Moynahan syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Moynahan syndrome. ORPHA:2574

A rare, genetic, epilepsy syndrome characterized by congenital alopecia, early-onset epilepsy, intellectual disability and speech delay. Large stature, delayed bone development and abnormal electroencephalogram have also been associated.