POST-TEST

What Clinicians Want to Know About Toxicity Considerations Associated with BTK Inhibitors

THE CORRECT ANSWER IS INDICATED WITH YELLOW HIGHLIGHTING.

- 1. Which of the following drugs is classified as a reversible Bruton tyrosine kinase (BTK) inhibitor?
 - a. Ibrutinib
 - b. Acalabrutinib
 - c. Zanubrutinib
 - d. Pirtobrutinib
- 2. Which of the following nonhematologic adverse events is NOT commonly observed with the available BTK inhibitors?
 - a. Arthralgia
 - b. Diarrhea
 - c. Infection
 - d. Alopecia
- 3. Which of the following BTK inhibitors is now available in a tablet formulation that allows its coadministration with proton pump inhibitors or its administration to patients unable to swallow capsules?

a. Acalabrutinib

- b. Ibrutinib
- c. Zanubrutinib
- d. Pirtobrutinib

- 4. Which of the following statements best reflects cardiotoxicity observations from trials evaluating ibrutinib or acalabrutinib for patients with chronic lymphocytic leukemia/small lymphocytic lymphoma (CLL/SLL)?
 - a. The incidence of atrial fibrillation observed in trials of acalabrutinib was comparable to that observed in trials of ibrutinib
 - b. The incidence of atrial fibrillation observed in trials of acalabrutinib was higher than that observed in trials of ibrutinib
 - c. The incidence of atrial fibrillation observed in trials of acalabrutinib was lower than that observed in trials of ibrutinib
- 5. Which of the following statements best characterizes the incidence of treatment-related headache observed among patients with CLL/SLL receiving pirtobrutinib in the BRUIN trial?
 - No patients were observed to experience headache as a treatmentrelated toxicity with pirtobrutinib
 - b. Incidence of any-grade and Grade ≥3 treatment-related headache was low
 - c. Incidence of any-grade and Grade ≥3 treatment-related headache was notably high