

Open Peer Review on Qeios

Sandhoff disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Sandhoff</u> <u>disease</u>. ORPHA:796

Sandhoff disease is a lysosomal storage disorder from the GM2 gangliosidosis family and is characterised by central nervous system degeneration.

Qeios ID: M9EEMT · https://doi.org/10.32388/M9EEMT