Lissencephaly syndrome, Norman-Roberts type

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Lissencephaly syndrome, Norman-Roberts type. ORPHA:89844

Lissencephaly syndrome, Norman-Roberts type is characterised by the association of lissencephaly type I with craniofacial anomalies (severe microcephaly, a low sloping forehead, a broad and prominent nasal bridge and widely set eyes) and postnatal growth retardation.