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 Synonimy Choroba LCC

> Zespół Labrunego Labrune syndrome

 Kod ORPHA
 Kod OMIM
 Kod ICD10

 542310
 614561
 I67.8

Kod ICD11

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*Źródło

orphanet