Microlissencephaly

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Microlissencephaly. ORPHA:1083

Microlissencephaly describes a heterogenous group of a rare cortical malformations characterized by lissencephaly in combination with severe congenital microcephaly, presenting with spasticity, severe developmental delay, and seizures and with survival varying from days to years.