COG4-CDG

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. COG4-CDG. ORPHA:263501

COG4-CDG is an extremely rare form of CDG syndrome (see this term) characterized clinically in the single reported case to date by seizures, some dysmorphic features, axial hyponia, slight peripheral hypertonia and hyperreflexia.