Open Peer Review on Qeios

COG1-CDG

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. COG1-CDG. ORPHA:263508

COG1-CDG is an extremely rare form of CDG syndrome (see this term) characterized clinically in the few cases reported to date by variable signs including microcephaly, growth retardation, psychomotor retardation and facial dysmorphism.