ALLOGENEIC HAEMATOPOIETIC CELL TRANSPLANTATION IN PATIENTS AGED <60 YEARS WITH ACUTE MYELOID LEUKEMIA

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BACKGROUND AND IMPORTANCE

Allogeneic hematopoietic cell transplantation is a potentially curative therapeutic modality for acute myeloid leukemia, but it still carries high morbidity and mortality; there are limited data regarding outcomes, so it is important to research its results, and the factors that influence them.



To assess the survival of allo-HCT in AML patients age <60 years, describe its characteristics, and identify factors that are related to the best outcomes.

MATERIAL AND METHODS



Overall survival and progression-free survival were analyzed using Kaplan-Meier and Long-Rank test





51% intermediate-risk, 43% high-risk

70% in first completer remission

92% patients KPS score over 90%

54% HCT-CI 0-2, 8<mark>1% EBMTscore ≤4</mark>

65% related donor

37 patients (43% HLA-identical and **22% haploidentical**)

Mean age 45 years 35% unrelated donor

65% women

(22% HLA-identical, 11% HLA 9/10, and 3% HLA 8/10)

70% allogeneic peripheral blood stem cell transplantation

65% reduced-intensity conditioning

16% retransplantation

Most donors were men >30 years



38% received posttransplantation treatment with

cyclophosphamide, tacrolimus, and mycophenolate mofetil

19% CMV-mismatch (patient pos/ donor neg)

57% ABO-compatible

54% development chronic GVHD and 40% acute GVHD

43% didn't require related hospitalization



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Median PFS and OS were not reached. The median follow-up for PFS was 33 months [1-69] and 34 months [1-69] for OS. PFS was significantly higher in patients in 1st CR, EBMTscore ≤4, and lower-risk.

CONCLUSION

Patients undergoing allo-HCT show encouraging survival, although more extended follow-up is required to define more accurately their prognosis.

