Familial infantile myoclonic epilepsy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial infantile myoclonic epilepsy. ORPHA:352582

A rare, genetic, infantile epilepsy syndrome disease characterized by neonatal- to infancy-onset myoclonic focal seizures occurring in various members of a family, associated in some with mild dysarthria, ataxia and borderline-to-moderate intellectual disability.