27th February, 2025		
Venue: Hotel Taj Ganges, Varanasi		

Time	Topic	Speaker
2:00 PM - 4:30 PM	Workshop 2A: Botulinum Neurotoxin in Movement Disorders	Incharge: Pankaj Agarwal (Mumbai) Jasloveleen (Mohali)
		Deepika Joshi (Varanasi)
		Hrishikesh Kumar (Kolkata)
		Anil Venkitachalam (Mumbai)
5:00 PM - 7:30 PM	Workshop 1A: Hands-on Genetics for the Movement Disorder Clinician	Incharge: Vikram Holla (Bangalore) Roopa Rajan (New Delhi)
		Gautam Arunachal (Bengaluru)
		Ramprasad VL (Bengaluru)
		Sneha Kamath (Bengaluru)
2:00 PM - 4:30 PM	Workshop 2B: Deep Brain Stimulation for Movement Disorders	Incharge: Rukmini Mridula (Hyderabad) Shejoy Joshua (Kochi)
		Aditya Gupta (Gurugram)
		Rajesh Alugolu (Hyderabad)
		Pettarusp Wadia (Mumbai)
5:00 PM - 7:30 PM	Workshop 1B: Cutting edge Neurophysiology for Movement Disorders	Incharge: Nitish Kamble (Bengaluru) Divya MR (New Delhi)
		Robert Chen (Toronto)
		Shweta Prasad (Bengaluru)
		Ashish Susvirkar (Gujarat)
		Supriyo Chowdhury (Kolkata)

28th February, 2025 Day 1 (Hall A)		
Time	**Topic**	**Speaker / Panelist**
8:00 AM - 8:45 AM	Registration and Breakfast	
8:45 AM - 9:00 AM	Welcome address by MDSI president	
9:00 AM - 10:30 AM	**Session: Plenary 1**	
	Chairpersons: Kalyan Bhattacharya (Kolkata), Madhuri Behari (New Delhi), Man Mohan Mehndi Ratta (Delhi), Navneet Kumar (Kanpur)	
9:00 AM - 9:30 AM	Approach to parkinsonian syndromes	U Meenakshisundaram (Chennai)
9:30 AM - 10:00 AM	Approach to hyperkinetic movement disorders	Sanjay Pandey (Faridabad)
10:00 AM - 10:30 AM	An overview about “Trial readiness” – what is required for a successful clinical trial	Susanne Schneider (Germany)
10:30 AM - 10:45 AM	**Tea Break**	
10:45 AM - 1:00 PM	**Session: Plenary 2**	
	Chairpersons: Asha Kishore (Kochi), GM Wali (Karnataka), Pramod Pal (Bengaluru), Ravi Yadav (Bengaluru)	
10:45 AM - 11:30 AM	MDSI Oration: Plasticity in Movement Disorders	Robert Chen (Toronto)
	Chairpersons: SK Poddar (Varanasi), PK Maheswari (Agra), Deepak Arjundas (Mumbai), Rajesh Singh (Varanasi)	
11:30 AM - 12:00 PM	Neuromodulation for movement disorders: present and future	Rupam Borgohain (Hyderabad)
12:00 PM - 12:30 PM	Assistive Technologies in Movement Disorders: What Clinicians Should Know	Roongroj Bhidiyasiri (Thailand)
12:30 PM - 1:00 PM	Autosomal Dominant Spino Cerebellar Ataxia: what we have learned and future pathway	Hrishikesh Kumar (Kolkata)
1:00 PM - 2:00 PM	**Lunch Symposium**	
	MDSICON Quiz	**Panelist:** **Mitesh Chandranna (Ahmedabad)** **Udit Saraf (Palghar)** **Kanchana Pillai (Mumbai)** **Varun Kumar Singh (Varanasi)**
2:00 PM - 4:00 PM	**Parallel Session 1A: Clinical skills - Video based phenomenology**	
	Chairpersons: Avinash Chandra Singh (Varanasi), Rajniti Prasad (Varanasi), Abu Zafar Ansari (Varanasi), Pahari Ghosh (Kolkata)	
2:00 PM - 2:30 PM	Updates in the evaluation and treatment of chorea	Deepika Joshi (Varanasi)
2:30 PM - 3:00 PM	Unusual and perplexing phenonemenologies in movement disorders (stereotypies, myorrhythmia, hyperekplxia, startle syndromes etc)	Suvorit Bhowmick (Vadodara)

3:00 PM - 3:30 PM	Facial movement disorders	Susanne Schneider (Germany)
3:30 PM - 4:00 PM	Clinical Use of Red Flags in Parkinsonism: Evidence vs. Experience	Roongroj Bhidiyasiri (Thailand)
4:00 PM - 4:30 PM	**Tea Break**	
4:30 PM - 6:30 PM	**Parallel Session 2A: Technology in Movement Disorders**	
	Chairpersons: Ashish Varma (Varanasi), Vinay Agarwal (Raebareli), Hardeep Malhotra (Lucknow), Chandan Singh (Varanasi)	
4:30 PM - 5:00 PM	Rating scales in movement disorders	Vijay Shankar P (Chennai)
5:00 PM - 5:30 PM	Transcranial magnetic stimluation and tDCS in Movement Disorders	Nitish Kamble (Bengaluru)
5:30 PM - 6:00 PM	Transcranial ultrasound stimulation: Novel non-invasive neuromodulation for movement disorders	Robert Chen (Toronto)
6:00 PM - 6:30 PM	Technology for rehabilitation in Movement Disorders	Rajinder Dhamija (New Delhi)
6:30 Onwards	**Inauguration & Fellowship**	
28th February, 2025 Day 1 (Hall B)		
Time	**Topic**	**Speaker / Panelist**
2:00 PM - 4:00 PM	**Parallel Session 1B: Tremor and Dystonia**	
	Chairpersons: Ashish Duggal (New Delhi), Manish Bhartiya (Pune), L K Prasanth (Bengaluru), Varun Kumar Singh (Varanasi)	
2:00 PM - 2:30 PM	Dissecting tremor syndromes: clinical and electrophysioloigcal clues	Shweta Prasad (Bengaluru)
2:30 PM - 3:00 PM	Updates in the treatment of tremor	Divya MR (New Delhi)
3:00 PM - 3:30 PM	Dystonia: is genetics replacing phenomenology?	Roopa Rajan (New Delhi)
3:30 PM - 4:00 PM	Current consensus in dystonia management	Narendra Barad (Ahemdabad)
4:00 PM- 4:30 PM	**TEA BREAK**	
4:30 PM - 6:30 PM	**Parallel Session 2B: Secondary Movement Disorders**	
	Chairpesons: Abhishek Pathak (Varanasi), Ashok Kumar (Patna), Niraj Kumar (Telangana), Anand Kumar (Varanasi)	
4:30 PM - 5:00 PM	Movement disorders in systemic diseases	Sahil Mehta (Chandigarh)
5:00 PM - 5:30 PM	Drug induced movement disorders	Bhaskar Ghosh (Kolkata)
5:30 PM - 6:00 PM	Peripherally induced movement disorders	Divya K P (Thiruvananthpuram)
6:00 PM - 6:30 PM	Stiff person spectrum disorders	Netravathi M (Bengaluru)

