50ème Réunion de la SENP+

50th SENP+ Meeting

Société Européenne de Neurologie Pédiatrique

14, 15 & 16 mars 2024

Milan, Italie

Programme Préliminaire / Preliminary Program

JEUDI 14 MARS / THURSDAY, MARCH 14 – COURSE DEDICATED TO PEDIATRICIANS

Emergencies in child neurology

13:30 – 14:00	Accueil/Welcome
14:00 – 14:30	**Migraine and headache** Caroline Ménache - *Hirslanden Clinique des Grangettes, Geneva, Switzerland*
14:30 - 15:00	**Prise en charge d'une 1ère crise épileptique chez l'enfant** Ilhem Ben Youssef Turki - *Université Tunis El Manar, Tunis, Tunisia.*
15:00-15:30	**Status epilepticus and recurrent seizures** Alec Aeby - *Université libre de Bruxelles, Brussels, Belgium*
15:30-16:00	**Ataxies aigues** François Rivier - *Université de Montpellier, Montpellier, France*
16:00-16:30	**Pediatric ischemic stroke: from initial symptoms to clinical management** Federica Teutonico - *Ospedale Niguarda Ca' Granda, Milan, Italy*
16:30-17:00	**Functional Neurological disorders** Nardo Nardocci - *Istituto Neurologico C Besta, Milan, Italy*
18:00	Présentation du Livre/Book Presentation
19:00	Cérémonie d'ouverture/Opening Ceremony
20:00	Cocktail de Bienvenu/Welcome Cocktail

VENDREDI 15 MARS / FRIDAY, MARCH 15

Hereditary disease of the cerebral white matter

07.30	Accueil/Welcome
08.00	Ouverture du congrès/Congress opening
08.30 – 10.30	**CONFERENCES/ LECTURES**
08.30 –9.00	**The history of leukodystrophies: from disease definition to understanding and therapy** Marjo van der Knaap - *Amsterdam University Medical Centers, Amsterdam, The Netherlands*
9.00-9.30	**Understanding leukodystrophies at its core: the vital contribution of neuropathology** Marianna Bugiani - *Amsterdam University Medical Centers, Amsterdam, The Netherlands*
9.30-10.00	**From discovery to hope: a journey through the history of metachromatic leukodystrophy** Francesca Fumagalli - *San Raffaele Hospital, Milan, Italy*

10.00-11.00 **COMMUNICATIONS ORALES / ORAL COMMUNICATIONS**

10.00-10.12 **Clinical and genetic study of leukodystrophies in Tunisian cohort**
Ichraf Kraoua - Lr18sp04 and Department of Pediatric Neurology. National Institute Mongi Ben Hmida of Neurology of Tunis, Tunisia

10.12-10.24 **Preliminary data from the analysis of neuroradiological findings in Type I Alexander Disease**
Ylenia Vaia - Unit of Pediatric Neurology, C.o.a.l.a. (center For Diagnosis and Treatment of Leukodystrophies), V. Buzzi Children's Hospital, Università Degli Studi Di Milano, Italy

10.24-10.36 **Multicenter International Cohort of p.Ala177Thr AGS2 homozygous mutated patients: clinical features, disease evolution and search for prognostic features**
Costanza Varesio - Department of Brain And Behavioral Sciences, University Of Pavia, Department Of Child Neurology And Psychiatry, Irccs Mondino Foundation, Italie

10.36-10.48 **Clinical investigation coupled with molecular dissection as the personalized approach to pediatric GNAO1 encephalopathies. Drug discovery and the potential to other rare diseases.**
Vladimir Katanaev - University of Geneva, Geneva, Switzerland

10.48-11.00 **Presentation Modes in Alexander Disease : A Multicenter Retrospective Study of 94 Patients**
Emma Perrier - Service De Neuropédiatrie, Crmr Neurogénétique, Hôpital Armand Trousseau-La Roche Guyon, Aphp, Sorbonne University, Paris, France

11.00-12.00 SYMPOSIUM SPONSORISÉ PAR L'INDUSTRIE / INDUSTRY SPONSORED SYMPOSIUM

(See page 6 for the detailed program)

12.00-13.00 ***LUNCH & poster visit***

13.00-14.00 SYMPOSIUM SPONSORISÉ PAR L'INDUSTRIE / INDUSTRY SPONSORED SYMPOSIUM

(See page 6 for the detailed program)

