Optimizing early diagnosis in lysosomal storage diseases and innovation for the future
*Chair: Julia Hennermann, Mainz, Germany*

The importance of early diagnosis in Fabry disease and potential innovative approaches
*Julia Hennermann, Mainz, Germany*

Screen4Care: shortening the path to rare disease diagnosis using newborn genetic screening and digital technologies
*Edith (Sky) Gross, Paris, France*

Early diagnosis of leukodystrophies: recognizing key signs and symptoms
*Ingeborg Krägeloh-Mann, Tübingen, Germany*

Newborn screening with a focus on mucopolysaccharidoses (MPS)
*Shuan-Pei Lin, Taipei, Taiwan*

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