

Changing trends in indications for Therapeutic Apheresis (TA) in tertiary care centre *(and its comparison with published report)*

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Introduction: *TP and Cytapheresis*

- Therapeutic Apheresis (TA) is an established procedure and its use has **grown substantially** and rapidly over the past **decades**
- TA can be categorized into three types: **Plasmapheresis**, **Leukapheresis**, and **Cell Depletion**
- TP is far more common and is divided into three types:
 1. **Continuous Renal Replacement Therapy (CRRT)**
 2. **Continuous Hemofiltration (CH)**
 3. **Immuno-Adsorption (IA)**



Introduction — ASFA adds & redefines!

- ASFA evaluates **potentially newer** for apheresis and categorizes the indications into Category I to IV on the basis of available case-reports, case series, and randomized control trials
- Categorization is dynamic and ASFA continues to **add** [*Medical conditions went up from 68 (ASFA 2010) to 87 (ASFA 2016)*] **and re-define** categories for various indications by consistently applying the **Grading of Recommendations Assessment Development and Evaluation (GRADE)** system definitions. Inclusion of fact sheets of certain newer diseases such as Henoch-Schonlein purpura, Heparin Induced Thrombocytopenia (HIT), etc.

Aim and Objectives

- **Evaluation** of our institute's **TA data** with respect to **indications**, **clinical response** and **adverse events**
- We also compared this data with an earlier published Indian report [**INDEX STUDY**-*Sharma et al (Trans Aph Sci 2011:45;239-45)*]

Materials And Methods

Materials and Methods – *Type/Consent/Procedure*

- **Retrospective** analysis of TA procedures; performed from Jan 2014 to Jun 2016 in Transfusion Medicine department at large tertiary care hospital
- Primary Physician in consultation with TMS assessed **patient suitability**
- Informed **consent** was obtained
- **Com.Tec**; PL1 kit for TP, modified PL1 for CP and P1R for IA, respectively
- **Central venous access** with double-lumen catheter was used in all patients
- Estimated plasma volume (in litres) = $TBV * (1 - \text{haematocrit})$
- **Injectable calcium gluconate** (10%; 10 ml diluted in 100 ml normal saline through intra-venous drip) for **every 1000ml to 1200ml** of plasma volume processed as prophylaxis to prevent citrate related adverse reactions

Materials and Methods - *Data Collection*

- Patient's details such as **age, gender, weight, clinical indication, ASFA category, clinical response** and **adverse events** were recorded
- Pre-procedure and post-procedure haematological parameters such as **Hb, Hct, platelet count, serum calcium, coagulation** parameters
- All procedures and **adverse reaction management** were done according to the departmental SOP
- Data from present study was compared with index study

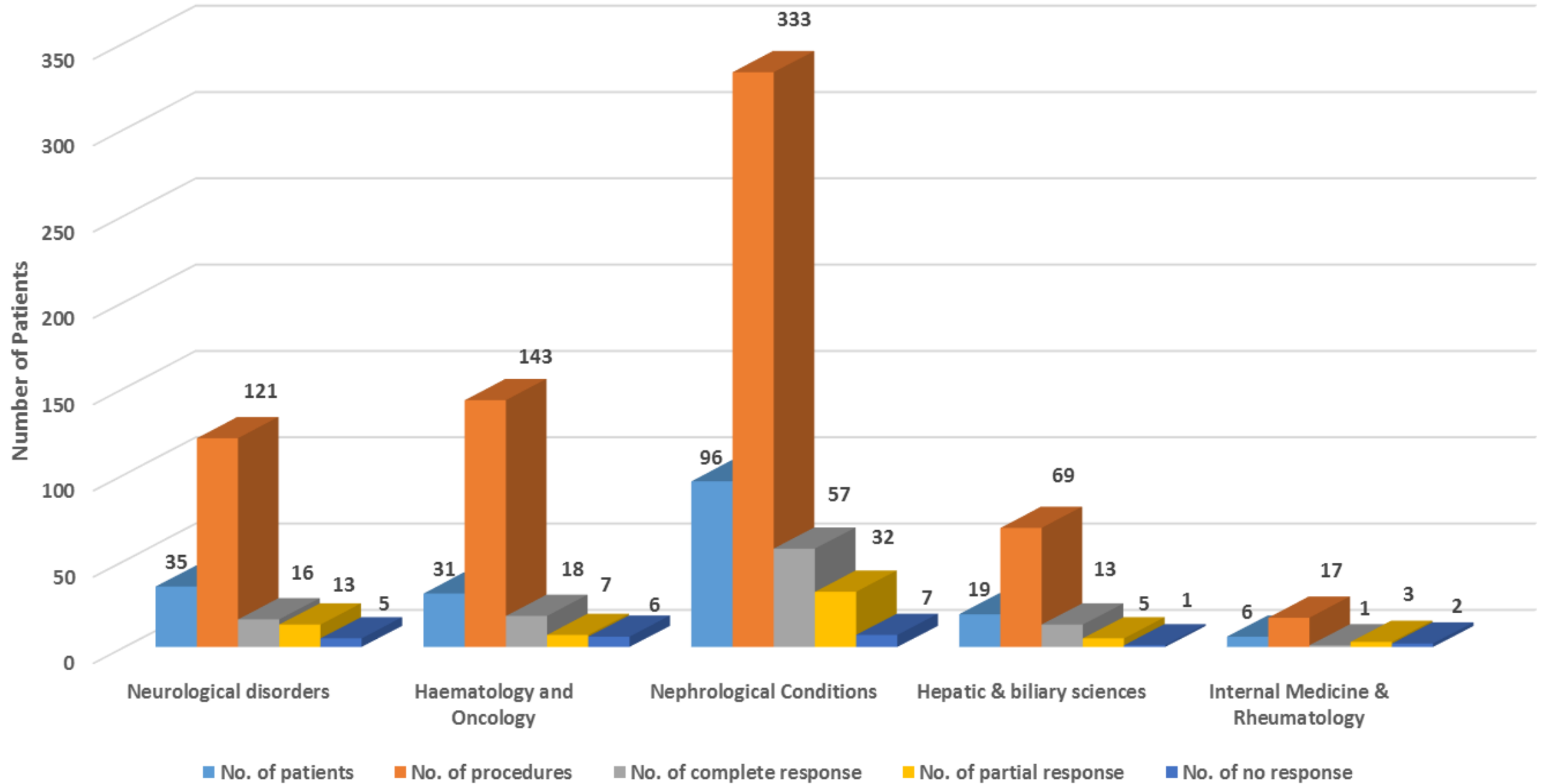
TA frequency, duration and clinical response criteria

