 of the 100 patients with a clinical diagnosis of CIDP according to the criteria by Dyck also had a MGUS. No clinical or electrophysiological differences were demonstrated between the groups of CIDP patients with an IgG, IgM or IgA MGUS. Motor conduction studies were similar in CIDP patients with or without a MGUS. However, CIDP patients with a MGUS were older than the patients without a MGUS, they had on average more frequent sensory loss with less severe weakness, and the course of disease was more indolent. Most CIDP-MGUS patients (21/26) were treated with prednisone or PE. Eleven patients improved, which was significantly less than in the CIDP group without a MGUS. Whether this is due to a possible worse response in the IgM MGUS subgroup is not clear. Several patients who did not respond to one of these treatments were subsequently treated with IVlg or azathioprine. “Most” poor responders after prednisone or PE treatment improved after treatment with a “second or third” modality. One of the CIDP-MGUS patients improved spontaneously. The results obtained from small numbers of patients in open studies, including our own limited experience in CIDP patients with an IgG or IgM monoclonal protein, indicate that the patients with an IgG, but probably also some patients with a
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IgM monoclonal gammopathy, can improve after IVlg treatment. Whether a gammopathy is a prognostic factor for the chance of improvement after IVlg treatment is at present unknown.

**IVlg mechanism of action in inflammatory demyelinating polyneuropathies**

Studies on various auto-immune diseases show that IVlg can interfere with the regulation of the immune-network, either by stimulation, blockage, or down-regulation. Reviews of the clinical use of IVlg, and the various mechanisms of immunomodulation by IVlg have recently been published. Studies on the mechanism of IVlg in CIDP are limited. In vitro studies have shown that IVlg contains anti-idiotypic antibodies that recognise a cross-reactive idiotype on anti-neuroblastoma (NBL) cell line antibodies present in roughly 50% of patients with GBS or CIDP. F(ab)2 antibodies present in serum from patients that have recovered from GBS can inhibit anti-NBL antibody activity in serum from patients with CIDP or with active GBS. These inhibitory antibodies which develop during recovery from GBS are also present in IVlg. Affinity chromatography revealed that these anti-idiotypic antibodies constitute less than 1% of the total IgG antibodies present in IVlg. It is, however, not proven that the various antibodies, including anti-NBL antibodies (that are present in GBS and in CIDP patients) have a direct pathogenic role in the process of disease. Whether a large pool of donors is essential for improvement from GBS or CIDP is questionable since we have observed that improvement from CIDP can also occur after intermittent infusions of fresh frozen plasma (FFP) obtained from a limited number (20–28) of donors. Although the mechanism of action of IVlg in patients with GBS or CIDP is far from clarified, there are arguments obtained from in vitro studies that V-region-dependent interaction may play a role in the regulation of GBS and CIDP.

**Conclusion**

Patients with CIDP can improve after IVlg treatment. The proportion of patients that improve after IVlg is difficult to establish and seems to be dependent on the selection of patients. Further studies should focus on prognostic criteria in relation to improvement after IVlg. Studies comparing the effect of IVlg with PE or steroids are eagerly awaited. These studies, however, should not only look for the initial clinical significant improvement, but should also study the necessity of maintenance treatment. Since most patients with a CIDP need treatment for a long period of time, cost-benefit studies need to focus not only on the initial phase of treatment but also on the effects of long-term treatment including the potential side-effects of steroid treatment. We have such a study in progress now. Studies on small numbers of CIDP patients with a monoclonal gammopathy suggest that these patients may respond after IVlg as well. However, controlled studies are urgently required.

The mechanism of action of IVlg in CIDP is far from being completely understood. It seems that immune-regulation via V-region interaction at the antibody level may play a role. Some other mechanisms by which IVlg might interact are the induction of T-suppressor cells, down-regulation of TNF-alpha and other cytokines, interference with Fc-receptors either on nervous tissue or on B cells, and a specific action on B cells that bear the pathogenic idiotype. Further studies on the effect of IVlg in CIDP may reveal important features that can be of help in unravelling the pathogenesis of the inflammatory demyelinating polyneuropathies.

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