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Associazione Malattie Rare
"Mauro Baschirotto" Onlus
Sezione Umbra

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Fondazione Malattie Rare
«Mauro Baschirotto»

Workshop

Gene Therapy for Lafora Disease: State of the Art and Emerging Perspectives

Perugia, 27th February 2026

*Section of Neurology – Neurophysiopatologia,
Department of Medicine and Surgery,
University of Perugia*

*Room 9, Building B, floor -2,
Gambuli Square 1, Perugia*



SCIENTIFIC RATIONALE

Lafora disease is a rare neurodegenerative disease associated with progressive myoclonus epilepsy, for which no disease-modifying therapy is currently available. Over the past few years, substantial progress has been made in understanding its molecular mechanisms, developing advanced experimental models, and exploring innovative therapeutic strategies, particularly in the field of gene therapy.

This workshop aims to bring together international experts working across basic, preclinical, and clinical research to critically review the state of the art in Lafora disease research. Special emphasis will be placed on experimental models, pathogenic mechanisms, preclinical proof-of-concept studies, translational biomarkers, and future clinical applications. By stimulating dialogue among scientists, clinicians, patient organizations, and families, the event seeks to promote knowledge exchange, collaboration, and the translation of research advances into tangible clinical benefit.

Scientific director

Cinzia Costa

Scientific secretariat

Lorenzo Gaetani; Valentina Imperatore; Andrea Mancini; Miriam Sciacaluga

FACULTY

Laura Bellingacci

Researcher, Department of Medicine and Surgery, University of Perugia

Barbara Bettegazzi

Researcher, Vita-Salute San Raffaele University, Milan

Barbara Cellini

Associate Professor, Department of Medicine and Surgery, University of Perugia

Davide Chiasserini

Associate Professor, Department of Medicine and Surgery, University of Perugia

Cinzia Costa

Associate Professor, Head of Neurophysiopathology Unit, Department of Medicine and Surgery, University of Perugia

Lorenzo Gaetani

Neurologist, Neurology Unit, Perugia University Hospital

Nerea Iglesias-Cabeza

Researcher, Instituto de Investigación Sanitaria-Fundación Jiménez Díaz, Universidad Autónoma de Madrid

Valentina Imperatore

Researcher, Department of Medicine and Surgery, University of Perugia

Andrea Mancini

Neurologist, Neurology Unit, Perugia University Hospital

Sabata Martino

Full Professor, Department of Chemistry, Biology, and Biotechnology, University of Perugia

Lucilla Parnetti

Full Professor, Head of Neurology Unit, Department of Medicine and Surgery, University of Perugia

Paolo Prontera

Head of Medical Genetics and Rare Diseases Unit, Perugia University Hospital

Marina P. Sánchez

Researcher, Instituto de Investigación Sanitaria-Fundación Jiménez Díaz, Universidad Autónoma de Madrid

José M. Serratos

Head of Epilepsy Unit, Instituto de Investigación Sanitaria-Fundación Jiménez Díaz, Universidad Autónoma de Madrid

Miriam Sciacaluga

Associate Professor, Department of Life Science, Health, and Health Professions, Link University, Rome

Luis Zafra-Puerta

Researcher, Instituto de Investigación Sanitaria-Fundación Jiménez Díaz, Universidad Autónoma de Madrid

Program

12:30 – 13:30 | Registration

13:30 – 14:00 | Welcome

14:00 – 14:30 | Institutional Remarks

Introduction: *L. Parnetti – C. Costa*

14:30 - 15:30 Session I – Clinical Context and Molecular Profiling in Lafora Disease

Chairs: C. Costa – B. Cellini

From patient to gene and back to the patient

J. M. Serratosa

Biochemical pathways in Lafora Disease

D. Chiasserini

Comparative Electrophysiological Characterization of Lafora Disease Models

L. Bellingacci

Discussion

15:30 – 16:50 Session II – Gene Therapy: Preclinical Proof-of-Concept and Key Findings

Chairs: M. Sciaccaluga – B. Bettegazzi

Intracerebroventricular EPM2A and EPM2B therapies

M. P. Sánchez

Optimizing EPM2A therapy through intraparenchymal administration

N. Iglesias-Cabeza

Intravenous delivery of EPM2A and EPM2B genes

L. Zafra Puerta

Gene therapy and neuronal function: electrophysiological insights from preclinical models

M. Sciaccaluga

Discussion

16:50 – 17:10 Coffee break

Session III – Translational Integration and Future Perspectives

17:10 – 18:10

Chairs: C. Costa – P. Prontera

From Preclinical Models to Clinical Perspectives in Lafora Disease

A. Mancini

Fluid Biomarkers of Neurodegeneration in Lafora Disease

L. Gaetani

Gene Therapy for Monogenic Disease: What Are We Learning?

S. Martino

Discussion

Community Voices - Final Discussion with Families and Associations

18:10 – 18:40

G. Baschirotto – M. Gatti