Opis choroby *

Definicja

An extremely rare and highly aggressive neoplasm, usually manifesting in the third to fourth decade of life, affecting males and females equally, and characterized by the onset of high fever, weight loss, jaundice, skin infiltration, lymphadenopathy, hepatosplenomegaly, and severe anemia. It has a fulminant and rapidly fatal disease course with the progressive appearance of multiorgan failure and disseminated intravascular coagulation.

Dane

Klasyfikacja
Choroba

Synonimy
ANKCL
Agresywny chłoniak z komórek NK
ANKCL
Białaczka limfocytarna z dużych ziarnistych komórek NK
NK-cell LGL leukemia
Aggressive NK-cell lymphoma
NK-cell LGL leukemia
NK-cell large granular lymphocyte leukemia

Kod ORPHA
86873

Kod OMIM
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Kod ICD10
C94.7

Kod ICD11
2A90.3

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*Źródło

orphanet