

Opis choroby *

Definicja

A rare genetic disorder of copper metabolism presenting with non-specific hepatic, neurologic, psychiatric or ophthalmologic manifestations due to impaired biliary copper excretion and consecutive excessive copper deposition in the body.

Dane

Klasyfikacja

Choroba

Synonimy

Hepatolenticular degeneration

Zwyrodnienie soczewkowo-wątrobowe

Kod ORPHA

905

Kod OMIM

277900

Kod ICD10

E83.0

Kod ICD11

5C64.00

[*Źródło](#)

orphanet