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Familial or sporadic hemiplegic migraine

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Familial or sporadic hemiplegic migraine</u>. ORPHA:569

Hemiplegic migraine (HM) is a rare variety of migraine with aura characterized by the presence of a motor weakness during the aura. Hemiplegic migraine has two main forms depending on the familial history: patients with at least one first- or second-degree relative who has aura including motor weakness have familial hemiplegic migraine (FHM); patients without such familial history have sporadic hemiplegic migraine (SHM).

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