

Opis choroby *

Definicja

A rare large granular lymphocyte leukemia characterized by persistent (> 6 months) natural killer cell lymphocytosis in the absence of clinical diagnosis of leukemia/lymphoma, autoimmune disease, or chronic viral infections. The clinical course is variable, but generally indolent. Patients often remain asymptomatic, or may present with clinical manifestations including vasculitic skin lesions, neutropenic infections, musculoskeletal symptoms, peripheral neuropathy, or splenomegaly.

Dane

| Klasyfikacja | Synonimy |
|--------------|--|
| Choroba | CLPD-NK |
| | CNKL |
| | CLPD-NK |
| | Przewlekła limfocytoza NK |
| | Przewlekła limfocytoza z komórek NK |
| | Przewlekła choroba limfoproliferacyjna z komórek NK |
| | Choroba proliferacyjna limfocytów ziarnistych z linii NK |
| | CNKL |
| | Chronic NK lymphocytosis |
| | Chronic NK-cell lymphocytosis |
| | Chronic lymphoproliferative disorder of NK-cells |
| | NK-cell lineage granular lymphocyte proliferative disorder |

Kod ORPHA
512017

Kod OMIM
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Kod ICD10
C91.7

Kod ICD11
2A90.2

*[Źródło](#)

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