Mucopolysaccharidosis type 6

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Mucopolysaccharidosis type 6. ORPHA:583

Mucopolysaccharidosis type 6 (MPS 6) is a lysosomal storage disease with progressive multisystem involvement, associated with a deficiency of arylsulfatase B (ASB) leading to the accumulation of dermatan sulfate.