COG5-CDG

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. COG5-CDG. ORPHA:263487

COG5-CDG is an extremely rare form of CDG syndrome (see this term) characterized clinically in the single reported case to date by moderate mental retardation with slow and inarticulate speech, truncal ataxia, and mild hypotonia.