6:30 Onwards	**Inauguration & Fellowship in Hall A**	
28th February, 2025 Day 1 (Hall C)		
Time	**Topic**	**Speaker / Panelist**
2:00 PM - 4:00 PM	**Parallel Session 1C: Lightning talks - 12 talks 120 minutes (6 talks + 10 minutes) - 2 Themes - Basic Sciences/Clinical**	
	Chairpersons: Ashutosh Tiwari (Gorakhpur), Pratibha Prasad (Patna), Sooraj Patil (Belgavi), Archana Ojha (Prayagraj), Priyanka Kashyap (Bhopal), Rajesh Sonkar	
2:00 PM - 2:10 PM	Long read sequencing in movement disorders	Debjyoti Dhar (Lucknow)
2:10 PM - 2:20 PM	Role of cerebellum in dystonia	Sarika Patil (Rajkot)
2:20 PM - 2:30 PM	Glucocerebrosidase in Parkinson's disease	Ashish Vijayaragahvan (Thiruvanathapuram)
2:30 PM - 2:40 PM	Glymphatic system in PD	Mansi Shah (Mumbai)
2:40 PM - 2:50 PM	Alpha-synuclein seed amplification assay	Arindam Ghosh (Kolkata)
2:50 PM - 3:00 PM	Movalepsy	Anand Kumar (Varanasi)
3:00 PM - 3:10 PM	Troubleshooting diphasic dyskinesia	Adrish Mukherjee (Kolkata)
3:10 PM - 3:20 PM	Microvascular decompression in HFS	Sreenivas U M (Chennai)
3:20 PM - 3:30 PM	GLP-1 agonists in PD	Heli Shah (Ahemdabad)
3:30 PM - 3:40 PM	CANVAS and SCA27B	Shivani Rath (Cuttuck)
3:40 PM - 3:50 PM	Programming tips for FoG in PD	Vaibhav Mathur (Jaipur)
3:50 PM - 4:00 PM	Pallidothalamic tract (PTT) lesioning in movement disorders	Anish Mehta (Bengaluru)
4:00- 4:30 PM	**TEA BREAK**	
4:30 PM - 6:30 PM	**Parallel Session 2C: PD Management Capsule**	
	Chairpersons: Rajeev Verma (Varanasi), Rakesh Kumar Mishra (Bhopal), Ajai Kumar Singh (Lucknow), Ravi Yadav (Bengaluru)	
4:30 PM - 5:00 PM	Current updates in treatment of early PD	Syam Krishnan (Thiruvananthpuram)
5:00 PM - 5:30 PM	Infusion therapies for Parkinson's disease	Moeed Syed Zafer (Vishakhapatnam)
5:30 PM - 6:00 PM	Cognitive and neuropsychiatric symptoms in PD: screening and management	Atanu Biswas (Kolkata)
6:00 PM - 6:30 PM	Evidence based management of autonomic dysfunction in Parkinsonian disorders	Rukmini Mridula Kandadai (Hyderabad)
6:30 Onwards	**Inauguration & Fellowship in Hall A**	

01st March, 2025 Day 2 (Hall A)		
Time	**Topic**	**Speaker / Panelist**
7:30 AM - 8:30 AM	**PLATFORM SESSION (CLINICAL SCIENCE)**	
7:30 AM - 7:40 AM	Efficacy of propranolol in tremors in patients with Spinocerebellar Ataxia 12	Prachi Mohapatra
7:40 AM - 7:50 AM	Deciphering differences between Functional tic-like movements versus Organic tics	Ajith Cherian
7:50 AM - 8:00 AM	Insights into Functional Movement Disorders: A Retrospective Review at a Tertiary Care Centre	Kartika Gulati
8:00 AM - 8:10 AM	Profile of onset of non-motor symptoms in relation to the motor symptoms in patients with Parkinson's disease – A cross sectional single centre experience	Tarunya Nagaraj
8:10 AM - 8:20 AM	Comparison of Globus Pallidus Internus Deep brain stimulation and Sub Thalamic Nucleus Deep brain stimulation for tremor control in Parkinson's disease	Thejus B
8:20 AM - 8:30 AM	Deep brain stimulation surgery in Parkinson's disease - Hurdles and maladies!	Nikhil Korah Paul
8:30 AM - 10:30 AM	**Session: Plenary 3**	
	Chairpersons: Rajesh Verma (Lucknow), Dinesh Khandelwal (Jaipur), Ramakant Yadav (Etawah), SP Gorthi (Pune)	
8:30 AM - 9:00 AM	Defining and revising PD definition in 2024-25	Achal K Srivastava (New Delhi)
9:00 AM - 9:30 AM	Medications for PD: Evidence and Experience	Charu Sankhla (Mumbai)
9:30 AM - 10:00 AM	Sleep and Movement Disorders	Ravi Yadav (Bengaluru)
10:00 AM - 10:30 AM	Autoimmune-Related Movement Disorders: Advancements in Phenomenology and Treatment	Yih-Ru Wu (Taiwan)
10:30 AM - 10:45 AM	**TEA BREAK**	
10:45 AM - 12:30 PM	**Session: Plenary 4**	
	Chairpersons: GM Wali (Belgavi), Pramod Pal (Bengaluru), Ravi Yadav (Bengaluru)	
10:45 AM - 11:25 AM	Presidential Oration: Unity in Diversity	Asha Kishore (Kochi)
	Chairpersons: Hrishikesh Kumar (Kolkata), Bhaskar Ghosh (Kolkata), Shanker Prasad Saha (Kolkata)	
11:25 AM - 12:00 PM	Shyamal Kumar Das Oration: The Copper Story	Mohit Bhatt (Mumbai)
	Chairpersons: Parimal Das (Varanasi), Sudhir Shah (Ahemdabad)	
12:00 PM - 12:30 PM	Gut and PD	Pramod Pal (Bengaluru)

12:30 PM - 1:30 PM	**LUNCH**	
	Lunch Symposium - Closed loop DBS (Adaptive DBS) - principles and pluses	Asha Kishore (Kochi)
1:30 PM - 2:30 PM	**ANNUAL GENERAL BODY MEETING**	
2:30 PM - 4:00 PM	**Parallel Session 3A: Practice Essentials**	
	Chairpersons: Samhita Panda (Jodhpur), Mrinal Kanti Ray (Kolkata), Harmohan Sahoo (Varanasi)	
2:30 PM - 3:00 PM	Eye examination in movement disorders	Niraj Kumar (Telangana)
3:00 PM - 3:30 PM	Conundrum of lower body parkinsonism: vascular or not?	Pankaj Agarwal (Mumbai)
3:30 PM - 4:00 PM	Paediatric movement disorders: a video odyssey	Arushi Saini (Chandigarh)
4:00 PM - 4:30 PM	**TEA BREAK**	
4:30 PM - 6:30 PM	**Parallel Session 4A: Clinical Skills**	
	Chairpersons: Gopeshwar Narayan (Varanasi), Surinder Kumar (Ranchi), Mrityunjay Kumar Singh (Rishikesh), Vivek Sharma (Varanasi)	
4:30 PM - 5:00 PM	Disability assessment and benefits in movement disorders	Rohan Mahale (Bengaluru)
5:00 PM - 5:30 PM	Genetic counselling in movement disorders: practical steps	Sheela Nampoothiri (Kochi)
5:30 PM - 6:00 PM	Strategies for Engaging Patients with Spastic Paraparesis	Yih-Ru Wu (Taiwan)
7:00 PM - 9:30 PM	**VIDEO AKHADA**	**Co-ordinators** Hrishikesh Kumar (Kolkata) Roopa Rajan (New Delhi) +10 Faculty
01st March, 2025 **Day 2 (Hall B)**		
Time	**Topic**	**Speaker / Panelist**
2:30 PM - 4:00 PM	**Parallel Session 3B: Functional Movement Disorders**	
	Chairpersons: Ajay Kumar Singh (Patna), Mona Srivastava (Varanasi), Ashutosh Mishra (Raebareli)	
2:30 PM - 3:00 PM	How do I diagnose a functional movement disorder?	Pettarusp Wadia (Mumbai)
3:00 PM - 3:30 PM	Pathophysiological basis of FMD: is it really real?	Jacky Ganguly (Kolkata)
3:30 PM - 4:00 PM	Investigations and management of functional movement disorders	Dhruv Batra (Nagpur)

