GM1 gangliosidosis type 3

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. GM1 gangliosidosis type 3. ORPHA:79257

GM1 gangliosidosis type 3 is a mild, chronic, adult form of GM1 gangliosidosis (see this term) characterized by onset generally during childhood or adolescence and by cerebellar dysfunction.