Takeda, Thursday September 1, 2022, 12.45-13.45, Hall Porto

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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