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# X-linked intellectual disability, Golabi-Ito-Hall type

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. X-linked intellectual disability, Golabi-Ito-Hall type. ORPHA:93947*

Golabi-Ito-Hall syndrome is an X-linked intellectual disability syndrome (XLMR) characterized by intellectual deficiency, microcephaly and short stature. It belongs to the group of disorders collectively referred to as Renpenning syndrome.