4:00 PM - 4:30 PM	**TEA BREAK**	
4:30 PM - 6:30 PM	**Parallel Session 4B: Treatable Movement Disorders**	
	Chairpersons: Jaya Chakravorty (Varanasi), Ruchika Tandon (Lucknow), LP Meena (Varanasi), VN Mishra (Varanasi)	
4:30 PM - 5:00 PM	Treatable ataxias: how not to miss?	Sujit Ovallath (Kannur)
5:00 PM - 5:30 PM	Movement disorders in intensive care	Soaham Desai (Anand)
5:30 PM - 6:00 PM	Movement disorders related to acute infections	Jayantee Kalita (Lucknow)
6:00 PM - 6:30 PM	Movement disorders related to chronic infections	Hardeep Malhotra (Lucknow)
7:00 PM - 9:30 PM	**VIDEO AKHADA IN HALL A**	**Co-ordinators** Hrishikesh Kumar (Kolkata) Roopa Rajan (New Delhi) +10 Faculty
01st March, 2025 Day 2 (Hall C)		
Time	**Topic**	**Speaker / Panelist**
7:30 AM - 8:30 PM	**PLATFORM SESSION (APPLIED SCIENCE)**	
7:30 AM - 7:40 AM	Effects of low intensity focused ultrasound stimulation combined with functional electrical stimulation on upper extremity motor symptoms in Parkinson disease	Naaz Desai
7:40 AM - 7:50 AM	Non-invasive transcranial ultrasound stimulation of the pedunculopontine nucleus as a treatment of freezing of gait in Parkinson's disease	Amitabh Bhattacharya
7:50 AM - 8:00 AM	Functional connectivity patterns in Imagined Writing (IW) as compared to actual Writing task (WT): a functional MRI study.	Deblina Biswas
8:00 AM - 8:10 AM	Exploring Sleep Spindle Dynamics in Essential Tremor and Essential Tremor Plus: A Comparison with Healthy Controls	Ravi Prakash Singh
8:10 AM - 8:20 AM	Electrophysiological efficacy of low frequency rTMS in patients with Functional Gait disorders	Sattwika Banerjee
8:20 AM - 8:30 AM	Differentiating two movement disorder groups from upper limb reaching kinematics: A machine learning approach	Asit Baran Bayen
2:30 PM - 4:00 PM	**Parallel Session 3C: Surgery**	
	Chairpersons: Anurag Sahu (Varanasi), Rakesh Singh (Varanasi), Kulwant Singh (Varanasi), Monika Porwal (Indore)	
2:30 PM - 3:00 PM	Lesioning surgeries in movement disorders: back to the future?	Manas Panigrahi (Hyderabad)
3:00 PM - 3:30 PM	Deep Brain Stimulation in PD: GPi vs STN	Anandh Balasubramaniam (Faridabad)

3:30 PM - 4:00 PM	Deep Brain Stimulation beyond Parkinson's Disease (dystonia, tremor)	Paresh Doshi (Mumbai)
4:00 PM - 4:30 PM	**TEA BREAK**	
4:30 PM - 6:30 PM	**Parallel Session 4C: Genetics Module**	
	Chairpersons: Alok Ojha (Varanasi), Kamlesh Sonkar (Prayagraj), Dhanvantri Shukla (Varanasi), Ravishanker Prasad (Varanasi)	
4:30 PM - 5:00 PM	Clinical spectrum of paroxysmal dyskinesias	G M Wali (Belgavi)
5:00 PM - 5:30 PM	Early onset ataxic disorders- clinico-genetic approach	Vikram V Holla (Bengaluru)
5:30 PM - 6:00 PM	Evolving spectrum of neurodegeneration with brain iron accumulation	Divyani Garg (New Delhi)
6:00 PM - 6:30 PM	Current understanding of Hereditary Spastic Paraparesis	Dinkar Kulshreshta (Lucknow)
7:00 PM - 9:30 PM	**VIDEO AKHADA IN HALL A**	**Co-ordinators** Hrishikesh Kumar (Kolkata) Roopa Rajan (New Delhi) +10 Faculty

02nd March, 2025 Day 3 (Hall A)		
Time	**Topic**	**Speaker / Panelist**
7:30 AM - 8:30 AM	**AWARD PAPER PRESENTATION**	
7:30 AM - 7:40 AM	Pediatric Onset Hereditary Spastic Paraplegia: Clinico-radio-genetic correlation	Avinash Sanap
7:40 AM - 7:50 AM	Real time Quaking induced conversion assay from skin and cerebrospinal fluid for diagnosis of Parkinson's disease	Sudharshana Prakash
7:50 AM - 8:00 AM	Comparison of effect of deep brain stimulation of subthalamic nucleus and globus pallidus internus for freezing of gait in Parkinson's disease	Manu S G
8:00 AM - 8:10 AM	Cortical Excitability in Patients with selected Spinocerebellar Ataxia Types: A TMS Study	Suchismita Majumdar
8:10 AM - 8:20 AM	REM Sleep Indices in Progressive Supranuclear Palsy – a potential biomarker?	Sathish Kumar
8:30 AM - 9:00 AM	**THESIS & AOMD AWARD PAPER**	
9:00 AM - 9:30 AM	**Session: Plenary 5**	
	Chairpersons: Royana Singh (Varanasi), RN Chaurasia (Varanasi)	
9:00 AM - 9:30 AM	Artificial intelligence in Movement Disorders	L K Prasanth (Bengaluru)
09:30 AM - 10:30 AM	**Knowledge Café (10 min per case) 10 min buffer**	
	Moderators: Jasloveleen (Mohali) Vysakh K V (Thiruvananthapuram)	Annu Aggarwal (Mumbai) Shweta Pandey (Lucknow) Sreeram Prasad (Kochi) Lulup Sahoo (Bhuvneshwar) C C Sanjeev (Bengaluru)
10:30 AM - 11:15 AM	**VALEDICTORY AND LUNCH**	
11:15 AM - 11:30 AM	**TEA BREAK**	
11:30 AM - 12:00 PM	GRAND ROUNDS	Yih-Ru Wu (Taiwan)
12:00 PM - 12:30 PM	GRAND ROUNDS	Charu Sankhla (Mumbai)
12:30 PM - 1:00 PM	DEBATE-1 Genetic testing for PD in the clinic: To do or not to do?	Propose 1 (Yes): Sneha Kamath (Bengaluru) Oppose 1 (No): Elavarasi A (New Delhi)

1:00 PM - 1:30 PM	DEBATE-2 Dopamine agonists are overutlized in PD treatment	Propose 2 (Yes): Kuldeep Shetty (Bengaluru) Oppose 2 (No): Anjali Chouksey (Jabalpur)
1:30 PM - 2:30 PM	**LUNCH**	

