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Mucopolysaccharidosis type 4

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base.

Mucopolysaccharidosis type 4. ORPHA:582

Mucopolysaccharidosis type IV (MPS IV) is a lysosomal storage disease belonging to the group of mucopolysaccharidoses, and characterised by spondylo-epiphyso-metaphyseal dysplasia. It exists in two forms, A and B.