Original Research Paper



Health Science

TO STUDY THE EFFECT OF THERAPEUTIC PLASMA EXCHANGE IN PATIENTS WITH MYASTHENIA GRAVIS PATIENTS

Dr Kuldeep Jareda	Junior resident -3 Department of Immunohematology and Transfusion Medicine SMS Medical College Jaipur
Dr Shailendra Singh*	Junior Resident -3 Department of Immunohematology and Transfusion Medicine SMS Medical College Jaipur.*Corresponding Author
Dr. Sunita Bundas	Senior Professor Department of Immunohematology and Transfusion Medicine SMS Medical College Jaipur

ABSTRACT

BACKGROUND- An integrated understanding of therapeutic plasma exchange (TPE) effects on immunoglobulins, autoantibodies, and natural or acquired (vaccine) protective antibodies in patients with autoimmune myasthenia gravis (MG) is lacking. Prior studies measured TPE effects in healthy volunteers or heterogeneous

autoimmune disease populations.

METHODS- This descriptive cross-sectional study was conducted in Department of Immunohematology & Transfusion Medicine, Blood Bank and Department of Neurology SMS hospital, Jaipur.

RESULTS- Out of total 9 patients of MG 8 patients were improved and 1 patient failed to respond to therapy. CONCLUSION- Therapeutic plasma exchange is a first line of management for most of the neuroimmunological disorder.

KEYWORDS : TPE, Auto-immune, Outcome

INTRODUCTION

Neuro-immunological disorder consists of disease in which the immune system seems to attack the nervous system. In order to consider TPE as a therapeutic option, two condition need to be present, a disease state related to the presence of a pathological substance in the plasma and the possibility of removing the substance in a sufficient amount to permit resolution of the disease. TPE is used to treat immunologically mediated peripheral neuropathies including chronic inflammatory demyelinating polyneuropathy, guillain-barre syndrome and other disorder such as myasthenia gravis.

In a recent report of the Therapeutics and Technology and Assessment Subcommittee of the American Academy of Neurology, TPE was established as an effective course of treatment for many diseases; it is offered in cases of severe acute inflammatory demyelinating polyneuropathy (AIDP)/GBS, in the short-term management of CIDP (Category-1), and is probably effective and should be considered for mild AIDP/GBS. There have been some other case reports and small studies in which it was claimed that TPE might be effective for some other neurologic diseases such as multiple sclerosis, neuromyelitisoptica, acute disseminated encephalomyelitis (ADEM), Stiff-man syndrome, Bickerstaff's encephalitis and hemorrhagic leucoencephalitis . Additionally, it has been suggested that TPE may also be successful in treating complications of the central nervous system resulting from systemic hematologic diseases such as thrombocytopenic purpura.²

MATERIAL AND METHODS

STUDY AREA: Department of Immunohematology & Transfusion Medicine, Blood Bank and Department of Neurology SMS hospital, Jaipur.

STUDY PERIOD: Over a period of one year after approval of plan research review board of institute from June 2019 till desired sample size is achieved.

TYPE OF STUDY: Cross-sectional

DESIGN OF STUDY: Descriptive

STUDY GROUP: Patients with myasthenia gravis

INCLUSION CRITERIA- Patients with myasthenia gravis

EXCLUSION CRITERIA-Disorder in which role of apheresis therapy is not established.

Patient who have experienced a serious adverse event associated with the first TPE procedure.

Patients who have not given informed consent.

Patients who is medically not fit for TPE

BESULTS

Table 1. Clinical outcome

Clinical outcome	No of patients	Percentage
Improved	8	88.89
Failed to response	1	11.11
Total	9	100.00

Out of total 9 patients of MG 8 patients were improved and 1 patient failed to respond to therapy.

DISCUSSION

Myasthenia gravis is an autoimmune disease which results from antibodies that block or destroy nicotinic acetylcholine receptors (AChR) at the junction between the nerve and muscle. This prevents nerve impulses from triggering muscle contractions. Most cases are due to immunoglobulin G1 (IgG1) and IgG3 antibodies that attack AChR in the postsynaptic membrane, causing complement-mediated damage and muscle weakness.^{3,4}

Newsom-Davis (1979) $^{\rm 5}$ compared the long-term effect of plasma exchange plus immunosuppressive drug in seven participants with myasthenia gravis to the effect of immunosuppressive drug alone in seven participants with myasthenia gravis. Plasma exchange was associated with improvement (100%) in all seven participants which has a higher outcome when compared to our study.

Olarte MR (1978-1980) 6 studied effect of plasmapheresis in myasthenia gravis among 21 patients. 350 plasma exchanges were performed on 21 MG patients, in each exchange about two litres of plasma were exchanged for two weeks. No adverse effects were attributed to the procedure, except transient thrombocytopenia. Out of 21 patients, 17 patients (81%) improved and 4 patients failed to improve after TPE for 2 weeks which was very similar to our result

CONCLUSION

Therapeutic plasma exchange is a first line of management for most of the neuroimmunological disorder.

REFERENCES

- 1. Corina Roman-Filip. Therapeutic plasma exchange and double filtration
- plasmapheresis in severe neuroimmune disorders. Lehmann HC, Hartung HP, Hetzel GR, St€uve O, Kieseier BC. Plasma exchange in neuroimmunological disorders. Part 1: rationale and treatment 2. of inflammatory central nervous system disorders. Arch Neurol 2006; 63:930-5.
- "Myasthenia Gravis Fact Sheet". NINDS. 10 May 2016. Archived from the original on 27 July 2016. Retrieved 8 August2016. Phillips WD, Vincent A (1 January 2016). "Pathogenesis of myasthenia gravis: 3.
- 4. update on disease types, models, and mechanisms". F1000Research. 5: 1513.
- Newsom-Davis J, Wilson SG, Vincent A, Ward CD. Longterm effects of repeated plasma exchange in myasthenia gravis. Lancet. 1979;1(8114):464-8. Antozzi C, Gemma M, Regi B, Berta E, Confalonieri P, Peluchetti D, et al. A 5.
- 6. short plasma exchange protocol is effective in severe myasthenia gravis. J Neurol. 1991;238(2):103-7