02nd March, 2025 Day 3 (Hall C)		
Time	**Topic**	**Speaker / Panelist**
7:30 AM - 8:30 AM	**PLATFORM PRESENTATIONS (BASIC SCIENCE)**	
7:30 AM - 7:40 AM	Investigating the Impact of LRRK2 I1371V Mutation on Calcium Homeostasis and Mitochondrial Dysfunction in Patient-Derived Astrocytes	Roon Banerjee
7:40 AM - 7:50 AM	Generation and Characterization of 3D Midbrain Organoids from LRRK2-I1371V Parkinson's Disease Patient derived iPSCs: Electrophysiological Analysis Using Multielectrode Array.	Mokshada Balaskar
7:50 AM - 8:00 AM	Targeting Hub Genes in Spinocerebellar Ataxia: An Integrative Genomics and Molecular Dynamics Approach to Drug Discovery	Surbhi Singh
8:00 AM - 8:10 AM	Uncovering novel genomic alterations in dystonia using gene burden analysis	Arti Saini
8:10 AM - 8:20 AM	Differential Expression of PPP2R2B Transcript Variants in SCA12: Insights from Patient-Derived PBMCs	Jyoti Rungta
8:20 AM - 8:30 AM	Altered Gene Expression Profiles for Mitochondrial Dysregulation in Peripheral Blood Mononuclear Cells in SCA12 Patients	Sabbir Ansari

28th February, 2025 Location: Outside Main Hall			
Time	**Poster No**	**Name**	**Title**
9:00 AM - 1:30 PM	**Session 1 - Parkinsonism**		
	Reviewers: Dinkar Kulshreshta (Lucknow), R. N. Chaurasia (Varanasi), Shweta Pandey (Lucknow), Ruchika Tandon (Lucknow)		
	P01	Swati yadav	Neuroleptic malignant syndrome
	P02	Manali Chandra	A Study on Altered Functional Connectivityin IPD patients with & without Cognitive Impairment With fMRI in resting state
	P03	Bhavani Madduluri	A New Dawn: Assessing Sleep DBS for Parkinson's Disease and Dystonia in a Tertiary Care Setting
	P04	Harshdeep Singh	Movements: The Soothsayer of Cognition, How Motor Timings Predict Cognitive Control in Parkinson's Disease
	P05	Sricharan Vijayakumar	Patient and caregiver profile in long-duration Parkinson's disease (> 10 years)
	P06	Neetu Chaurasia	REVERSIBLE METABOLIC ENCEPHALOPATHY WITH PARKINSONISM
	P07	Anand Kumar Rai	A prospective observational study for autonomic dysfunction
	P08	Navodaya salwe	Pimavanserin improves sleep quality in Parkinson's disease psychosis: Post hoc analysis based on duration of psychosis
	P09	Arthik Shetty	Efficacy of Pimavanserin in Parkinson's disease psychosis in Indian patients: A post-hoc analysis of phase-3 study based on Clinical Global Impression-Severity (CGI-S) scores
	P10	Vignesh Sampath Iyer	Characterising the clinical symptomatology of pain as a non-motor symptom in Parkinsons Disease
	P11	Phalguni Anand Alladi	Evaluation of Fibrinogen and α-Synuclein as biomarkers for Parkinson's disease.
	P12	Dr. Jivan Morey	Young woman who got slow in daily routine activities...
	P13	Rupam Borgohain	Delayed Normal Pressure Hydrocephalus Following Deep Brain Stimulation: Coincidence or Complication
	P14	VVSRK Prasad	Idiopathic Parkinson's disease and Normal pressure Hydrocephalus – a tale of two sisters
	P15	Rashmi rajur	Interesting case of parkinsonian hyperpyrexia syndrome
	P16	Madhavi Karri	SPG7 mutation – Novel phenotypic presentation mimicking idiopathic Parkinson's disease
	P17	Pranjali Batra	Behavior disturbances and parkinsonism syndrome is not always Wilsons : A case of PLA2G6 in young female.
	P18	Jagdeep Singh	Cataract as a clinical clue in the diagnosis of DJ-1 Parkinsonism
	P19	Athira P M	Neuropsychological insight in a cohort of Atypical Parkinsonism from ATPARK study.
	P20	Subhajit Roy	Clinical, investigational and genetic profiles of seven patients with PARK-SYNJ1: An experience from a tertiary care center in India
	P21	Vasavi Biruduraju	Central pontine and extrapontine myelinolysis presenting as subacute atypical parkinsonism in a patient with chronic alcoholism
	P22	Geethu TV	Caregiver distress in a cohort of patients with Atypical Parkinsonism: Preliminary insights from ATPARK Study.

	P23	Madhav Bahadur	AN ASSESSMENT OF NON-MOTOR SYMPTOMS IN LEVODOPA NAÏVE PATIENTS OF PARKINSON'S DISEASE
	P24	Shatabdi Choudhury	ELISA-based assessment of NLRP3 Inflammasome-associated markers in Serum of Parkinson's Disease Patients
	P25	Ajay Emani	Deciphering the role of iron status as a potential biochemical marker and their association with the clinical characteristics of patients with Parkinson's disease
	P26	Dhanya sureddy	Finding the anatomical sweetspot - Beta activity mapping to guide precision in DBS
	P27	Dhanya sureddy	High Beta, Better outcome - local field potential stimulation in DBS
	P28	Dhanya sureddy	Shape lock technology in Deep Brain Stimulation
	P29	Shubha Bhat GS	Comparison of pain scores in patients of Parkinson's disease with and without REM sleep behavior disorder.
	P30	DK Samartha	A Study of Metabolic Profile in patients with Early onset and Late onset Parkinson's Disease
	P31	Sandeep	Establishment a protocol for RNA isolation for transcriptomics from human brains using deep brain stimulation microelectrodes
	P32	Lakhan Parajiya	A RARE CASE OF METRONIDAZOLE INDUCED AKINETIC RIGIDIT STATE
	P33	Mahima Bharadwaj	Comparison of neuropsychological profiles in patients with young and late onset Parkinson's disease
	P34	Megha Shri N	Altered sleep pattern in patients with Parkinson's Disease
	P35	Jayasree Manikinda	EFFECT OF LOW FREQUENCY VERSUS HIGH FREQUENCY STIMULATION ON FREEZING OF GAIT IN PATIENTS WITH PARKINSON'S DISEASE POST DEEP BRAIN STIMULATION
	P36	Anjali	Neuronal and microglial numbers in Substantia nigra pars compacta of middle-aged mice differ based on the inherent susceptibility of the mice strain to MPTP
	P37	Niraj Kumar Srivastava	A Complex Interplay of Parkinsonism Linked with Neurocysticercosis: Representation of Three Unusual Cases
	P38	G Manvitha	A STUDY ON QUALITY OF LIFE, ITS PREDICTORS, AND NON-MOTOR SYMPTOMS IN PATIENTS WITH IDIOPATHIC PARKINSON'S DISEASE, PROGRESSIVE SUPRANUCLEAR PALSY, AND MULTIPLE SYSTEM ATROPHY.
	P39	Siddhartha Sankar Mondal	Neuromorphometric Alteration and Blood Neurofilament Light Chain in Progressive Supranuclear Palsy: A Pilot Study
	P40	Bishmita Biswas	Study of Gene Expression Linked to Mitochondrial Dysregulation in Patients with PLA2G6 Mutation
	P41	Ravi Yadav	Understanding progression and natural history of patients with Atypical Parkinsonism (ATPARK): A longitudinal follow-up study
	P42	Pawan Kumar Verma	DEEP BRAIN STIMULATION FOR PARKINSON'S DISEASE
	P43	Jayasree Manikinda	A RARE PRESENTATION OF HYPERTROPHIC PACHYMENINGITIS IN ASSOCIATION WITH LEVODOPA RESPONSIVE PARKINSON'S DISEASE
	P44	Tanmoy Maiti	Automatic Segmentation of the Subthalamic Nucleus and Globus Pallidus Internus: Can it support planning and programming in Deep brain stimulation

