DISEASE:
MORM syndrome

NAME:	MORM syndrome
DESCRIPTION:	A rare genetic syndromic intellectual disability characterized by language delay and mild to moderate intellectual disability associated with truncal obesity, congenital nonprogressive retinal dystrophy with poor night vision and reduced visual acuity, and micropenis in males. Cataracts may occur in the second or third decade of life.
ORPHACODE:	75858
SYNONYMS:	Intellectual disability-truncal obesity-retinal dystrophy-micropenis syndrome
XREF(S):	Orphanet ICD-10 MeSH OMIM
ANALYTE(S):	INPP5E
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

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