Niedobór mieloperoksydazy
Kod Orpha: 2587 Kod OMIM: 254600

Opis choroby *

Definicja
A rare primary immunodeficiency due to a defect in innate immunity characterized by a marked decrease or absence of myeloperoxidase activity in neutrophils and monocytes. Clinically, most patients are asymptomatic. Occasionally, severe infectious complications may occur, particularly recurrent candida infections, being especially severe in the setting of comorbid diabetes mellitus.
Dane

Klasyfikacja
Choroba

Synonimy
MPO deficiency
Niedobór MPO

Kod ORPHA
2587

Kod OMIM
254600

Kod ICD10
E80.3

Kod ICD11
4A00.0Y

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.