Opis choroby *

Definicja

A rare red cell disorder classified principally into two clinical phenotypes: autosomal recessive congenital (or hereditary) methemoglobinemia types I and II (RCM/RHM type 1; RCM/RHM type 2).

Dane

Klasyfikacja
Choroba

Synonimy
Autosomal recessive methemoglobinemia
Methemoglobinemia wrodzona
Autosomalna recesywna methemoglobinemia
Congenital methemoglobinemia

Kod ORPHA
621

Kod OMIM
250800

Kod ICD10
D74.0

Kod ICD11
3A92

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*Źródło

orphanet