Laurence-Moon syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Laurence-Moon syndrome. ORPHA:2377

Laurence-Moon syndrome (LMS) is a very rare genetic multisystemic disorder characterized by pituitary dysfunction, ataxia, peripheral neuropathy, spastic paraplegia, and chorioretinal dystrophy.