## Lymphoma and Chronic Lymphocytic Leukemia Update — Volume 1, Issue 2

## THE CORRECT ANSWER IS INDICATED WITH YELLOW HIGHLIGHTING.

- 1. Which of the following categories reflects the mechanism of action of obinutuzumab?
  - a. Anti-CD20 monoclonal antibody
  - b. Immunomodulatory drug
  - c. Anti-PD-1/PD-L1 antibody
  - d. Proteasome inhibitor
- 2. Which of the following observations was made in the Phase III GALLIUM study evaluating obinutuzumab- versus rituximab-based induction and maintenance therapy for previously untreated FL?
  - a. No difference in PFS
  - b. PFS favored rituximab
  - c. PFS favored obinutuzumab
- Hospitalization for the purpose of monitoring for TLS is required for all patients starting therapy with venetoclax.
  - a. True
  - b. False
- 4. Which of the following categories reflects the mechanism of action of copanlisib?
  - a. Anti-PD-1/PD-L1 antibody
  - b. Bruton tyrosine kinase inhibitor
  - c. CAR-T therapy
  - d. PI3K inhibitor
- 5. Results of the Phase III AETHERA trial evaluating brentuximab vedotin versus placebo as consolidation therapy after ASCT for patients with HL at risk of relapse or disease progression demonstrated a statistically significant improvement in \_\_\_\_\_ with brentuximab vedotin.
  - a. Overall survival
  - b. PFS
  - c. Both a and b
  - d. Neither a nor b

- The Phase III LyMa trial \_\_\_\_\_ a statistically significant overall survival advantage with rituximab maintenance therapy after ASCT for younger patients with MCL.
  - a. Demonstrated
  - b. Did not demonstrate
- 7. Which side effect is of the greatest concern for patients with acute lymphomas receiving CAR-T therapy?
  - a. Cytokine release syndrome
  - b. Renal failure
  - c. TLS
- 8. The majority of patients with del(17p) CLL
  - a. Present up front with the 17p deletion
  - b. Acquire the 17p deletion over the course of their disease
- 9. Venetoclax is dosed and administered in which of the following manners?
  - a. 20 mg once daily
  - b. 400 mg once daily
  - Initiated at 20 mg and gradually escalated to the target dose of 400 mg once daily
- is an orally bioavailable inhibitor of the delta isoform of PI3 kinase that is approved by the FDA for the treatment of relapsed CLL.
  - a. Copanlisib
  - b. Ibrutinib
  - c. Idelalisib
  - d. TGR-1202