GM1 gangliosidosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. GM1 gangliosidosis. ORPHA:354

GM1 gangliosidosis is a rare lysosomal storage disorder characterized biochemically by deficient beta-galactosidase activity and clinically by a wide range of variable neurovisceral, ophthalmological and dysmorphic features.