DISEASE:
COG5-CDG

NAME:	COG5-CDG
DESCRIPTION:	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.
ORPHACODE:	263487
SYNONYMS:	CDG syndrome type IIi CDG-IIi CDG2I Carbohydrate deficient glycoprotein syndrome type IIi Congenital disorder of glycosylation type 2i Congenital disorder of glycosylation type IIi
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	COG5
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