**Agenda**

6:15 AM Welcome and Introductions  
*Uma Ramaswami, Chair*

6:20 AM Early diagnosis: The need, the challenges and new tools  
*Roberto Giugliani*

6:35 AM Monitoring and management: Guidelines, gaps and good news  
*Ozlem Goker-Alpan*

6:50 AM A rapidly evolving treatment landscape: New approvals, clinical trials, the future of Fabry disease and you  
*Dominique Germain*

7:05 AM Panel discussion and audience Q&A

**Target Audience**

This educational activity is designed to deliver up-to-date education to pediatricians, geneticists, internal medicine and family practice specialists, nephrologists, nurse practitioners, physician assistants, nurses, genetic counselors and other healthcare providers who manage patients with Fabry disease.

**Learning Objectives**

At the end of this session, the participants will be able to:

1. Delineate issues in diagnosis of Fabry disease patients.
2. Utilize current recommendations for managing patients with Fabry disease.
3. Review current and emerging therapies for Fabry disease.

**Facility**

Uma Ramaswami, FRCPCH, MD (Chair)  
Consultant in Inherited Metabolic Disorders
Clinical Lead, Lysosomal Disorders Unit
Institute of Immunity and Transplantation
Royal Free London NHS Foundation Trust  
London, United Kingdom

Dominique Germain, MD, PhD  
Professor of Medical Genetics,  
University of Versailles - Paris Saclay University  
Head of Division of Medical Genetics,  
Director - Referral Center for Fabry disease  
Raymond Poincare Teaching Hospital, Garches, France

Roberto Giugliani, MD, PhD  
Professor, Department of Genetics, UFRGS  
Founder and Member, Medical Genetics Service, HCPA  
Coordinator, Brazilian Institute of Population Medical Genetics, INAGEMP  
Porto Alegre, Brazil

Ozlem Goker-Alpan, MD  
Founder and Chief Medical Officer  
Lysosomal & Rare Disorders Research & Treatment Center  
Fairfax, Virginia, USA

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