	P45	Cheshta Arora	Gastrointestinal Dysfunction in Parkinson's Disease: A Prospective Study of 150 Patients
	P46	Syed Tazeem Fathima	5HTTLPR (44bp Ins/Del) polymorphism: Serotonergic subtype of Parkinson's Disease
	P47	Madhavi Karri	Efficacy and Outcomes of deep brain stimulation in elderly Parkinson's disease: An institutional based study
2:00 PM - 4:30 PM	**Session 2 - Ataxia**		
	Judges: V. N. Mishra (Varanasi), Anand Verma, Abhishek Pathak (Varanasi), Hardeep Malhotra (Lucknow)		
	A01	Farsana Mustafa	Case Series of Nine cases of Cerebrotendinous Xanthomatosis
	A02	Shalaj Jain	A RARE NOVEL CASE OF SPASTIC-ATAXIA DUE TO KIF1C MUTATION: BROADENING SPG 58 PHENOTYPE
	A03	Sapna Mittal	Double Trouble
	A04	Laxmi Patil	Affection Of Dentato-Rubro-Thalamo-Cortical Tracts Causing Pancerebellar Syndrome in Multiple Sclerosis
	A05	Aniruddha Kundu	Tale of a Wobbly Child
	A06	Shreshtha Gupta	Anti Ampiphysin Cerebellar syndrome: An Insight
	A07	Neha Pandey	OPSOCLONUS MYOCLONUS ATAXIA SYNDROME (OMAS): Case series: Clinical features, cause, treatment & prognosis
	A08	Kavya R	Spectrum of Primary Ciliopathies in the Central Nervous System: Insights from Eleven Pediatric Cases
	A09	Apsara P S	Genetically proven Autosomal Recessive Spastic Ataxia of Charlevoix-Saguenay
	A10	Dhanush M	A rare presentation of paraneoplastic opsoclonus and cerebellar ataxia related to anti-Ma2 antibody: a case report
	A11	Rojina Choudhury	CONFUSION OF CLUMSY
	A12	Shreshtha Gupta	Anti Ampiphysin Cerebellar syndrome: An Insight
	A13	Angshuman Mukherjee	Sensory Ataxic Variant of GBS : A Rare Case
	A14	Prithwiraj Patra	A rare case of central Diabetes Insipidus with Autosomal Recessive Spinocerebellar Ataxia-8
	A15	MIT ANKUR RAVAL	Characterizing Digenic TBP/STUB1-Associated Spinocerebellar Ataxia Type 17: Clinical, Neuroimaging, and Genetic Insights from a Case Series
	A16	SULAKSHNA DAHIYA	Neuroleptospirosis - A rare case of cerebellitis
	A17	Harishma R S	Exploring Synaptic nuclear envelope (SYNE1) related ataxia in Asian Oceanian Populations: Case Series and Review of literature
	A18	Amira Patel	Cognitive profile in patients with Spinocerebellar ataxia
	A19	Ashutosh Gupta	A rare case of Neuronal Ceroid Lipofuscinosis [LCN2]: A novel pathogenic variant in the TPP1 gene
	A20	Ashutosh Gupta	Hypogonadotropic Hypogonadism in Spinocerebellar Ataxia 21: A novel finding
	A21	Raghunandan nadig	Dissecting the clues to diagnosis Cerebro tendinous Xanthomatosis

	A22	Nishka Mishra	Investigating cognitive impairment in patients with spinocerebellar ataxia type 12 (SCA12) using neuropsychological assessments: a cross-sectional study
	A23	Swarnava Sengupta	Abnormal Amyloid Beta Peptides and Proinflammatory Processes Detected in Plasma of Patients with SCA12
	A24	Esha Basu	Potential Blood-based Biomarkers for Redox Modulation and Neurodegeneration in Spinocerebellar Ataxia Type 12
	A25	Surbhi Singh	Screening Natural Compounds for VCP Modulation: A Promising Strategy for Reducing PolyQ Protein Aggregation in SCAs
	A26	Swapnil Samadhiya	SPINOCEREBELLAR ATAXIA 15 MIMICKING BULBAR ONSET AMYOTROPIC LATERAL SCLEROSIS
	A27	Vikram Jayant	Hereditary Spastic Paraplegia SPG78: A Case Contributing to Limited Documentation

1st March, 2025 Location: Outside Main Hall			
Time	**Poster No**	**Name**	**Title**
9:00 AM - 1:30 PM	**Session 1 - Dystonia + MaChoreaStereotyp**		
	Judges: Hardeep Malhotra (Lucknow), Abdul Quavi, Ajai Kumar Singh (Lucknow), Jayanti Kalita (Lucknow)		
	D01	Arushi Gahlot Saini	Movement disorder in children with NDUFV1-related mitochondrial complex 1 deficiency
	D02	Sneh Jain	"Severe Blepharospasm with Ptosis in a case of Artery of Percheron Infarct : A Therapeutic Challenge and success story of Botulinum Toxin Intervention "
	D03	Vikas Lakhanpal	MPAN presenting with movement disorder and peripheral motor neuropathy
	D04	Ankur Vivek	Clinical-Genetic Profiling of Dystonia Patients
	D05	Amlan Kusum Datta	Phenomenology and treatment response of movement disorders in paediatric anti NMDAR encephalitis: A cohort from Eastern India
	D06	Piyali Bhattacharya	Clinical Spectrum Of Dystonia in Parkinsonism
	D07	Laxmi Patil	Diagnostic Delays and Phenotypic Diversity in Dopa-Responsive Dystonia Due to GCH1 Mutations
	D08	Abhisek Guin	Camptocormia as an initial manifestation of IgLON5 antibody associated disease
	D09	Farsana MK	Clinical, Radiological and Therapeutic Profile of Patients With DYT-TOR1A from a single tertiary care centre from India
	D10	UJJAWAL ROY	Ultrasound as a key in the Evaluation of Tardive Syndrome: A Case Series
	D11	Nayana Bhuyan	Unraveling the mystery of a twisted male, a rare case of truncal dystonia
	D12	Gurdeep Kumar Rajan	A Study of Sleep Architecture in Patients with Cervical Dystonia Using Polysomnography
	D13	Archita Makharia	A Viral Symphony: “Piano-Playing Dystonia” as a Post Encephalitic Phenomenon in Dengue

