

## Pediatric disease risk index for acute leukemia

To obtain credit, you should first read the journal article. After reading the article, you should be able to answer the following, related, multiple-choice questions. To complete the questions (with a minimum 75% passing score) and earn continuing medical education (CME) credit, please go to <http://www.medscape.org/journal/blood>. Credit cannot be obtained for tests completed on paper, although you may use the worksheet below to keep a record of your answers. You must be a registered user on <http://www.medscape.org>. If you are not registered on <http://www.medscape.org>, please click on the "Register" link on the right hand side of the website. Only one answer is correct for each question. Once you successfully answer all post-test questions you will be able to view and/or print your certificate. For questions regarding this activity, contact the accredited provider, [CME@medscape.net](mailto:CME@medscape.net). For technical assistance, contact [CME@medscape.net](mailto:CME@medscape.net). American Medical Association Physician's Recognition Award (AMA PRA) credits are accepted in the US as evidence of participation in CME activities. For further information on this award, please go to <https://www.ama-assn.org>. The AMA has determined that physicians not licensed in the US who participate in this CME activity are eligible for *AMA PRA Category 1 Credits™*. Through agreements that the AMA has made with agencies in some countries, AMA PRA credit may be acceptable as evidence of participation in CME activities. If you are not licensed in the US, please complete the questions online, print the AMA PRA CME credit certificate, and present it to your national medical association for review.

**Qayed M, Ahn KW, Kitko CL, Johnson MH, Shah NN, Dvorak C, Mellgren K, Friend BD, Verneris MR, Leung W, Toporski J, Levine J, Chewning J, Wayne A, Kapoor U, Triplett B, Schultz KR, Yanik GA, Eapen M. A validated pediatric disease risk index for allogeneic hematopoietic cell transplantation. *Blood*. 2021;137(7):983-993.**

1. **Your patient is a 7-year-old boy with acute myeloid leukemia (AML). According to the study by Qayed and colleagues, which of the following statements about development of the pediatric disease risk index (DRI) for stratifying children and adolescents with AML and acute lymphocytic leukemia (ALL) undergoing hematopoietic cell transplantation (HCT) into risk groups according to patient and disease characteristics is correct?**
  - For ALL, B- and T-cell lineage and cytogenetic risk were significantly associated with leukemia-free survival (LFS)
  - Hematopoietic comorbidity score and performance score were significantly associated with LFS for AML and ALL
  - Two independent predictors were associated with lower LFS for AML
  - For AML, 4 risk groups were identified on the basis of age, cytogenetic risk, and disease status, including minimal residual disease status at transplantation
  
2. **According to the study by Qayed and colleagues, which of the following statements about validation of the pediatric DRI for stratifying children and adolescents with AML and ALL undergoing HCT into risk groups on the basis of patient and disease characteristics is correct?**
  - For AML, 5-year LFS for low (0 points), intermediate (2, 3, 5), high (7, 8), and very high (> 8) risk groups was 78%, 53%, 40%, and 25%, respectively ( $P < .0001$ )
  - For ALL, the 3 risk groups did not differ significantly in 5-year LFS
  - The DRI did not predict outcome among patients with AML aged 12 to 18 years
  - For AML, compared with patients with low risk, overall survival was 50% lower (hazard ratio = 1.5) for individuals with high risk or very high risk
  
3. **According to the study by Qayed and colleagues, which of the following statements about clinical implications of the pediatric DRI for stratifying children and adolescents with AML and ALL undergoing HCT into risk groups on the basis of patient and disease characteristics is correct?**
  - The pediatric DRI is not feasible or reliable for use in clinical settings
  - Donor type affects the usefulness of the pediatric DRI
  - The DRI can stratify heterogeneous populations of children and adolescents in HCT trials on the basis of their risk for relapse or mortality
  - The pediatric DRI is likely to be useful only within the United States and is not more effective in children than the adult DRI