



UNLOCKING THE POWER OF THE PATIENT VOICE

Holistic care for patients with
Fabry disease and alpha-mannosidosis

JOIN US TO HEAR THE PANEL'S PERSPECTIVE

Thursday, February 8 from 5:15 – 6:15 pm
Manchester Grand Hyatt San Diego, Grand Hall AB

In this symposium, the presenters will discuss **patient and caregiver perception of disease burden, monitoring, and management** of lysosomal disorders, with a focus on **Fabry disease and alpha-mannosidosis**.

Speakers will provide insights on ways to incorporate patient perspectives in clinical decision-making to adopt a holistic patient-care approach. The presentation will include case studies and a dynamic panel discussion with patient advocates for Fabry disease and alpha-mannosidosis. The symposium will end with a Q&A session

SPEAKERS



Lisa Berry
Genetic Counselor
Rare Genetic Disease Program
Cincinnati Children's Hospital



Prof. Julia B. Hennermann
Head of the Villa Metabolica
(Department of Inborn Errors of Metabolism)
University Medical Center Mainz



Sabina Kineen
Patient advocate
Fabry Disease



Sophie Thomas
Patient advocate
MPS and related lysosomal disorders

This session is sponsored by Chiesi Global Rare Diseases.
This is not a continuing education (CE) activity.

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