COG8-CDG

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. COG8-CDG. ORPHA:95428

The CDG (Congenital Disorders of Glycosylation) syndromes are a group of autosomal recessive disorders affecting glycoprotein synthesis. CDG syndrome type IIh is characterised by severe psychomotor retardation, failure to thrive and intolerance to wheat and dairy products.