## 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