Lissencephaly with cerebellar hypoplasia type E

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Lissencephaly with cerebellar hypoplasia type E. ORPHA:100015

A rare, genetic, lissencephaly with cerebellar hypoplasia subtype characterized by the presence of lissencephaly with an abrupt transition, near the boundary between the frontal and parietal cortex, from frontal agyria to posterior gyral simplification, associated with cerebellar hypoplasia which predominantly affects the midline vermis.