Clinical Disorder	Frequency and duration	Response criteria	
		Complete	Partial
Neurological disorders (AIDP, MG, CIDP)	Every alternate day (5 cycles up to 2 weeks)	a. Muscle power = regained to normal b. Weaning off Mechanical ventilation – Yes c. Improvement in vital capacity over base line – $\geq 50\%$	a. Not fully recovered b. Prolonged c. $\leq 50\%$
Haematological disorders (TTP and HUS)	Daily till a sustained remission attained (6 days to 2 weeks)	a. Platelet count $\geq 1.5 \times 10^3 / \mu\text{l}$ b. Serum LDH < 250 U/L c. RFT within normal range	a. $\geq 50 \times 10^3 / \mu\text{l}$ b. < 500 U/L c. Partial recovery
Renal disorders (ABOi/HLAi, AMR)	Every alternate day (5 to 6 cycles)	Recovery of RFT - Complete	Partial
Hepatic Disorders (ABOi)	Every alternate day	Recovery of LFT - Complete	Partial
Internal Medicine & Rheumatology (SLE, Purpura Fulminans)	Daily (3 to 4 cycles)	a. Urinalysis Haematuria= No Proteinuria = No b. Recovery of Renal Functions- complete	Yes Yes Partial

RESULTS

- No. of patients: **187** (118 Males and 69 females)
- Mean age: **41** years (7 months - 73 years)
- TA procedures: **683**
 - i. 504 TPE (73.7%) → 144 patients
 - ii. 176 CP (25.7%) → 41 patients
 - iii. 03 IA (0.43%) → 2 patients
- **Indication category** in the present study according to 2013 ASFA guidelines:
 - Category I → 99
 - Category II → 59
 - Category III → 29

Response in patients — Complete/Partial/No response



Parameters	Present Study	Sharma et al (Trans Aph Sci 2011:45;239-45)
Procedures	683	492
Number of Patients	187	Discussion
Males	118 (63.1%)	
Females	69 (36.9%)	
ASFA Category	Discussion- Adverse Reactions	
I	365 (53.4%)	en years' [492] data - ↑
II	225 (32.9%)	Much lower rate 15 (0.02%) of adverse reactions as compared to index AIDP/GBS] (75.4% Vs.
III	93 (13.6%)	publication 73 (14.83%)
Procedures Performed for- Nephrology	333 (48.9%)	1. Change in replacement fluid from FFP to albumin in almost all indications
Haematology & Oncology	143 (20.9%)	2. Very little (300 ml) in CP or no fluid loss in IA - only minimal/no replacement fluid
Neurology	121 (17.7%)	3. Fewer Hypocalcaemic episodes as IV calcium gluconate given prophylactically
Hepatic & Biliary Sciences	69 (10.1%)	4. Central venous catheter :100% Vs. 69%.
Internal Medicine and Rheumatic disorders	17(2%)	
Response Rate	76.2% (133/174)	
Encouraging Response Rate	ABOi & aHUS (100%)	
Adverse Events	Neurology (88.4%)	
Common type	15 (0.02%) procedure Hypotension, hypocalcaemia, nausea.	73 (14.83%) procedures Hypotension, catheter dysfunction, hypocalcaemia

Conclusion

- In the last decade there has been **increase in number and spectrum of indications** for therapeutic apheresis and frequency of these procedures.
- **Safety** of these procedures have also improved.

THANK YOU