



7th international symposium on paediatric movement disorders

Barcelona, February 9 - 11, 2022

PROGRAM

Wednesday 9 February

15h45 Introduction Emilio Fernández-Alvarez and Nardo Nardocci

Session 1: Opening talks

16h00-16h25 Lecture 1: Is phenomenology still important in the era of exome and genome sequencing? *Jonatham Mink (Rochester USA)*

16h30-16h55 Lecture 2: Future challenges on epilepsy and movement disorders *Alexis Arzimanoglou (Lyon, France)*
17h00-17h30 Discussion

17h30-18h00 Video / Short communication session 1

17h30-17h45 NGLY1 deficiency *Khaterinne Mackenzie (Palo Alto, USA)*

17h45-18h00 Pending

18h00-18h30 Coffee Break

Session 2 : New insights into the characterization of clinical syndromes (1)

18h30 – 18h55 Lecture 3: Genetics mimics in cerebral palsy *Michael Krueer (Phoenix USA)*

19h00 – 19h25 Lecture 4: Functional movement disorders *Nardo Nardocci (Milan, Italy)*

19h30 – 20h00 Discussion

Thursday February 10th, 2022

Session 3 New insights into the characterization of clinical syndromes (2)

8h45 – 9h10 Lecture 5: Benign polymorphous paroxysmal movement of infancy *Emilio Fernandez-Alvarez (Barcelona Spain)*

9h15 – 9h40 Lecture 6: Hypokinetic movement disorders and parkinsonism in infancy and early childhood *Roser Pons (Athens, Greece)*

9h45-10h10 Lecture 7: Childhood onset chorea : an overview of genetic etiologies about a serie of 85 french children *Diane Doummar (Paris, France)*

10h15 – 10h45 Discussion

10h45 – 11h15 Coffee Break

11h15-11h45 Video / Short communication Session 2

11h15 - 11h30 Functional tic-like movements in adolescent females *Tammy Hedderley (London, UK)*

11h30 - 11h45 "Phenomenological patterns with multiple 'parents'." > A short observational study. *Jean Pierre Lin (London, UK)*

Session 4: Recent advances in genetic neurodevelopmental diseases with abnormal movements (1): GNAO1 Session

Chairs: Jennifer Friedman (US)

11h45 – 12h10 Lecture 8 Mechanistic role of GNAO1 in striatal control of movements *(Kirill A Martemyanov Jupiter USA)*

12h15-12h40 Lecture 9 Movement disorders in GNAO1 defect *Vincenzo Leuzzi (Rome, Italy)*

12h45-13h10 Lecture 10 Deep brain stimulation in GNAO1-related neurodevelopmental disorder *Dario Ortigoza-Escobar (Barcelona, Spain)*

13h15-13h45 Discussion

13h45 – 14h35 Lunch & Posters

14h35 – 15h05 Video/ short communication session-3

14h35-14h50 Pending

14h50-15h05 Exome sequencing in dystonia - single-gene disorders and clinical implications *Michael Zech (Munchen, Germany)*

Session 5: Recent advances in genetic neurodevelopmental diseases with abnormal movements

15h10-15h35 Lecture 11 HOPSAND spectrum (HOPS-associated neurological disorders VPS41/VPS16/VPS11 *Mamju Kurian (London, UK)*

15h40-16h05 Lecture 12 ATP1A2- and ATP1A3-associated early profound epileptic encephalopathy and polymicrogyria *Renzo Guerrini (Firenze, Italy)*

16h10- 16h35 Lecture 13 Clinical characterization of genet disorders of the glutamatergic synapse: hyperfunction versus hypofunction *Angels Garcia- Cazorla (Barcelona, Spain)*

16h40-17h10 Discussion

17h10 – 18h25 Video /short communication session-4

17h10 -17h25 Pending

17h25-17h40 Effects of deep brain stimulation in pediatric patients with dyskinetic cerebral palsy - a prospective multicenter trial. *Anne Koy (Cologne Germany)*

17h40 – 17h55 Pending

17h55 – 18h10 "Non-motor effects of neurotransmitter replacement therapy in secondary biogenic amine deficiency" *Gabielle Horvath (Vancouver, Canada)*

18h00-18h25 Pending

Friday February 11th, 2022

Session 6 Clinical spectrum

Chairs: Pending

8h45 – 9h10 Lecture 14 Immune-brain interactions and epigenetic risk factors of tic disorders in childhood *Russell Dale (Sidney Australia)*

9h15-9h40 Lecture 15 Psychiatric and motor manifestations in autoimmune encephalitis *Tiziana Granata (Milan, Italy)*

10h15-10h45 Discussion

10h45 – 11h15 Coffee Break

Session 7: Therapeutics

Chairs: Pending

11h15-11h40 Lecture 16 Outcomes of gene therapy in pediatric movement disorders *Toni Pearson (USA)*

11h45-12h10 Lecture 17 Long term outcome of functional neurosurgery for pediatric progressive dystonia: the Besta case series *Giovanna Zorzi (Milan, Italy)*

12h15-13h10 Lecture 18 Focal Ultrasound: An expanding approach to treat neurometabolic disorders *José Obeso (Madrid Spain)*

13h15-13h45 Discussion

13h45 Closing ceremony

MAIN TOPICS

New dystonia genes
Benign Hereditary chorea
Hyperkinetic Movement Disorders
Epilepsy and Movement Disorders
Deep Brain Stimulation
Gene therapy for Movement Disorders
Neurotransmitters Disorders
Stereotypes
Tic and Tourette syndrome

Organizing Committee

Alexis ARZIMANOGLU (Lyon)
Jaume CAMPISTOL (Barcelona)
Russell DALE (Sydney)
Emilio FERNANDEZ-ALVAREZ (Barcelona)
Angels GARCIA-CAZORLA (Barcelona)
Manju KURIAN (London)
Nardo NARDOCCI (Milano)

Symposium Organization

ANT Congrès
paedmovdisorders@ant-congres.com



www.paedmovdissymposium.com