

Opis choroby *

Definicja

A rare, non-syndromic cerebral malformation due to abnormal neuronal migration characterized by variable clinical manifestation depending on the location, size and thickness of subcortical bands. Clinical presentation ranges from mild cognitive deficit to developmental delay with severe intellectual disability, seizures and behavioral problems.

Dane

Klasyfikacja	Synonimy
Wada morfologiczna	Subcortical laminar heterotopia Podkorowa heterotopia warstwowa

Kod ORPHA	Kod OMIM	Kod ICD10
99796	607432	Q04.3

Kod ICD11
LD20.1

*Źródło

orphanet