

## 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