

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

*Źródło

orphanet