

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	Synonimy
Choroba	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	Kod OMIM	Kod ICD10
86873	-	C94.7

Kod ICD11
2A90.3

*Źródło

orphanet