RESEARCH ARTICLE

Plasmapheresis in neurological disorders: six years experience from University Clinical center Tuzla [version 1; referees: 2 approved]

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Abstract

Background: Therapeutic plasma exchange (TPE) is an extracorporeal blood purification technique that is designed to remove substances with a large molecular weight. The TPE procedure includes removal of antibodies, alloantibodies, immune complexes, monoclonal protein, toxins or cytokines, and involves the replenishment of a specific plasma factor. The aim of the study was to describe the clinical response to TPE in various neurological patients, and to assess the clinical response to this therapy.

Methods: The study was retrospective. We analyzed the medical records of 77 patients who were treated at the Department of Neurology, University Clinical Center (UCC) Tuzla from 2011 to 2016.

Results: 83 therapeutic plasma exchanges were performed in the 77 patients. There was a slight predominance of male patients (54.5%), with an average age of 51±15.9 years. The most common underlying neurological diseases were Guillain–Barré syndrome (GBS) (37.7%), then chronic inflammatory demyelinating polyneuropathy (CIDP) (23.4%), multiple sclerosis (MS) (11.7%) and myasthenia gravis (10.4%). Less frequent neurological diseases that were encountered were paraneoplastic polyneuropathies (5.2%), neuromyelitis optica (also known as Devic’s disease) (3.9%), motor neuron disease (3.9%), polymyositis (2.6%) and multifocal motor neuropathy (1.2%).

Conclusions: Six years experience of therapeutic plasma exchange in neurological patients in our department have shown that, following evidence-based guidelines for plasmapheresis, the procedure was most effective in patients with GBS, CIDP and myasthenia gravis.

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Referee Status: ✔ ✔

Invited Referees

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**Background**
Therapeutic plasma exchange (TPE) is an extracorporeal blood purification technique designed for the removal of large molecular weight substances. The basic premise of the treatment is that removal of these substances will allow for the reversal of the pathologic processes related to their presence. In Asia and Australia, TPE is most commonly used for treatment of digestive system diseases, whereas in Europe and USA neurologic disorders prevail. While first experiences with TPE relate to acute life-threatening conditions, such as treatment of Guillain-Barre syndrome (GBS) or myasthenic crisis, therapeutic success has also been shown for chronic diseases where immunosuppressive therapy is often required for long-term management. The TPE procedure includes removal of antibodies, alloantibodies, immune complexes, monoclonal protein, toxins or cytokines, and involves the replenishment of a specific plasma factor.

The aim of the study was to describe the clinical response to TPE in various neurological patients, and to assess the clinical response to this therapy.

**Methods**
This study is retrospective, and examines medical records of patients who were treated at the Department of Neurology, University Clinical Center (UCC) Tuzla from January 2011 to December 2016. We recorded the patient demographics, the neurological findings of patients on admission, the diagnosis that prompted treatment with TPE, comorbidities, and any medical complications that took place. Hematological parameters including blood counts, serum proteins, electrolytes and coagulation profiles were monitored after every TPE. The neurological state of patients and the recovery and outcome of therapy were assessed again when discharged. The study received institutional ethical approval from the University Clinical Center Tuzla, and also written informed consent was obtained from all patients who were treated with TPE.

**Results**
83 therapeutic plasma exchanges were performed, in 77 patients, over the course of six years (2011 – 2016) at the Department of Neurology, University Clinical Center (UCC) Tuzla. Some of the patients received more than one course of plasmapheresis. There was a slight predominance of male patients (54.5%), with an average age of 51±15.9 years. The youngest patients were 16, and the oldest 78 (Table 1). Most patients were from the Tuzla Canton, but 28 of them were from other cantons of the Bosnia and Herzegovina, and one patient was from Croatia.

