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Landau-Kleffner syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. [Landau-Kleffner syndrome](#). ORPHA:98818

Landau-Kleffner syndrome (LKS) is an age-related epileptic encephalopathy where developmental regression occurs mainly in the language domain and the electroencephalographic (EEG) abnormalities are mainly localized around the temporal-parietal regions. The term acquired epileptic aphasia describes the main features of this condition.