



## Seconda Conferenza Europea GNAO1 (2020)

### 1 OCTOBER 2020 | RESEARCH UPDATES SESSION

**Kirill Martemyanov** | Scripps Research

*Insights into GNAO1 function in the brain and pathogenic mechanisms of its disease variants*

**Vladimir Katanaev** | University of Geneva

*GNAO1 encephalopathy: molecular etiology and animal models.*

**Simone Martinelli** | Italian National Institute of Health

*Modelling GNAO1-related diseases in *Caenorhabditis elegans**

**Frank Baas** | Leiden University

*Gene Identification and Studies in Neurodevelopmental Disorders*

**Maryana Bardina** | Marlin Biotech

**Timofei Zatsepin** | Skolkovo Institute of Science and Technology

**Alexey Wolfson** | Advirna

*Early development of antisense oligonucleotide therapeutics to treat G203R GNAO1 encephalopathy*

### 2 OCTOBER 2020 | CLINICIANS/RESEARCHERS ROUND TABLE

In this session clinicians and researchers who have experience with GNAO1 gene mutation will have the chance to share knowledge and ideas with each other. The aim is to establish a network of experts from all over the world and facilitate the constitution of international teams who will work together in order to better understand the neurological mechanisms underlying the disease and define the next steps to take to move the scientific research forward.

### 3 OCTOBER 2020 | CLINICAL/REHABILITATION SESSION

**Amy Viehoveer** | Washington University in St. Louis

*GNAO1 Natural History Study (title TBC)*

**Serena Galosi** | Sapienza University of Rome

*GNAO1 related disorders: update on clinical spectrum*

**Erika Axeen** | University of Virginia

*Preliminary results of the GNAO1 International Registry*

**Giovanna Zorzi** | Foundation I.R.C.C.S Carlo Besta Neurological Institute

*Update on treatment options in GNAO1-associated movement disorder*

**Alessandro Capuano** | Bambino Gesù Children's Hospital

**Alice Bergonzoli** | Sirio Medical

*Eye-tracking for communication skills assessment and training in GNAO1 patients: a single center experience*

**Karen Balk** | Washington University in St. Louis

*Occupational Therapy strategies for children affected by GNAO1 gene mutation (title TBC)*

## Relatori

**Kirill Martemyanov**

*Associate Professor, Department of Metabolism & Aging Faculty*

*Chair of Department of Neuroscience*

*Scripps Research, Florida Campus*

**Vladimir Katanaev**

*Full Professor and Chair in Translational Oncology*

*Department of Cell Physiology and Metabolism*

*Translational Research Centre in Oncohaematology*

*Faculty of Medicine, University of Geneva*

**Simone Martinelli**

*In-staff scientist, Department of Oncology and Molecular Medicine*

*Istituto Superiore di Sanità (ISS)*

**Frank Baas**

*Head of Genome Diagnostics - Department of Clinical Genetics*

*Leiden University Medical Centre (LUMC)*

**Maryana Bardina**

*Research Fellow*

*Lab. of modeling and gene therapy of hereditary diseases, Group Leader*

*Marlin Biotech*

**Timofei Zatsepin**

*Associate Professor, Center of Life Sciences  
Skolkovo Institute of Science and Technology*

**Alexey Wolfson**

*CEO Advirna*

**Amy Viehoveer**

*Assistant Professor, Neurology, Washington University in St. Louis  
Division of Pediatric Neurology St. Louis Children's Hospital*

**Serena Galosi**

*Department of Human Neuroscience  
Sapienza Università di Roma  
Umberto I - Policlinico di Roma*

**Erika Axeen**

*Pediatric neurologists  
University of Virginia Medical Center*

**Giovanna Zorzi**

*U.O.C. Genetica Medica e Neurogenetica  
Dipartimento di Diagnostica e Tecnologia Applicata  
Fondazione IRCCS Istituto Neurologico "Carlo Besta"*

**Alessandro Capuano**

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Ospedale Pediatrico Bambino Gesù di Roma*

**Alice Bergonzoli**

*TNPEE - Esperta in Comunicazione Aumentativa e Alternativa  
Sirio Medical*

**Karen Balk**

*Occupational Therapist  
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