	D14	Noopur Navanda	SUCCESS STORY OF DEEP BRAIN STIMULATION IN PEDIATRIC PATIENTS WITH PRIMARY DYSTONIA- A CASE SERIES
	D15	Ashish Sharma	Varied Presentations of MPAN (Mitochondrial Membrane Protein-Associated Neurodegeneration): A case series
	D16	Kavya R	Genetic Pediatric Dystonias : A 14-Year Analysis of 108 Cases
	D17	Kavya R	Spectrum of Primary Ciliopathies in the Central Nervous System: Insights from Eleven Pediatric Cases
	D18	Archita Makharia	Central Nervous System Tuberculosis and Movement Disorders: An Overlooked Nexus
	D19	Farsana MK	Rescue Pallidotomy and Thalamotomy in a patient with PLA2G6 associated Refractory Dystonic Storm
	D20	PHANINDHER MLN	SPECTRUM OF TREATABLE NON-WILSONIAN GENETIC DYSTONIAS
	D21	Shikha Priya	From Spasticity to Dystonic Tremor: A 15-Year Journey of Hereditary Spastic Paraplegia in a Young Adult
	D22	Dipankar Pal	Excessive Laugh! Good or Bad? A rare case of Niemann Pick type C
	D23	Monika Shailesh	A Case Report of Myoclonus Dystonia: A Rare syndrome with a Novel Mutation of the SGCE gene.
	D24	Vivek Chouhan	Pallidal Deep Brain Stimulation in VPS16-Related Dystonia: A Case Report
	D25	Hemanga Kumar Dhing	A case of a broken mandible due to chronic neglected oromandibular dystonia
	D26	PRATIBHA PRASAD	idipathic isolated oromandibular dystonia: diagnosis of exclusion
	CH27	Kuljeet Singh Anan	COMPLEX MOTOR STEREOTYPIES SECONDARY TO ACUTE ISCHEMIC STROKE
	CH28	Dr. Md. Karimulla	Whispers of the Vanishing Mind: Unravelling the Rapid Dementia and Chorea in the Wake of Nutritional Deficits
	CH31	Swansu Batra	Paraballism in Non-ketotic Hyperglycemia – Unmasking Primary Diabetes Mellitus
	CH29	Farsana MK	Five-Year Analysis of Sydenham Chorea Cases at a Tertiary Health Care Centre in South India
	CH30	Sathish Kumar	Sleep Architecture Alterations in Neurodegenerative Disorders: A Comparative Study of Huntington's Chorea and Progressive Supranuclear Palsy.
	CH32	Talika Sibal	Huntington's Disease Registry, India
2:00 PM - 4:30 PM	**Session 2 - Tremor + Myoclonus**		
	Judges: Divya MR (New Delhi), Divyani Garg (New Delhi), Pratibha Prasad (Patna)		
	T01	Atrikumar Patel	Functional Orthostatic tremor in MOG myelitis
	T02	Dipanwita Santra	Title: Task-Specific Modulation of Postural Tremor: Changes in Peak Frequency, Amplitude, and Stability Index under Cognitive and Motor Load
	T03	Mrityunjay Patidar	Development and Validation of Integrated Yoga Module for the patients with Essential Tremor
	T04	Swansu Batra	From Wings to the Trunk : Rediscovering ' Beating Tremors ' in Wilsons Disease- A Pandora' s Box.
	T05	Tanmoy Maiti	Deep brain stimulation for dystonic tremor

	T06	Jayasree Manikinda	Magnetic Resonance guided Focused Ultrasound Thalamotomy for patients with Essential tremor and Tremor dominant Parkinson's disease – Case series
	T07	DK Samartha	A Study of Metabolic Profile in patients with Early onset and Late onset Parkinson's Disease
	T08	Sandeep	Establishment a protocol for RNA isolation for transcriptomics from human brains using deep brain stimulation microelectrodes
	T09	Lakhan Parajiya	A RARE CASE OF METRONIDAZOLE INDUCED AKINETIC RIGIDIT STATE
	T10	Mahima Bharadwaj	Comparison of neuropsychological profiles in patients with young and late onset Parkinson's disease
	T11	Megha Shri N	Altered sleep pattern in patients with Parkinson's Disease
	T12	Jayasree Manikinda	EFFECT OF LOW FREQUENCY VERSUS HIGH FREQUENCY STIMULATION ON FREEZING OF GAIT IN PATIENTS WITH PARKINSON'S DISEASE POST DEEP BRAIN STIMULATION
	T13	Anjali	Neuronal and microglial numbers in Substantia nigra pars compacta of middle-aged mice differ based on the inherent susceptibility of the mice strain to MPTP
	T14	Niraj Kumar Srivastava	A Complex Interplay of Parkinsonism Linked with Neurocysticercosis: Representation of Three Unusual Cases
	T15	G Manvitha	A STUDY ON QUALITY OF LIFE, ITS PREDICTORS, AND NON-MOTOR SYMPTOMS IN PATIENTS WITH IDIOPATHIC PARKINSON'S DISEASE, PROGRESSIVE SUPRANUCLEAR PALSY, AND MULTIPLE SYSTEM ATROPHY.
	T16	Siddhartha Sankar Mondal	Neuromorphometric Alteration and Blood Neurofilament Light Chain in Progressive Supranuclear Palsy: A Pilot Study
	T17	Bishmita Biswas	Study of Gene Expression Linked to Mitochondrial Dysregulation in Patients with PLA2G6 Mutation
	T18	Ravi Yadav	Understanding progression and natural history of patients with Atypical Parkinsonism (ATPARK): A longitudinal follow-up study
	T19	Pawan Kumar Verma	DEEP BRAIN STIMULATION FOR PARKINSON'S DISEASE
	T20	Jayasree Manikinda	A RARE PRESENTATION OF HYPERTROPHIC PACHYMENINGITIS IN ASSOCIATION WITH LEVODOPA RESPONSIVE PARKINSON'S DISEASE
	T21	Tanmoy Maiti	Automatic Segmentation of the Subthalamic Nucleus and Globus Pallidus Internus: Can it support planning and programming in Deep brain stimulation
	T22	Cheshta Arora	Gastrointestinal Dysfunction in Parkinson's Disease: A Prospective Study of 150 Patients
	T23	Syed Tazeem Fathima	5HTTLPR (44bp Ins/Del) polymorphism: Serotonergic subtype of Parkinson's Disease
	T24	Madhavi Karri	Efficacy and Outcomes of deep brain stimulation in elderly Parkinson's disease: An institutional based study
	MY25	Shweta Pandey	Tumefactive Lesion in Fulminant Subacute Sclerosing Panencephalitis Presenting with Unilateral Myoclonus
	MY26	Shubham Kaudinya	RISING SURGE OF SUBACUTE PANENCEPHALITIS IN PANDEMIC ERA
	MY27	Surabhi P	UNRAVELING THE DANCE OF DEGENERATION: THE PATTERN OF MOVEMENT DISORDERS IN CREUTZFELDT-JAKOB DISEASE

	MY28	SHARAN	AN UNUSUAL CAUSE OF REFRACTORY MYOCLONUS
	MY29	Drishti Desai	TYPHUS ON THE MOVE : UNRAVELLING MOVEMENT ABNORMALITIES DUE TO SCRUB TYPHUS
	MY30	Dr Sachin Giri	Opsoclonus Myoclonus Ataxia Syndrome (OMAS)
	MY31	Kousik Karmakar	A Case Series of Parainfectious Opsoclonus Myoclonus Syndrome
	MY32	Arnab Adhya	Interferon in Subacute Sclerosing Pan Encephalitis: An underestimated option?
	MY33	Anand Vardhan	Neurocysticercosis masquerading as myoclonus with rapidly progressive Dementia
	MY34	Shivani Singh	AN UNUSUAL CASE OF SUBACUTE SCLEROSING PANENCEPHALITIS PRESENTING AS MYOCLONUS AND CERVICAL DYSTONIA

