Opis choroby *

Definicja

A rare genetic cerebral small vessel disease characterized by leukoencephalopathy and cerebral calcification and cysts due to diffuse cerebral microangiopathy resulting in microcystic and macrocystic parenchymal degeneration. The condition can present at any age from early childhood to late adulthood and manifests as a progressive cerebral degeneration. Symptoms are variable, but restricted to the central nervous systems, and include, among others, slowing of cognitive performance, seizures, and movement disorder with a combination of pyramidal, extrapyramidal, and cerebellar features.

Dane

Klasyfikacja
Choroba

Synonimy
LCC
Zespół Labrunego
Labrune syndrome

Kod ORPHA
542310

Kod OMIM
614561

Kod ICD10
I67.8

Kod ICD11
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*Źródło

orphanet