Myoclonic-astastic epilepsy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Myoclonic-astastic epilepsy. ORPHA:1942

Myoclonic Astatic Epilepsy (MAE) is a rare epilepsy syndrome of childhood characterized by the occurrence of multiple different seizure types including myoclonic-astatic, generalized tonic-clonic and absence seizures, usually in previously healthy children.