

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

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.