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Autosomal recessive primary microcephaly

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Autosomal recessive primary microcephaly</u>. ORPHA:2512

Autosomal recessive primary microcephaly (MCPH) is a rare genetically heterogeneous disorder of neurogenic brain development characterized by reduced head circumference at birth with no gross anomalies of brain architecture and variable degrees of intellectual impairment.

Qeios ID: B2M6RE · https://doi.org/10.32388/B2M6RE