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

orphanet