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

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*Źródło

orphanet