Familial recurrent peripheral facial palsy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial recurrent peripheral facial palsy. ORPHA:2809

Familial recurrent peripheral facial palsy is a rare peripheral neuropathy characterized by an acute onset of unilateral facial muscle weakness with Bell's phenomenon. It is non-progressive, resolves spontaneously, and it might be recurrent with no obvious precipitating factors.