Opis choroby *

Definicja

A subtype of acute myeloid leukemia with recurrent genetic abnormalities, characterized by clonal proliferation of myeloid blasts harboring somatic mutations of the CEBPA gene in the bone marrow, blood and, rarely, other tissues. It can present with anemia, thrombocytopenia, and other nonspecific symptoms related to ineffective hematopoesis (fatigue, bleeding and bruising, recurrent infections, bone pain) and/or extramedullary site involvement (gingivitis, splenomegaly).

Dane

Klasyfikacja
Choroba

Synonimy
AML with CEBPA somatic mutations
AML z somtycznymi mutacjami CEBPA

Kod ORPHA
319480

Kod OMIM
601626

Kod ICD10
C92.0

Kod ICD11
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*Źródło

orphanet