TPE is usually carried out across three sessions. In 27 patients, it was carried out in five sessions, and in one case of severe poliradiculoneuritis in a young patient with tetraplegia, seven sessions were carried out. The most common underlying neurological disease was poliradiculoneuritis, Guillain-Barré syndrome (GBS), presenting in 29 patients. These patients had a very good response to the therapy, and a good recovery of motor strength was observed. All patients with paraparesis or quadriparesis recovered some movement, even those with quadriplegia. All patients continue physical therapy in stationary conditions.

| Table 1. Demographic and clinical characteristics of the neurological patients that were treated with therapeutic plasma exchange at the Department of Neurology, University Clinical Center (UCC) Tuzla. |
|---------------------------------|------------------|
| **Patient characteristics**     | **Value**        |
| Sex (female/male)               | 35/42            |
| Mean age (years)                | 51±15.9          |
| Average number of plasmapheresis sessions per patient | 3 (every other day) |
| Neurological disease (n/%)      |                  |
| Guillain–Barré syndrome         | 29/37.7          |
| Chronic inflammatory demyelinating polyneuropathy | 18/23.4 |
| Multiple sclerosis              | 9/11.7           |
| Myasthenia gravis               | 8/10.4           |
| Paraneoplastic polyneuropathy   | 4/5.2            |
| Neuromyelitis optica (Devic’s disease) | 3/3.9         |
| Motor neuron disease            | 3/3.9            |
| Polymyositis                    | 2/2.6            |
| Multifocal motor neuropathy     | 1/1.2            |
| Most common comorbidity         |                  |
| Hypertension                    | 27               |
| Diabetes                        | 9                |
| Cancer                          | 3                |
| Heart disease                   | 3                |
| Thrombovascular event           | 3                |
Two patients had complications, including deep vein thrombosis, but after treatment continued with physical therapy. One patient developed pneumonia, due to immobility (hypostatic pneumonia), not related to TPE. Unfortunately, one patient died after the third session of plasmapheresis.

18 patients with chronic inflammatory demyelinating polyneuropathy (CIDP) underwent TPE. Due to the disease being chronic, improvement generally was slower than in the acute form of demyelinating polyneuropathy. All patients felt recovery of motor strength and their sensory ability increased, after treatment. In some of these patients TPE had been repeated over the years.

Patients with severe forms of myasthenia gravis, with generalized muscle weakness, and in some of them respiratory failure, also underwent TPE. All these procedures had no complications, and all the patients recovered motor strength, except one, where no benefits were observed.

TPE was also carried out on patients with demyelinating diseases, including nine with chronic progressive form of MS and three with neuromyelitis optica. Patients who were treated with TPE had progressive forms of MS, with a high score on the Expanded Disability Status Scale (EDSS > 7.0), and we accomplished improvements in symptoms such as tremors, spasms or paresthesias, and there was slightly improvement in motor strength. One MS patient in the progressive stage of disease died. One patient with neuromyelitis optica died, but after the treatment, on palliative care. The other two patients with Devic’s disease, with spastic quadriplegia, managed to take a few steps after treatment with an orthopedic tool after discharge, during physical therapy.

Significant improvements after TPE were observed in two patients with polymyositis, including better mobility and pain reduction, and in four patients with paraneoplastic syndromes improvements in motor strength and reduced paresthesia were observed. One of the patients had a diagnosis of cerebellar paraneoplastic disorder, caused by breast cancer, with distal weakness, tremor, ataxia and loss of perception, and inability to walk. After three courses of plasmapheresis, of five sessions each, the patient was able to walk for short distances with help. A patient with multifocal motor neuropathy had severe muscle weakness and milder atrophy, but after treatment he noticed improvement in muscle strength. However, in three patients with motor neuron disease, plasmapheresis had no effect.

Alongside the TPE, patients were receiving treatment for their underlying neurological condition (steroids, immunosuppressive agents). Also, it is important to emphasize that all patients continued with physical therapy. A good outcome of the procedure was observed in 87% of patients (improvements were registered in 25 out of 29 GBS patients, in 18 patients with CIDP, 8 with MS, 7 with myasthenia gravis, 4 with paraneoplastic disorders, 2 with Devic’s disease, 2 with polymyositis and in one patient with multifocal motor neuropathy). Only one complication was observed (pneumotorax), but the patient fully recovered. Death was registered in three patients: two had severe, progressive forms of demyelinating disease and one patient with GBS experienced sudden death. The deaths were not directly related to plasmapheresis, but rather as results of complications associated with the disease.

