# MORM syndrome

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. MORM syndrome. ORPHA:75858*

MORM syndrome is characterised by the association of intellectual deficit, truncal obesity, retinal dystrophy and micropenis. It has been described in 14 individuals from a consanguineous family. It is transmitted in an autosomal recessive manner. The causative locus has been mapped to chromosome region 9q34.