GM1 gangliosidosis type 1

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. GM1 gangliosidosis type 1. ORPHA:79255

GM1 gangliosidosis type 1 is the severe infantile form of GM1 gangliosidosis (see this term) with variable neurological and systemic manifestations.