2nd March, 2025 Location: Outside Main Hall			
Time	**Poster No**	**Name**	**Title**
9:00 AM - 1:30 PM	**Session 1 - Miscellaneous**		
	Judges: Harmohan Sahoo (Varanasi), Avinash Chandra Singh (Varanasi), Ashutosh Mishra (Raebareli), Varun Kumar Singh (Varanasi)		
	M1	Pratibha Prasad	Relapse of AE was defined as new onset or worsening of symptoms after an initial improvement or stabilization of at least 2 months .[1] . Very few cases are reported regarding the relapse of NMDAR encephalitis studies are reported in Indian literature so relapse and outcome are poorly understood.
	M2	Priyanka Samal	Gait dysfunction is a common abnormality in patients with basal ganglia dysfunction. Many patients with parkinsonism use internal or external cues to aid walking. We present a case with a peculiar gait with unique maneuver to aid walking.
	M3	Dr Shikha Priya	4H Leukodystrophy is a hypomyelinating disorder caused by mutations in the POLR3A gene, typically presenting in childhood. Adult-onset cases are exceedingly rare and often more severe. We report a case of a young adult female presenting with this progressive neurodegenerative condition.
	M4	Prabhakara Sommana	Next Generation Sequencing for diagnosis of Paroxysmal Movement Disorders.
	M5	Revathi Sampath	Whole Exome Sequencing is a new Diagnostic approach for Paroxysmal Kinesigenic Dyskinesia.
	M6	Nishanth Gowda	Delineating the Movement Disorder Spectrum in Fahr's Disease/Syndrome
	M7	Neetu Rnai Dhiman	Unravelling the Genetic Influence of TPH2 on Physiotherapy Outcomes in Persistent Postural-Perceptual Dizziness: A study protocol
	M8	Vedang Desai	The prevalence of Movement disorders with Epilepsy in Immune Mediated Epilepsy Syndromes
	M9	ROHAN MAHALE	Quantification and clinical correlation of posterior cranial fossa cerebrospinal fluid volume in primary hemifacial spasm using magnetic resonance imaging

	M10	DR SANGAM SINGH	Acute onset movement disorders in children are well-recognized neurological conditions commonly encountered in clinical practice.The etiology is often diverse, with an even broader spectrum of presentations.Genetic disease like Wilsons disease can have such acute presentation with encephalitis.(1)
	M11	Dr. Anand Kumar	Genetic Determinants of Antipsychotic-Induced Tardive Dyskinesia: A Systemic Review and Meta-Analysis
	M12	Shubham Kaudinya	Stiff person syndrome (SPS) is an extremely rare autoimmune entity resulting due to absence of check mechanism over excitatory neurotransmitters in central nervous system (CNS) causing abnormal muscle contraction such as spasm, contractures and rarely myoclonic jerk.
	M13	Suman Kushwaha	Study of genetic analysis of Wilsons diseases in North Indian Cohort
	M14	Dr Chaithra	Antibodies against Leucine-rich glioma-inactivated protein 1 and contactin-associated protein-like 2 known as Morvan syndrome present with peripheral nerve hyperexcitability, insomnia, autonomic dysfunction, encephalopathy and sometimes movement disorders.
	M15	Dr Anand Kumar Rai	Neuro-rehabilitation in Movement Disorders – A Prospective Study from Tertiary Care Health Centre AIIMS, Patna
	M16	Shrikrishna Kamthankar	Autoimmune movement disorders encapsulate large & diverse group of neurologic disorders occurring either in isolation or accompanying more diffuse autoimmune encephalitic illnesses with diverse presentations.1 We present the mystery of dancing eyes & shaking limbs with Positive anti GAD65 antibody.
	M17	Sanchali Chakraborty	Etiological and clinical spectrum of movement disorders in a tertiary care hospital
	M18	Amarnath Chavan	Spastic paraplegia 11 (SPG11) is defined as progressive spasticity and weakness of the lower limbs and associated with mild intellectual disability with learning difficulties in childhood.
	M19	Avinash Ganapule	Anti-IgLON5 disease is an evolving entity which lies at the confluence of autoimmunity and neurodegeneration. Apart from the five classical presentations described, a wider spectrum is being increasingly recognised, across which this disorder has to be evaluated for.
	M20	Manali Chandra	Limb shaking transient ischaemic attack (TIA) is a rare phenomenon described in severe steno occlusive disease of the carotid artery . Limb shaking TIA frequently masquerades movement disorders like tremor , focal myoclonus , chorea or focal motor seizure.
	M21	Avinash Sanap	The adaptor protein-4 (AP-4) complex, encoded by AP4B1, AP4M1, AP4E1, and AP4S1, governs protein trafficking from the trans-Golgi to endosomes. Mutations cause SPG47, SPG50, SPG51, and SPG52, leading to developmental delays, spasticity, microcephaly, foot deformities, and epilepsy.
	M22	Divya K P	Comparison between Functional movement disorders in children and adults

	M23	Arunmozhimaran Elavarasi	Clinicoradiological profile, response to lumbar tap test and outcomes of patients with Normal Pressure Hydrocephalus
	M24	Chandrasekhar Enuguri	Primary Familial Brain Calcification (PFBC), or Fahr's disease, is a rare neurodegenerative disorder marked by progressive bilateral calcifications in brain regions like the basal ganglia, thalamus, and cerebellum
	M25	Swati Parida	TBC1D24 gene mutations are known to cause a spectrum of disorders, including epileptic encephalopathies. However, their association with paroxysmal movement disorders, such as paroxysmal kinesigenic dyskinesia (PKD), remains largely unexplored.
	M26	Kirthika Kannan	In this presentation, 7 cases are highlighted with a common clinical syndrome of the presence of movement disorder with cognitive disturbances and infection occurring in Adolescence and old age. Hence we have brought together jacob-creuzfeldt disease and subacute sclerosing panencephalitis.
	M27	Shivani Rath	ATP1A3 is associated with a spectrum of neurologic disorders, which continues to expand beyond the initially defined phenotypes of alternating hemiplegia of childhood, ROPD, and CAPOS and includes childhood-onset schizophrenia, epileptic encephalopathy, cerebellar ataxia, optic atrophy and SNHL .
	M28	Siddharth Khanna	A cross-sectional study of the spectrum of movement disorders in children with neurometabolic and neurodegenerative disorders
	M29	S Swathy	Metachromatic leukodystrophy (MLD) is an autosomal recessive demyelination of central and peripheral nervous system secondary to arylsulfatase-A enzyme defect. The rare adult-onset phenotype is a diagnostic-challenge as the presenting symptoms can overlap with other conditions commoner in adults.
	M30	Vineeta Singh	The Interplay Between Cognition and Gait: A Systematic Review of Shared Mechanisms and Clinical Implications
	M31	Debayan Dutta	Free-living amoeba are unicellular, aerobic,mitochondriate,eukaryotic protists also called amphizoic amoebae for their ability to exist as both a parasite and free-living organism.They have a mortality of over 90%.Parkinsonism and myoclonus are very rare presenting feature.
	M32	Ritesh Kumar	Tardive syndromes are a group of hyperkinetic and hypokinetic movement disorders that occur after some delay following exposure to dopamine receptor blocking agents such as antipsychotic and anti-emetic drugs. The severity of these disorders ranges from mild to disabling or even life-threatening.
	M33	Manali Chaudhari	Clinicoradiologic-genetic Profile of 20 patients with NBIA (Neurodegeneration with brain iron accumulation) from Western India
	M34	Naresh Chinthala	Painful legs and moving toes syndrome (PLMT) is a rare syndrome characterized by neuropathic pain in the lower extremities and involuntary movements of single or multiple toes.

