Mucopolysaccharidosis type 1

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Mucopolysaccharidosis type 1. ORPHA:579

Mucopolysaccharidosis type 1 (MPS 1) is a rare lysosomal storage disease belonging to the group of mucopolysaccharidoses. There are three variants, differing widely in their severity, with Hurler syndrome being the most severe, Scheie syndrome the mildest and Hurler-Scheie syndrome giving an intermediate phenotype.