

Opis choroby *

Definicja

A rare autosomal recessive primary immunodeficiency characterized by susceptibility to Epstein-Barr virus (EBV)-associated lymphoproliferative disorders such as malignant B-cell proliferation, Hodgkin lymphoma, B-cell lymphoma, lymphoid granulomatosis, hemophagocytic lymphohistiocytosis, and smooth muscle tumor. Patients present persistent symptoms of infectious mononucleosis including recurrent febrile episodes, lymphadenopathies, and hepatosplenomegaly, accompanied by high EBV viral load in the blood. Additional manifestations are autoimmune diseases like hemolytic anemia or renal disease.

Dane

Klasyfikacja

Choroba

Synonimy

Autosomal recessive lymphoproliferative disease due to ITK deficiency

Autosomal recessive lymphoproliferative disease due to ITK deficiency

ITK deficiency

Kod ORPHA

538963

Kod OMIM

613011

Kod ICD10

D82.3

Kod ICD11

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[*Źródło](#)

orphanet