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GM2 gangliosidosis, AB variant

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. [GM2 gangliosidosis, AB variant](#). ORPHA:309246

GM2 gangliosidosis, AB variant is an extremely rare, severe genetic disorder characterized by progressive neurological decline due to ganglioside activator deficiency.