

## Opis choroby \*

### Definicja

A rare acute myeloid leukemia (AML) characterized by the presence of acute leukemia with at least 20% peripheral blood or bone marrow blasts with morphological features of myelodysplasia, or occurrence in patients with a prior history of a myelodysplastic syndrome (MDS) or myelodysplastic/myeloproliferative neoplasm, with MDS-related cytogenetic abnormalities, in the absence of specific genetic abnormalities characteristic of AML with recurrent genetic abnormalities. Prior cytotoxic or radiation therapy for an unrelated disease must be excluded. The condition occurs mainly in elderly patients and is rare in children. Patients often present with severe pancytopenia. Prognosis is generally poor.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

AML with multilineage dysplasia

AML z dysplazją wieloliniową

AML with myelodysplasia-related features

Acute myeloid leukemia with multilineage dysplasia

#### Kod ORPHA

86845

#### Kod OMIM

601626

#### Kod ICD10

C92.8

#### Kod ICD11

2A60.1

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#### [\\*Źródło](#)

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