Opis choroby *

Definicja

A subtype of acute myeloid leukemia with recurrent genetic abnormalities characterized by leukocytosis, thrombocytosis and nonspecific symptoms related to ineffective hematopoiesis (fatigue, bleeding and bruising, recurrent infections, bone pain), with frequent extramedullary involvement typically presenting as gingival hyperplasia and lymphadenopathy. The disease is characterized by clonal proliferation of myeloid blasts harboring mutations of the NPM1 gene in the bone marrow, blood and other tissues. It is associated with multilineage dysplasia, involving the myeloid, monocytic, erythroid, and megakaryocytic cell lineages.

Dane

Klasyfikacja
Choroba

Synonimy
AML with NPM1 somatic mutations
AML z somatycznymi mutacjami NPM1

Kod ORPHA
402026

Kod OMIM
-

Kod ICD10
C92.0

Kod ICD11
-

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*Źródło

orphanet