Opis choroby *

Definicja

Autosomal recessive severe congenital neutropenia due to CXCR2 deficiency is a rare, genetic, primary immunodeficiency disorder characterized by recurrent bacterial infections (including septic thrombophlebitis and subacute bacterial endocarditis) and neutropenia without lymphopenia or warts, resulting from recessively inherited mutations in <i>CXCR2<i/>.

Dane

Klasyfikacja
Choroba

Kod ORPHA
420699

Kod OMIM
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Kod ICD10
D70

Kod ICD11
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*Źródło

orphanet