

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 hematopoiesis (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