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Where the world comes for answers

MPS Awareness Webinar

An Overview of Mucopolysaccharidoses for Families and Providers
June 9, 2023 from Noon–1 PM (EST)

[Click here to join meeting – preregistration not required: JOIN MPS WEBINAR](#)

NERGG, the Boston Children's Hospital Lysosomal Disorders (BoLD) Program, and the National MPS Society are pleased to present this free presentation on ZOOM. Join us as we discuss various types of conditions in this group of inherited metabolic disorders. This information-packed session will include:

- **Overview of the Diagnosis, Clinical Challenges, and Management of MPS**
 - Walla Al-Hertani, MD, MSc, FRCPC, FCCMG, FACMG
Harvey Levy Chair and Director of the Metabolism Program, Lysosomal Disorders (BoLD) Program and the Glycogen Storage Diseases Program
Division of Genetics and Genomics, Boston Children's Hospital
- **Resources Offered by the National MPS Society**
 - Leslie Urdaneta, MSW, LCSW, Director of Family Support and Communication
 - Carol Bryant, MSW, LCSW, Pathways Program Director
- **A Family's Experience with Sanfilippo Syndrome (MPS III)**
 - Joanne Huff, BSN, RN New Hampshire Consumer Representative on the NERGG Board of Directors, and the parent of an adult daughter with MPS III
- **Supporting a Patient's Journey**
 - Kathy Cody, RD, Account Manager Skeletal Dysplasia (MPS & Achondroplasia)
BioMarin Pharmaceutical Inc.
 - Speaker from Takeda Pharmaceutical Company

For any questions, please contact NERGG's Executive Director [Cindy Ingham](#).

We look forward to seeing you and sharing information about MPS!

BIO·MARIN

Many thanks to our exhibitors

