Opis choroby *

Definicja

A rare genetic cerebral small vessel disease characterized by subcortical ischemic events associated with cognitive decline and gait disturbance with an age of onset typically in the sixth or seventh decade of life. Imaging reveals white matter hyperintensities, status cribrosus, lacunar infarcts, and sometimes microbleeds. Extra-neurological manifestations are absent.

Dane

Klasyfikacja
Choroba

Synonimy
HTRA1-related autosomal dominant cerebral angiopathy
Autosomalna dominująca angiopatia mózgu zależna od HTRA-1

Kod ORPHA
482077

Kod OMIM
616779

Kod ICD10
I67.8

Kod ICD11
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*Źródło

orphanet