Hereditary neurocutaneous malformation

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hereditary neurocutaneous malformation. ORPHA:1062

Hereditary neurocutaneous angioma is characterised by the association of cerebral and cutaneous angiomatous lesions. It has been described in less than 10 families. Clinical manifestations of the cerebral lesions include epilepsy, cerebral haemorrhage, and focal neurological deficit. Transmission is autosomal dominant.