Myoclonic epilepsy in non-progressive encephalopathies

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Myoclonic epilepsy in non-progressive encephalopathies. ORPHA:86913

Myoclonic epilepsy in non-progressive encephalopathies is a rare epilepsy syndrome characterized by recurrent, long-lasting myoclonic status in infants and young children with a non-progressive encephalopathy, associated with transient and recurring motor, cognitive and/or behavioral disturbances.