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# Tay-Sachs disease

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Tay-Sachs disease. ORPHA:845*

GM2 gangliosidosis, variant B or Tay-Sachs disease is marked by accumulation of G2 gangliosides due to hexosaminidase A deficiency.