Sturge-Weber syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Sturge-Weber syndrome. ORPHA:3205

Sturge-Weber syndrome (SWS) is a rare congenital neurocutaneous disorder characterized by facial capillary malformations and/or cerebral and ocular ipsilateral vascular malformations that result in variable degrees of ocular and neurological anomalies.