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
-

Kod ICD10
D81.8

Kod ICD11
4A01.1Y

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.