

Opis choroby *

Definicja

A rare genetic cerebral small vessel disease characterized by amyloid deposition in the cerebral blood vessels leading to predominantly hemorrhagic strokes, focal neurological deficits, and progressive cognitive decline eventually leading to dementia.

Dane

Klasyfikacja

Choroba

Synonimy

HCHWA

HCHWA

Kod ORPHA

85458

Kod OMIM

605714

Kod ICD10

I68.0*

Kod ICD11

8B22.3

[*Źródło](#)

orphanet