## POST-TEST

Meet The Professor: Optimizing the Management of Acute Myeloid Leukemia and Myelodysplastic Syndromes — Part 1 of a 3-Part Series

## THE CORRECT ANSWER IS INDICATED WITH YELLOW HIGHLIGHTING.

- 1. An updated overall survival (OS) analysis of the Phase III ASCERTAIN trial evaluating oral decitabine/cedazuridine for intermediate- or high-risk myelodysplastic syndromes (MDS) demonstrated which of the following outcomes?
  - a. A median OS of 5.1 months
  - b. A median OS of 15.3 months
  - c. A median OS of 31.7 months
- 2. The Phase III COMMANDS trial evaluating luspatercept versus epoetin alfa for lower-risk transfusion-dependent MDS demonstrated which of the following outcomes?
  - a. Epoetin alfa was superior to luspatercept in increasing the likelihood of achieving transfusion independence and increasing hemoglobin levels
  - b. Luspatercept was superior to epoetin alfa in increasing the likelihood of achieving transfusion independence and increasing hemoglobin levels
  - No difference was observed between luspatercept and epoetin alfa in subsequent transfusion independence or increased hemoglobin levels

- 3. Data from the QuANTUM-First trial led to the recent FDA approval of quizartinib in combination with chemotherapy for patients with acute myeloid leukemia (AML) in which setting?
  - a. Newly diagnosed AML with an IDH1 mutation
  - b. Relapsed AML with an IDH1 mutation
  - c. Newly diagnosed AML with a FLT3-ITD mutation
  - d. Relapsed AML with a FLT3-ITD mutation
- 4. The Phase III placebo-controlled QuANTUM-First trial of quizartinib with chemotherapy for AML demonstrated which of the following outcomes?
  - a. A nonsignificant increase in OS with quizartinib/chemotherapy
  - b. A statistically significant doubling of OS with quizartinib/chemotherapy
  - No difference in OS between quizartinib/chemotherapy and placebo
- 5. Which of the following drug types best reflects the mechanism of action of olutasidenib?
  - a. Hypomethylating agent
  - b. FLT3 inhibitor
  - c. IDH1 inhibitor
    - d. Bcl-2 inhibitor