14.00-14.30 **SESSION POSTER FLASH / FLASH POSTER SESSION**

14.00-14.06 **Acute necrotizing encephalopathy associated with RANBP2 mutation: case presentation and literature review**
Irene Peterlongo - Department of Biomedical And Clinical Sciences, Postgraduate School Of Child Neuropsychiatry, University Of Milan, Italy

14.00-14.12 **Hypomyelinating leukodystrophies associated with GJC2 gene mutations: a Tunisian case series**
Ichraf Kraoua - Lr18sp04 and Department of Pediatric Neurology. National Institute Mongi Ben Hmida of Neurology of Tunis, Tunisia

14.12-14.18 **The first Italian single center cohort of Phelan McDermid syndrome patients: clinical and genetic aspects**
Alice Decio - Irccs Medea, Italy

14.18-14.24 **Neurodevelopmental profile in children born to mothers affected by Systemic Sclerosis**
Erika Loi - Department of Clinical and Experimental Sciences, University of Brescia; Unit Of Child Neurology And Psychiatry, Asst Spedali Civili Of Brescia, Italiy

14.24-14.30 **Cognitive skills in Aicardi Goutières Syndrome**
Annamaria Del Boca - Department of Brain and Behavioral Sciences, University of Pavia, Department of Child Neurology And Psychiatry, Irccs Mondino Foundation, Italy

14.30-15.30 COMMUNICATIONS ORALES / ORAL COMMUNICATIONS

14.30-14.42 **Clinical and genetic characterization of PLP1-related disorders: preliminary insights from an Italian cohort**
Fabio Bruschi - Pediatric Neurology Unit, Coala (center For Diagnosis and Treatment of Leukodystrophies) V. Buzzi Hospital, Università Degli Studi di Milano, Italy

14.42-15.54 **Stroke and stroke-like episodes as a new clinical manifestation in GLUT1 Deficiency Syndrome**
Roberto Previtali - University of Milan, Italy

14.54-15.06 **Study of children with neurological, metabolic and/or neurosurgical pathology transferred from Antilles Guyane to a Parisian university hospital from January 1, 2018 to January 1, 2023**
Léa Lacoffrette - Department of Pediatrics, Martinique University Hospital, University of The French West Indies and French Guiana, France

15.06-15.18 **Hidden Challenges: Mental Health Outcomes in Pediatric Stroke Survivors**
Ludovica Serafini - Neurology, The Hospital for Sick Children, Toronto, Canada

15.18-30 **Developmental coordination disorder and Cerebral visual impairment: convergences and divergences**
Serena Micheletti - Unit of Child Neurology and Psychiatry, Asst Spedali Civili of Brescia, Italy

15:30-16:00 **Coffee break**

16:00-17:00 **CONFERENCES/ LECTURES**

16:00-16:30 **Pelizaeus-Merzbacher Disease and "PLPathies" from clinic to therapies**
Odile Boespflug Tanguy - *Neurodiderot Université de Paris Cité, Paris, France*

16:30-17:00 **Challenges and advances in managing Aicardì-Goutières Syndrome**
Elisa Fazzi - *Università degli Studi di Brescia, Brescia, Italy*

17:00 – 17:30 ***X-Linked Adrenoleukodystrophy: Past, Present, and Future Directions***
Caroline Sevin - Hôpital du Kremlin Bicêtre, Paris, France

