

Niedobór NIK

Kod Orpha: 447731 Kod OMIM:

Opis choroby *

Definicja

A rare, genetic, primary combined T and B cell immunodeficiency characterized by recurrent, severe viral and bacterial infections. Immunologic findings include decreased immunoglobulin levels, decreased numbers of B and NK cells, reduced relative CD19+ B cells in peripheral blood, impaired memory responses to viral infections and defective antigen-specific T-cell proliferation.

Dane

Klasyfikacja

Choroba

Synonimy

Primary immunodeficiency with multifaceted aberrant lymphoid immunity
Primary immunodeficiency with multifaceted aberrant lymphoid immunity

Kod ORPHA

447731

Kod OMIM

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Kod ICD10

D81.8

Kod ICD11

4A01.1Y

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.