Open Peer Review on Qeios

STT3A-CDG

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. STT3A-CDG. ORPHA:370921

STT3A-CDG is a form of congenital disorders of N-linked glycosylation characterized by developmental delay, intellectual disability, failure to thrive, hypotonia and seizures. STT3A-CDG is caused by mutations in the gene STT3A (11q23.3).