

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 coproporphyrinuria, and hereditary deficiency 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

*Źródło

orphanet