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

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*Źródło

orphanet