Opis choroby *

Definicja

A distinct form of Acute myeloid leukemia (AML) in which this chromosomal anomaly is found de novo or in therapy-related AML cases, and is characterized by frequent extramedullary involvement (mainly hepatomegaly, splenomegaly, lymphadenopathies, cutaneous infiltration, but also gum, bone, central nervous system, testicles involvement), severe coagulation disorder (disseminated intravascular coagulopathy or primary fibrinolysis) and poor prognosis. Morphologically, a blast population with a myelomonocytic stage of differentiation is observed.

Dane

Klasyfikacja
Choroba

Synonimy
AML with t(8;16)(p11;p13) translocation
AML z translokacją t(8;16)(p11;p13)

Kod ORPHA
370026

Kod OMIM
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Kod ICD10
C92.0

Kod ICD11
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*Źródło

orphanet