

Acute Myeloid Leukemia and the General Medical Oncologist: New Agents and Treatment Strategies, Particularly for Older Patients — A Virtual Meet The Professor Series

THE CORRECT ANSWER IS INDICATED WITH YELLOW HIGHLIGHTING.

1. The ADMIRAL trial of gilteritinib compared to chemotherapy for patients with relapsed/refractory AML with a FLT3 mutation demonstrated which of the following clinical outcomes for the gilteritinib study arm?
 - a. Statistically significant improvement in overall survival when compared to chemotherapy
 - b. An increase in the proportion of patients achieving complete remission (CR) when compared to chemotherapy
 - c. No statistically significant improvement in overall survival when compared to chemotherapy
 - d. No increase in the proportion of patients achieving CR when compared to chemotherapy
 - e. Both a and b
 - f. Both c and d
2. The Phase I AG120-C-001 study evaluated first-line therapy for patients with AML with which IDH inhibitor that is FDA approved in the newly diagnosed setting?
 - a. Enasidenib
 - b. Ivosidenib
3. Differentiation syndrome associated with IDH1/2 inhibitors, such as ivosidenib and enasidenib, is characterized by which of the following features?
 - a. Nonspecific symptoms such as fever, weight gain, rash and rising leukocyte count
 - b. A median time to onset of approximately 30 days after the initiation of treatment
 - c. A high frequency of occurrence (>50%) at Grade 3 or higher severity
 - d. Both a and b
 - e. Both a and c
4. The Phase III VIALE-A trial evaluating azacitidine in combination with either venetoclax or placebo for patients with treatment-naïve AML not eligible for intensive therapy demonstrated which clinical outcome on the azacitidine/venetoclax arm?
 - a. Statistically significant improvement in overall survival
 - b. An increase in the proportion of patients achieving CR
 - c. Both a and b
 - d. Neither a nor b
5. Which of the following drug types best characterizes the mechanism of action of the novel agent pevonedistat?
 - a. Bcl-2 inhibitor
 - b. FLT3 inhibitor
 - c. IDH1/2 inhibitor
 - d. NEDD8-activating enzyme inhibitor