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Familial cervical artery dissection

INSERM

Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base*. [Familial cervical artery dissection](#). ORPHA:36382

Familial cervical artery dissection is a rare, genetic, neurological disorder characterized by dissection of the cervical artery in various members of a single family, presenting with variable manifestations which range from asymptomatic to the triad of ipsilateral pain in the head, neck, and face, Horner syndrome, and cerebral or retinal ischemic symptoms. Headache and cerebral ischemic features are most frequently observed.