

PROGRAMMA ECM IN PROGRESS

Disability Progression in MS, NMO and MOGAD: from bench to bedside

EVENTO RESIDENZIALE

Hotel Crowne Plaza VERONA

Data	20-21 MARZO 2025
Provider ECM 6207 Segreteria Organizzativa	Eolo Group Eventi srl, Via V. Veneto 11 – 35043 Monselice (PD) ID Provider 6207
Responsabile Scientifico	Prof. Massimiliano Calabrese The Multiple Sclerosis Center of University Hospital of Verona
Segreteria Scientifica	Dr. Damiano Marastoni Dr.ssa Roberta Magliozzi Dr.ssa Valentina Camera
RAZIONALE	The study of disability progression, including progression independent of relapse activity (PIRA), in multiple sclerosis (MS), neuromyelitis optica spectrum disorder (NMOSD), and myelin oligodendrocyte glycoprotein antibody-associated disease (MOGAD) is crucial for improving patient outcomes and developing targeted therapies. These disorders share a common feature of neuroinflammatory and neurodegenerative processes, but they differ significantly in their clinical course, pathophysiology, and response to treatment. Understanding the mechanisms behind disability progression in each condition is key to tailoring therapeutic strategies and predicting long-term outcomes.
	In MS, disability accumulation can occur due to both relapses and PIRA, where neurodegeneration occurs even in the absence of clinical relapses. Studying PIRA is essential as it reflects the underlying smoldering inflammation and neurodegeneration that continue despite immunomodulatory treatment. Early identification and intervention in patients at risk of progressive disability could significantly alter the disease trajectory.
	Similarly, while NMOSD and MOGAD are traditionally viewed as primarily relapse-driven diseases, emerging evidence suggests that some patients experience progressive disability even outside of acute attacks. Investigating the factors contributing to this progression, especially in PIRA-like patterns, can help distinguish subgroups of patients who may benefit from different therapeutic approaches. Moreover, identifying biomarkers that signal early progression can facilitate personalized treatment and improve long-term quality of life.





By studying disability progression and PIRA in these three diseases, we can better understand the shared and distinct pathways of neurodegeneration, which will aid in developing more effective, disease-specific treatments. This knowledge is also essential for designing clinical trials that more accurately assess treatment efficacy in preventing disability accumulation across the spectrum of these disorders.

Destinatari N.100

Medici chirurghi specializzati in neurologia, neurochirurgia, medicina generale, radiologia, geriatria, medicina interna, psichiatria, medicina fisica e riabilitazione.

Farmacisti, Logopedisti, Infermieri, Fisioterapisti, Terapisti della riabilitazione, Psicologi-Psicoterapisti, Biologi, Tecnici radiologi, Tecnici di neurofisiopatologia.

	Programma
	March 20, 2025
9.00	Registration
9.30	Welcome and Day Introduction Massimiliano Calabrese
	Session 1: NMOSD an update Chairs: Valentina Camera, Sara Mariotto
9.40	MOGAD/NMOSD: pathology Gabriele De Luca (Oxford)
10.00	NMOSD: Clinical and MRI diagnostic and prognostic features Rosa Cortese (Siena)
10.20	NMOSD: Monitoring and new and old treatments to prevent disability Ho Jin Kim (Seoul)
10.40	MOGAD/NMOSD diagnostic antibody assays and biomarkers predicting disease activity Matteo Gastaldi (Pavia)
11.00	Coffee Break and poster viewing
	Session 2: MOGAD an update Chairs: Alberto Gajofatto, Carla Tortorella
11.30	MOGAD: Clinical and MRI diagnostic and prognostic features Georgina Arrambide (Barcellona)
11.50	Novel insight in disability outcomes in MOGAD and NMOSD clinical trials Ruth Geraldes (Oxford)
12.10	MOGAD: Monitoring and new and old treatments to prevent disability Jackie Palace (Oxford)
12.45	Light Lunch and poster viewing
14.00	Greetings from Authorities



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t./f. 0429 767381

c. 392 6979059



	Session 3: Disability Progression in MS from neuropathology to Advanced MRI
	Chairs: Massimiliano Calabrese, Nicola De Stefano
14.20	Advanced MRI: from histopathology of progressive disease to novel biomarkers Cristina Granziera (Basel)
14.50	Chronic Active Lesions from neuropathology to MRI Martina Absinta (Bethesda)
15.20	Advanced Imaging Techniques for Assessing Remyelination in Multiple Sclerosis Alessandro Cagol (Basel)
15.50	Coffee Break and poster viewing
	Session 4: From advanced MRI to clinical applications Chairs: Francesca Benedetta Pizzini, Claudio Gasparini
16.30	Typical and atypical MRI lesions in MS Alex Rovira (Barcellona)
17.00	Translation of Advanced MRI to clinical application Agnese Tamanti (Verona)
17.30	Quantifying neurodegeneration: innovative tools and techniques in neurodegenerative diseases Federica Agosta (Milan)
18.00	Conclusion of the day
	March 21, 2025
9.00	Registration
9.40	Introduction of the day Massimiliano Calabrese
	Session 5: The concept of PIRA Chairs: Antonio Scalfari, Pietro Annovazzi
9.45	Keynote Lecture: What is MS and how we should treat it Massimo Filippi (Milan)
10.15	PIRA – Vino Vecchio in Botti Nuove? Ludwig Kappos (Basel)
10.45	Cognitive PIRA Stefano Ziccardi (Verona)
11.10	Coffee Break and poster viewing
	Session 6: MS from neuropathology to fluid biomarkers Chairs: Roberta Magliozzi, Bruno Bonetti



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11.40	Neuropathological Mechanisms of disease progression Simon Hametner (Wien)
12.05	Meningeal Inflammation in MS: an update Richard Reynolds (London)
12.30	Single-Cell Transcriptomics in Progressive Patients: Decoding the Cellular Landscape Damiano Marastoni (Verona)
12.55	Fluid Biomarkers (CSF and blood) of Progressive MS Massimiliano Di Filippo (Perugia)
13.20	Light Lunch and poster viewing
	Session 7: Progressive MS: from AI to personalized treatment Chairs: Massimiliano Calabrese, Paolo Preziosa
14.20	Keynote Lecture: How AI can help us Isabella Castiglioni (Milano)
14.50	Is it possible to predict the progression? Tom Fuchs (Amsterdam)
15.15	Smoldering MS: from immunology to treatment Roberto Furlan (Milan)
15.40	Poster awards Chairs: V. Camera, D. Marastoni, R. Orlandi
16.00	Conclusion Remarks and Welcome to Verona 2026
16.30	Compilazione del questionario ECM e questionario di gradimento

Ore Formative

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