Acute myeloid leukemia with myelodysplasia-related changes (AML-MRC) is an AML subtype with a poor prognosis

The WHO 2016 diagnostic criteria for AML-MRC^{1,2}



Defined as patients with AML who have ≥20% **blasts** in the peripheral blood or bone marrow and any of the following^{1,2}:



Previously documented MDS or MDS/MPN



MDS-related cytogenetic abnormalities

Multilineage dysplasia in \geq 50% of \geq 2 cell lineages in the absence of *NPM1* or biallelic *CEBPA* mutations ပိုလ

Excluding factors^{1,2}:

Any of the cytogenetic abnormalities qualifying for diagnosis of AML with recurrent genetic abnormalities, such as inv(3), t(6;9), or NPM1 mutation

Prior cytotoxic therapy for unrelated disease

AML-MRC is not included as a category in the ICC and WHO 2022 classifications^{3,4}:

ICC 2022: prior MDS or MDS/MPN is used as a diagnostic qualifier, AML with myelodysplasia-related cytogenetic abnormality and AML with myelodysplasia-related gene mutation are separate categories, and multilineage dysplasia is not included

WHO 2022: a type of AML with defining genetic abnormalities "AML-MR" replaces AML-MRC and includes AML transformation of MDS or MDS/MPN and AML with MDS-related cytogenetics or gene mutations; multilineage dysplasia is not included

MDS-related cytogenetic abnormalities sufficient to diagnose AML-MRC (WHO 2016 classification)^{a,1,2}



^aSufficient to diagnose when ≥20% peripheral blood or bone marrow blasts are present and prior therapy has been excluded.



Abbreviations: AML-MRC, acute myeloid leukemia with myelodysplasia-related changes; AML, acute myeloid leukemia; WHO, World Health Organization; MDS, myelodysplastic syndrome; MPN, myeloproliferative neoplasm; NPM1, nucleophosmin 1; CEBPA, CCAAT enhancer binding protein alpha; FISH, fluorescence in situ hybridizatior

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AML-MRC primarily occurs in older patients^{5,6}

AML-MRC represents up to 48% of all adult AML cases^{5,6,14}





Median age at diagnosis5: 68 vears

AML-MRC is associated with poor clinical outcomes⁷⁻¹²

AML-MRC treated with conventional chemotherapy:

Complete remission rate: **24%**¹³–61%⁷

Median overall survival7:



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