	M35	Ahana Bhattacharya	A quantitative assessment of galanin and neuropeptide-Y in the human locus coeruleus: significance in aging and neurodegeneration
	M36	SUMAN JAIN	Intermittent theta burst stimulation (iTBS) has been used to induce neuronal and synaptic plasticity by applying a magnetic field stimulation to the brain. However, the effect of iTBS in complete SCI patients on motor cortex plasticity is still elusive.
	M37	Koustubh Bavdhankar	Dentatorubral–pallidoluysian atrophy (DRPLA) is a hereditary disease caused due to trinucleotide repeat expansions in the ATN 1 gene with an autosomal dominant mode of Inheritance. It is rarely reported in the non-Japanese population. Here we present clinic-radiological features of 9 DRPLA cases.
	M38	Deeksha Patel	Rewiring Mobility: Intermittent Theta Burst Stimulation Modulating Cortical Excitatory-Inhibitory Imbalance in Complete Spinal Cord Injury
	M39	SUBHAJIT DAS	A STUDY ON RESTLESS LEGS SYNDROME IN PATIENTS WITH CHRONIC KIDNEY DISEASE
	M40	Jyoti Chaurasia	Movement disorders like dyskinesias are rare but critical in immunosuppressed patients, such as those with HIV or CKD. They stem from metabolic issues, neurotoxicity, or CNS involvement. Effective management requires multidisciplinary care, especially with co-infections.
	M41	Sanchit Shailendra Chouksey	Parry-Romberg syndrome (PRS) is a rare craniofacial disorder characterized by progressive hemifacial atrophy with systematic manifestations. The combination with hemimasticatory spasm (HMS) is rare, with only 9 patients reported before.
	M42	Arpan Mitra	Creutzfeldt jakob disease (CJD) is a rare prion disease. It usually presents as rapidly progressive dementia as well as motor abnormalities. It is relentlessly progressive, and death occurs within 9 months of onset. Paraneoplastic encephalitis is an important differential diagnosis of CJD.
	M43	Akansha	Twinkle (TWNK gene), a mitochondrial 5`-3` DNA helicase, whose defects are known to cause mitochondrial deletion syndromes with progressive external ophthalmoplegia.
	M44	Ibrahim Hussain	Hemifacial spasm is characterized by persistent and rhythmic spasms of the facial muscles. It can be primary or secondary. Here we describe a case of hemifacial spasm secondary to a CP angle cyst.
	M45	Harish Nigam	Isolated lingual hyperkinetic disorder (ILHD) is a hyperkinetic movement disorder involving the tongue in isolation with varied presentation and etiology.[1] The objective was to analyze the clinical characteristics, phenomenology, etiology, treatment response in ILHD, and review the literature

28th February, 2025 Video E-Poster Location: Outside Main Hall				
Time	**Screen No**	**Poster No**	**Name**	**Title**
9:00 AM - 1:30 PM	**Session 1**			
	Screen 1	SP01	Sreenivas UM	Immune check point inhibitor
	Screen 1	SP02	Anil Dash	Dystonia +LMN
	Screen 1	SP03	Bhavani Madduluri	HFS
	Screen 2	SP04	Sayoni Roy Chowdhury	Child with ataxia, sz, myoclonus, dystonia ?PMD responsive to phenytoin
	Screen 2	SP05	Manthan Dave	Adult with myogenic tremor
	Screen 2	SP06	Vaibhav Mathur	PMA, cherry red spot
	Screen 3	VP10	Farsana Mustafa	GSSD Phenotype with E200K Mutation
	Screen 3	VP11	Farsana Mustafa	Steroid responseive encephalopathy assosicated with autoimmmune thyroditis
	Screen 3	VP12	Farsana Mustafa	GABA B Receptor Encephalilis
	Screen 3	VP13	Farsana Mustafa	Type 1 Sialidosis
	Screen 4	VP20	Madhvi Karri	CLNS mutation late onset focal dystonia
	Screen 4	VP21	Madhvi Karri	PRRT2 mutation
	Screen 4	VP30	Shivam Mirg	VPS16 Dystonia 30
	Screen 4	VP31	Shivam Mirg	KCNC1- progressive myoclonic epilepsy
	Screen 5	VP38	Vaibhav Mathur	PD-FOG
	Screen 5	VP39	Vaibhav Mathur	Silandiosis 1
	Screen 5	VP40	Vaibhav Mathur	Fahn Syndrome
	Screen 5	VP41	Vedang Desai	NPC2
	Screen 6	VP42	Vibhor Upadhyay	PLA2G6
	Screen 6	VP43	Vijayan K	FXTAS/MRGFUS
	Screen 6	VP44	Vijayan K	PD/MRGFUS
	Screen 6	VP45	Vijayan K	CoQ8A

	Screen 7	VP02	Ananya Karle	Auto immune encephalitis LGI-1 Antibody Positive
	Screen 7	VP04	Arnab Adhya	Holme's tremor, Psot Stroke
	Screen 7	VP46	Vyshaka K V	Paroxysmal Kinesogenic Dyskinesia
	Screen 8	VP05	Farsana MK (Aysha)	Perrault syndrome
	Screen 8	VP06	Bhavani Madduluri	PORRETTI BOLTSHAEUR
	Screen 8	VP07	Chaitra Rajanna	Early onset isolated generalized dystonia
2:00 PM - 4:30 PM	**Session 2**			
	Screen 1	VP01	Aakansha Jain	Zech Boesch Syndrome
	Screen 1	VP03	Ankur Vivek	Pseudobulbar Affect
	Screen 1	VP15	Janki Makni	Progressive myocolnus ataxia
	Screen 1	VP16	Janki Makni	Cognitive decline and spascity
	Screen 2	VP08	Deepak Chadha	Adrenomyeloneurpathy
	Screen 2	VP09	Dhanya/Vijayashankar	Autoimmune chorea dystonia
	Screen 2	VP14	Hema Krishna	Paroxysmal Dyskinesia/ Hypocalcemia
	Screen 3	VP17	Joydeep Mukherjee	YOPD
	Screen 3	VP18	Kartika Gulati	SCA 2
	Screen 3	VP19	Laxmi	Jumping Stump
	Screen 4	VP22	Manthan Dave	Myogenic tremor
	Screen 4	VP24	Nayana Bhuyan	PNS
	Screen 4	VP25	Nayana Bhuyan	Sporadic CJD
	Screen 5	VP26	Noopur Navandar	Lesch nyhan variant - HPRT1
	Screen 5	VP27	Pranjali Batra	CJD
	Screen 5	VP28	Sangeethameena S	Multiplce Sclerosis
	Screen 6	VP29	Sayoni Roy Chowdhury 1	TBC1D24 mutation
	Screen 6	VP32	Shivani Rath	SCA7
	Screen 6	VP33	Shweta Pandey	Hemiatrophy- hemidystonia/hemiparkinsonism syndrome

	Screen 7	VP34	Siddhart	Neuronal ceroid Lipotuscinosis 7
	Screen 7	VP35	Suresh Chandran	RUL shaking TIA, LICA Occulsion, HT/CKD
	Screen 7	VP36	Swathy S	Diffused glioma-likely neoplastic
	Screen 8	VP23	Mridula	DYT 1
	Screen 8	VP37	Swati Parida	NDUFA mutation