Opis choroby *

Definicja

A rare, inherited disorder characterized by widespread calcifications of basal ganglia and cortex, developmental delay, small stature, retinopathy and microcephaly. The absence of progressive deterioration of the neurological functions is characteristic of the disease.

Dane

Klasyfikacja
Choroba

Kod ORPHA
178506

Kod OMIM
613658

Kod ICD10
G93.8

Kod ICD11
LD20.4

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*Źródło

orphanet