Opis choroby *

Definicja

A rare, syndromic, hyperpigmentation of the skin characterized by multiple lentigines and café-au-lait spots associated with hiatal hernia and peptic ulcer, hypertelorism and myopia. There have been no further descriptions in the literature since 1982.

Dane

Klasyfikacja
Choroba

Kod ORPHA
2069

Kod OMIM
137270

Kod ICD10
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Kod ICD11
EC23.1

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*Źródło

orphanet