Mucolipidosis type IV

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Mucolipidosis type IV. ORPHA:578

Mucolipidosis type IV (ML IV) is a lysosomal storage disease characterised clinically by psychomotor retardation and visual abnormalities including corneal clouding, retinal degeneration, or strabismus.