Lissencephaly with cerebellar hypoplasia type A

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Lissencephaly with cerebellar hypoplasia type A. ORPHA:100011

A rare, genetic, lissencephaly with cerebellar hypoplasia subtype characterized by classical lissencephaly with thickened cortical gray matter (with either no discernable gradient, a predominantly posterior gradient, or a predominantly anterior gradient) associated with variable, predominantly midline, cerebellar hypoplasia.