Opis choroby *

Definicja

A rare inflammatory and autoimmune disease with epilepsy characterized by unilateral hemispheric atrophy, associated with drug-resistant focal epilepsy, progressive hemiplegia, and cognitive decline. The disease mainly affects children and begins with a prodromal period with mild hemiparesis or infrequent seizures lasting up to several years. The acute stage is marked by frequent seizures arising from one cerebral hemisphere, followed by a residual stage with persistent severe neurological deficits and relapsing epilepsy.

Dane

Klasyfikacja
Choroba

Synonimy
Rasmussen syndrome
Zespół Rasmussena

Kod ORPHA
1929

Kod OMIM
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Kod ICD10
G04.8

Kod ICD11
8A62.Y

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*Źródło

orphanet