Opis choroby *

Definicja

A rare sub-group of porphyrias characterized by the occurrence of neuro-visceral attacks with or without cutaneous manifestations. Acute hepatic porphyrias encompass four diseases: acute intermittent porphyria (the most common), variagate porphyria, hereditary coproporphyria, and hereditary deficit of delta-aminolevulinic acid dehydratase (extremely rare).

Dane

Klasyfikacja
Grupa fenomenów

Kod ORPHA
95157

Kod OMIM
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Kod ICD10
E80.2

Kod ICD11
5C58.1Y

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*Źródło

orphanet