DISEASE:
COG1-CDG

NAME:	COG1-CDG
DESCRIPTION:	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.
ORPHACODE:	263508
SYNONYMS:	CDG syndrome type IIg CDG-IIg CDG2G Carbohydrate deficient glycoprotein syndrome type IIg Congenital disorder of glycosylation type 2g Congenital disorder of glycosylation type IIg
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	COG1
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