Opis choroby *

Definicja

A rare acute leukemia of ambiguous lineage characterized by clonal proliferation of primitive hematopoietic cells, primarily in the bone marrow and blood, lacking lineage-specific markers and detectable genotypic alterations. The patients present with leukocytosis, anemia, variable platelet count and a variety of nonspecific symptoms related to ineffective hematopoesis (fatigue, bleeding and bruising, recurrent infections, bone pain) and/or extramedullary site involvement (lymphadenopathy, splenomegaly, hepatomegaly).

Dane

Klasyfikacja
Choroba

Synonimy
Acute myeloid leukemia, minimal differentiation, FAB M0
Acute myeloid leukemia, minimal differentiation, FAB M0

Kod ORPHA
98835

Kod OMIM
601626

Kod ICD10
C95.0

Kod ICD11
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*Źródło

orphanet