DDOST-CDG

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. DDOST-CDG. ORPHA:300536

DDOST-CDG is a form of congenital disorders of N-linked glycosylation characterized by failure to thrive, developmental delay, hypotonia, strabismus and hepatic dysfunction. The disease is caused by mutations in the gene DDOST (1p36.1).