## POST-TEST

Preventing and Managing Toxicities Associated with Antibody-Drug Conjugates in the Management of Metastatic Breast Cancer

## THE CORRECT ANSWER IS INDICATED WITH YELLOW HIGHLIGHTING.

- 1. What was the primary objective of the Phase II PRIMED study of sacituzumab govitecan for advanced breast cancer?
  - a. To evaluate the efficacy and safety of sacituzumab govitecan for previously untreated PD-L1-negative metastatic triple-negative breast cancer (mTNBC)
  - b. To evaluate the effectiveness of prophylactic granulocyte colonystimulating factor (G-CSF) and loperamide in improving the tolerability of sacituzumab govitecan
  - c. To evaluate the effectiveness of various dose levels and schedules of sacituzumab govitecan in improving tolerability
- 2. Which of the following statements best describes the rates of discontinuation due to adverse events observed in the Phase III ASCENT-04/KEYNOTE-D19 trial for patients who received first-line sacituzumab govitecan with pembrolizumab compared to those who received pembrolizumab with chemotherapy for PD-L1-positive mTNBC?
  - a. The rate was higher with sacituzumab govitecan and pembrolizumab
  - b. The rates were similar between the treatment arms
  - c. The rate was higher with pembrolizumab and chemotherapy
- 3. Which of the following statements best describes the incidence of neutropenia observed with sacituzumab govitecan and pembrolizumab compared to pembrolizumab and chemotherapy in the Phase III ASCENT-04/KEYNOTE-D19 trial?
  - a. The incidence was higher with sacituzumab govitecan and pembrolizumab
  - b. The incidence was comparable between the treatment arms
  - c. The incidence was higher with pembrolizumab and chemotherapy

- 4. Which of the following prophylactic measures is recommended to manage oral mucositis/stomatitis associated with datopotamab deruxtecan?
  - a. Saline mouthwashes
  - b. Steroid mouthwashes
  - c. Benzocaine lozenges
  - d. No prophylaxis is needed for oral mucositis/stomatitis
- 5. A patient with advanced breast cancer receives sacituzumab govitecan but experiences their first occurrence of Grade 3 diarrhea. Aside from administering loperamide, what is the recommended course of action?
  - a. Continue sacituzumab govitecan uninterrupted at the same dose
  - b. Hold sacituzumab govitecan until diarrhea resolves to Grade ≤2, then resume at the same dose
  - c. Hold sacituzumab govitecan until diarrhea resolves to Grade ≤2, then resume at a dose reduced by 25%
  - d. Hold sacituzumab govitecan until diarrhea resolves to Grade ≤2, then resume at a dose reduced by 50%
  - e. Discontinue sacituzumab govitecan