Benign familial infantile epilepsy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Benign familial infantile epilepsy. ORPHA:306

Benign familial infantile epilepsy (BFIE) is a genetic epileptic syndrome characterized by the occurrence of afebrile repeated seizures in healthy infants, between the third and eighth month of life.