Discussion

The American Academy of Neurology proposed an evidence-based guideline for plasmapheresis in neurological disorders. According to these recommendations, there is strong evidence that treatment is beneficial in severe forms of GBS (severe enough to impair independent walking or to require mechanical ventilation), and also as a short-term treatment for patients with CIDP. Following these evidence-based guidelines, we treated 29 GBS patients with plasmapheresis (37.7%) and 18 CIDP patients (23.4%). Furthermore, according to these guidelines there was good evidence for treating polyneuropathy patients with plasmapheresis, and also that it had shown benefits as an adjunctive treatment in relapsing forms of MS. We treated MS patients with progressive forms of the disease, and we only achieved a mild improvement of symptoms, but no significant improvement of EDSS scores.

The study by Láinez-Andrés et al. concluded that TPE proved to be an effective alternative treatment for diseases such as GBS, CIDP and myasthenia gravis. In comparative studies with intravenous immunoglobulin, the efficacy of both therapies is similar. A recent, large meta-analysis by Ortiz-Sales et al. concluded that there is no evidence that either treatment is more effective or safe in the management of GBS and myasthenia gravis.

Conclusion

Six years experience of therapeutic plasma exchange in neurological patients in our department have shown that, following evidence-based guidelines for plasmapheresis, the procedure was the most effective in patients with GBS, CIDP and myasthenia gravis. We did not record any significant complications associated with the procedure itself, only complications associated within the course of the patients’ neurological disease.

Data availability

Dataset 1. Data of neurological patients that were treated with therapeutic plasma exchange, with demographic and clinical characteristics.

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Competing interests

No competing interests were disclosed.

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15. Sinanović O, Zukić S, Burgina A, et al.: Dataset 1 in: Plasmapheresis in neurological disorders: six years experience from University Clinical center Tuzla. F1000Research. 2017. Data Source
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Current Referee Status: ✔ ✔

Version 1

Referee Report 06 September 2017

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Hidenori Matsuo
Department of Clinical Research, Nagasaki Kawatana Medical Center, Nagasaki, Japan

The experience of therapeutic apheresis in Tuzia is described.

The authors should discuss more the use of TPE in patients with polymyositis or motor neuron disease. According to guidelines, TPE in these diseases seems ineffective. Why did the authors perform TPE for the patients, and how do they think this effects polymyositis?

Is the work clearly and accurately presented and does it cite the current literature?
Yes

Is the study design appropriate and is the work technically sound?
Yes

Are sufficient details of methods and analysis provided to allow replication by others?
Yes

If applicable, is the statistical analysis and its interpretation appropriate?
Not applicable

Are all the source data underlying the results available to ensure full reproducibility?
No source data required

Are the conclusions drawn adequately supported by the results?
Partly

Competing Interests: No competing interests were disclosed.

I have read this submission. I believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Referee Report 29 August 2017

doi:10.5256/f1000research.12795.r24544
David B. Vodusek  
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This report on a retrospective analysis of neurological patients treated with PE reflects the practice in a big neurological department in the years 2011-2016. As such it is intrinsically interesting as it provides data on neurological practice in an European region. It would add to the information if some additional general data were provided (population served by the department; number of patients treated per year).

The outcome of PE treatment should be - if possible - described in more detail, and only results, no explanation should be given in the Results section ("...due to disease being chronic..."). Improvements as reported subjectively by patients, and those resulting in objective improvement in function, should be differentiated.

In GBS, CIDP, MG and whenever else possible the authors should describe when in the time course of the disease the PE was given, what were the deficits before PE, when was the improvement noted and what was the final outcome in terms of function.

It is stated that the patients treated with PE had also immunosuppressive drugs - but it remains unclear whether all patients (GBS??), and which drugs, and whether the drug regime was changed during PE. It would be interesting to note why it was decided to give PE in MND.

It would be best to describe all untoward effects of PE in one paragraph.

It would be interesting to note whether IVIG has also been used in the department in the same time period.

Is the work clearly and accurately presented and does it cite the current literature?  
Yes

Is the study design appropriate and is the work technically sound?  
Yes

Are sufficient details of methods and analysis provided to allow replication by others?  
Yes

If applicable, is the statistical analysis and its interpretation appropriate?  
Not applicable

Are all the source data underlying the results available to ensure full reproducibility?  
Yes

Are the conclusions drawn adequately supported by the results?  
Yes

Competing Interests: No competing interests were disclosed.

I have read this submission. I believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.