POST-TEST

What Clinicians Want to Know: Addressing Current Questions and Controversies in the Management of Myelofibrosis

THE CORRECT ANSWER IS INDICATED WITH YELLOW HIGHLIGHTING.

- 1. Which of the following drug types best reflects the mechanism of action of imetelstat?
 - a. ALK2 inhibitor
 - b. JAK inhibitor
 - c. Telomerase inhibitor
 - d. XPO1 inhibitor
- 2. In the Phase II IMbark trial, imetelstat demonstrated a clinically meaningful benefit in which of the following populations?
 - a. Patients whose disease is treatment naïve
 - b. Patients whose disease is relapsed or refractory to JAK inhibition
 - c. Patients whose disease is heavily pretreated (after 4 or more prior lines of therapy)
 - d. Patients with platelet counts below 50,000/µL

3. Which of the following pathways and processes is affected by selinexor?

- a. JAK-STAT
- b. p53-driven cell death
- с. NF-кB
- d. Cell cycle arrest
- e. All of the above

- 4. Which of the following agents is under investigation in combination with ruxolitinib for JAK inhibitor-naïve myelofibrosis (MF) in a Phase III study?
 - a. Vorinostat
 - b. Selinexor
 - c. Galunisertib
 - d. Alisertib
- Which of the following JAK inhibitors has been associated with a survival benefit for patients with JAK inhibitor-naïve MF?
 - a. JAK inhibitors are not associated with a survival benefit in this setting
 - b. Ruxolitinib
 - c. Fedratinib
 - d. Pacritinib
 - e. Momelotinib