MPDU1-CDG

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. MPDU1-CDG. ORPHA:79323

The CDG (Congenital Disorders of Glycosylation) syndromes are a group of autosomal recessive disorders affecting glycoprotein synthesis. CDG syndrome type If is characterised by psychomotor delay, seizures, failure to thrive, and cutaneous and ocular anomalies.