Opis choroby *

Definicja

A rare constitutional hemolytic anemia characterized by a low 6-phosphogluconate dehydrogenase activity in the erythrocytes, which clinically manifests with a well-compensated chronic nonspherocytic hemolytic anemia and transient hemolytic periods with jaundice.

Dane

Klasyfikacja
Choroba

Kod ORPHA
99135

Kod OMIM
619199

Kod ICD10
D55.1

Kod ICD11
3A10.0Y

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*Źródło

orphanet