# <u>Agenda</u>

#### 08:30 - 09:00 - Gather together and On-site Registration

#### <u>09:00 – 09:10 - Welcome and Introduction</u>

Sumathi Iyengar & Amir Kedar, WAS organizations and Anna Villa, IRCCS Ospedale San Raffaele, Milan, Italy.

#### <u>09:10 - 09:20 - Families Story</u>

Family story - Gene Therapy - Valentina Pagani, Rosa Giglia.

#### 09:20 - 09:50 - Plenary Session A

Alessandro Aiuti - head of the Pediatric Immunohematology Unit at the IRCCS San Raffaele, Milano, Italy.

"The long path of gene therapy for Wiskott-Aldrich Syndrome"

#### <u>09:50 - 10:35 - Gene Therapy Session</u>

**Chair: Fabio Candotti** 

09:50 - 10:05 Francesca Ferrua - San Raffaele Telethon Institute for Gene Therapy, Italy.

Safety and efficacy of lentiviral hematopoietic stem cell Gene Therapy for Wiskott-Aldrich Syndrome

10:05 - 10:20 Maria Carmina Castiello - San Raffaele Telethon Institute for Gene Therapy, Italy.

Long-term efficacy of lentiviral vector-mediated gene therapy in restoring B-cell homeostasis in Wiskott-Aldrich syndrome patients

**10:20 - 10:35 Michaela Semeraro -** Centre d'Investigations Cliniques, INSERM CIC1419, Hôpital Necker Enfants malades - Université de Paris-Cité

Long-term Inflammatory manifestations post lentiviral hematopoietic stem/progenitor cell gene Therapy for Wiskott Aldrich syndrome

#### 10:35 - 11:05 - Gene Therapy Panel

**Chair:** Emma Morris

Participants: Alessandro Aiuti, Marina Cavazzana, Luigi Notarangelo, Suhag Parikh.

"Lentiviral Gene Therapy for WAS, why is there variability in outcome and how we can address it?"

#### 11:05 - 11:20 - Break

# 11:20 - 11:25 - Families Story

**Family story** – Gene Therapy – Priya Stephan (Short video)

#### 11:25 - 11:55 - Plenary Session B

Alessia Cavazza - Infection, Immunity & Inflammation Department, UCL Institute of child health, London, UK. Development of a safe and efficient CRISPR/Cas gene editing platform for the treatment of Wiskott-Aldrich Syndrome

#### 11:55 - 12:30 -Clinical International Session I

**Chair:** Anders Fasth

11:55 - 12:05 Carmem Bonfim- Pediatric Transplant and Cellular Therapy Program - Duke University, NC, USA.

Haploidentical cell transplantation using post-transplantation cyclophosphamide for patients with Wiskott-Aldrich: Excellent outcomes and emerging challenges



12:05 - 12:15 Deepti Suri - Department of Pediatrics, Postgraduate Institute of Medical Education and Research, Chandigarh, India

Correlation of Clinical Profiles, Immune Cells, Genotypes, and Regulatory Gene Expression in Indian Patients with XLT/WAS

12:15 - 12:25 Anahita Razaghian - IAARI and Hakim children hospital, Teheran, Iran. Online presentation
Clinical features and outcomes of 39 Wiskott-Aldrich syndrome patients from a single center of Iran

12:25 - 12:30 Questions and answers and discussion

12:30 - 12:40 - Families Story

Young Adult Family story – Emily Kinsel (Short video)

#### 12:40 - 13:10 - Plenary Session C - Panel Discussion

**Moderator:** Luigi Notarangelo - Immune Deficiency Genetics Section, National Institute of Health, USA. Participants: Alessandro Aiuti, Michael Albert, Fabio Candotti, Sumathi Iyengar, Emma Morris

"Evolving aspects of WAS: milder forms of the disease and carrier status – from disease perception to clinical implications"

# 13:10 - 14:10 - Lunch

#### 14:10 - 15:10 - Clinical Session II

**Chair:** Carmem Bonfim

14:10 - 14:30 Emma Morris - Invited speaker, Director, UCL Division of Infection and Immunity, London, UK. "Allogeneic HSCT for adults with WAS"

14:30 - 14:45 Tanja C. Vallée - Stem cell transplantation unit, Dr. von Haunersches University Children's Hospital, Munich Allogeneic HSCT with adjusted busulfan dosing results in excellent survival and cure rates for WAS

14:45 - 15:00 Yoji Sasahara – Dep. of Pediatrics & Pediatric Oncology, Tohoku University Graduate School of Medicine Efficacy of rituximab for the treatment and prevention of autoimmunity in patients with WAS and XLT

15:00 - 15:10 Questions and answers and discussion

#### 15:10 - 15:45 - Clinical Session III

Chair: Suhag Parikh

**15:10 - 15:25 Maddalena Migliavacca -** Pediatric Immunohematology and Bone Marrow Transplantation Unit, IRCCS San Raffaele Scientific Institute, Milan, Italy.

New insights into Wiskott Aldrich Syndrome carriers: A multicentric observation

15:25 - 15:40 Markus Walz - Associate Partner, Magnolia Innovation, New Jersy, USA.

An International Study of Patient and Caregiver-Reported Burden and Quality of Life in Wiskott-Aldrich Syndrome

15:40 - 15:45 Questions and answers and discussion

#### <u> 15:45 – 16:20 – Clinical Session Panel</u>

**Chair: Michael Albert** 

Participants: Alessandro Aiuti, Siobhan Burns, Fabio Candotti, Luigi Notarangelo



'Towards a consensus statement in the management of WAS'

# 16:20- 16:35 - Break

#### 16:35 - 17:10 - Research Session I

Chair: Anna Villa

**16:35 - 16:50 Rhaissa Vieira** - Department of Microbiology Tumor and Cell Biology, Karolinska Institutet, Sweden.

Therapeutic Candidate for Enhancing Platelet Production and Function in Wiskott-Aldrich Syndrome and X-Linked Thrombocytopenia

16:50 - 17:05 Lia Pinho - Department of Microbiology Tumor and Cell Biology, Karolinska Institutet, Sweden.

From Structure to Function: Understanding the impact of WASp Mutations on Actin Dynamics and T **Cell Activation** 

17:05 - 17:10 Questions & answers and discussion

#### 17:10 - 18:20 - Research Session II

Chair: Alessandro Aiuti

17:10 - 17:30 Yatin Vyas - Invited speaker, Professor and Chair, Department of Pediatrics, Penn State Health Children's Hospital, USA.

Functional pairing of WASp and Fanconi anemia proteins in the cell nucleus- co-opting cosmic quantum entanglement

17:30 - 17:45 Roberto Amadio - Cellular Immunology department, University of Trieste and ICGEB Trieste, Italy.

Actin polymerization by WASp limits nuclear envelope rupture and inflammation in macrophages

17:45 - 18:00 Jordan Chill - Department of Chemistry, Bar Ilan University, Israel.

Curation of a Structural WAS/XLT Database: Challenges and First Conclusions

18:00 - 18:15 Gianluca Baldanzi - Department of Translational Medicine, University of Piemonte Orientale, Novara, Italy.

DGKa is a negative regulator of megakaryopoiesis inhibited by WASp

18:15 - 18:20 Questions & answers and discussion

18:20 - 18:30 - Closing and Summary of the day Anna Villa, Fabio Candotti, Sumathi Iyengar, Amir Kedar

### 18:30 - 19:15 - Reception

An informal get-together to renew acquaintances and meet new colleagues will be held at the Meeting Venue. All registered participants are invited to attend.



The Wiskott Aldrich Syndrome Association (R.A)







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