17.30-18.30 **Assemblée Générale de la SENP/ SENP General Assembly**

19:00 – 20:00 **Visite Musée**

20:00 **Dîner de gala**

SAMEDI 16 MARS / SATURDAY, MARCH 16

Pediatric neuroinflammatory disorders of infectious etiology

8:00-8:30	Visite Poster/Poster visit
	CONFERENCES/ LECTURES
8:30 – 9:00	**Evolution and changing paradigms of diagnostic perspectives and patient management in pediatric infectious and post-infectious encephalitis** Yael Hacohen - *University College London, London, United Kingdom*
9:00-9:30	**From traditional pathology to modern challenges: the evolution of neuropathology in the field of infectious and parainfectious disorders of CNS** Marianna Bugiani - *Amsterdam University Medical Centers, Amsterdam, The Netherlands*
9:30-10:00	**Neuroradiological Aspects of Infectious Encephalitis: from the Typical Phenotype to the Challenge of Aspecific findings** Cecilia Parazzini - *Ospedale dei Bambini Buzzi, Milan, Italy*
10:00 -10:30	**Coffee break**
10.30-11.00	**Human genetic and immunological determinants of childhood HSV1 encephalitis: a 20-year long journey** Shen-Ying Zhang - *The Rockefeller University, New York, United States*
11:00 -11:30	**Pleomorphic neurological acute presentations of pediatric Lyme LNB and Varicella-Zoster virus: lessons from the past and on-going controversies** Joël Fluss – *Hôpitaux Universitaires Genève, Geneva, Switzerland*
11:30-12:00	**Acute flaccid myelitis: emerging polio-like disease in an era believed to be polio-free"?** Jelte Helfferich - *University Medical Center Groningen, Groningen, The Netherlands*
12:00-12:30	**Poster visit**
12:30-14:00	***LUNCH & poster visit***
14:00 – 15:00	**DISCUSSION DE CAS CLINIQUES / CLINICAL CASES DISCUSSION**
14:00 – 14:15	**Epilepsy surgery, ketogenic diet or acetazolamide in a boy with refractory epilepsy.** Benoit Semal - Université Libre De Bruxelles, Hôpital Universitaire De Bruxelles, Hôpital Universitaire Des Enfants Reine Fabiola, Department Of Genetics, Belgium
14:15 – 14:30	**Severe Inflammatory lesions with partial Kluver-Bucy Syndrome. New infectious or Autoimmune phenotype?** Gemma Olivé-Cirera - Hospital Parc Taulí De Sabadell / Hospital Clínic De Barcelona, Spain
14:30 – 14:45	**A deceiving diagnostic process in an infant with acute encephalopathy. When clinical findings, EEG and neuroimaging disagree: which path to follow?** Ylenia Vaia - Unit of Pediatric Neurology, C.o.a.l.a. (center For Diagnosis and Treatment of Leukodystrophies), V. Buzzi Children's Hospital, Università Degli Studi di Milano, Italy
14:45 – 15:00	**Subacute onset of encephalopathy with movement disorder: diagnostic challenges between old and new diagnosis** Caterina Zanus - Institute for Maternal and Child Health Burlo Garofolo, Italy

15:00-16:00 **COMMUNICATIONS ORALES / ORAL COMMUNICATIONS**

15:00-15:12 **Children with Autism Spectrum Disorder can imagine actions like their neurotypical peers?**
Jessica Galli - Department of Clinical And Experimental Sciences, University Of Brescia And Unit Of Child Neurology Psychiatry, Asst Spedali Civili Of Brescia, Brescia, Italy

15:12-15:24 **Electroclinical phenotype and possible genotype-phenotype correlations in Pallister Killian syndrome: preliminary data**
Giulia Ferrera - University Of Milan, Italy

15:24-15:36 **Early discontinuation of treatment in neonates with acute provoked seizures: a European, multicenter, retrospective study**
Margherita Bonino - Cliniques Saint-Luc U.C Louvain, Belgium

15:36-15:48 **Toward a phenotype characterization at onset in pediatric central nervous system immune-mediated disorders (pCNS-IMD): an Italian Single-Center Experience.**
Arianna Gadda – University of Milan, Milan, Italy

15:48-16:00 **Paroxysmal dyskinesia: analysis of a series of 80 children, back to the basis after genetic studies?**
Claudia Ravelli - Service De Neurologie Pédiatrique, Centre De Référence De Neurogénétique, Hôpital Armand Trousseau Ap-Hp.su, Sorbonne Université, Fhu I2-D2, France

16:00- 16:30 **Remise prix & farewell**

Symposia program

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Rare and Complex Epilepsies: Focus on Comorbidities and Cognitive Issues Moderator: Pierangelo Veggiotti

- Comorbidities in Rare and Complex Epilepsies (Pasquale Striano)
- Exploring Complexities: Cognitive Issues in Children with Epilepsy (Stefania Maria Bova)

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Seizure and Beyond: The Management of DEEs. Moderator: Aglaia Vignoli.

- Cannabidiol: a new treatment opportunity in DEEs (Francesca Ragona)
- Clinical Cases presentation (Enrico Alfei; Silvia Masnada)