

THE CORRECT ANSWER IS INDICATED WITH YELLOW HIGHLIGHTING.

- A Phase II study by Stilgenbauer and colleagues evaluating venetoclax monotherapy for patients with relapsed/refractory del(17p) CLL demonstrated an overall response rate of approximately 80% but a high incidence of clinical tumor lysis syndrome.

  - True
  - False
- The Phase I/II ACE-CL-001 trial evaluating acalabrutinib for relapsed CLL demonstrated \_\_\_\_\_.

  - A high response rate
  - A high incidence of bleeding
  - A favorable safety profile
  - Both a and b
  - Both a and c
- The ongoing Phase III GALLIUM trial is evaluating \_\_\_\_\_ with chemotherapy versus rituximab/chemotherapy followed by maintenance therapy with \_\_\_\_\_ or rituximab in patients with previously untreated FL.

  - Bortezomib
  - Ibritumomab tiuxetan
  - Obinutuzumab
- On the Phase II SORAML trial, the sequential addition of sorafenib to standard chemotherapy for younger patients with newly diagnosed AML resulted in a statistically significant improvement in \_\_\_\_\_ versus standard chemotherapy and placebo.

  - Event-free survival
  - Overall survival
  - Both a and b
  - Neither a nor b
- The Phase III RATIFY (CALGB-10603) trial for patients with newly diagnosed FLT3-mutated AML \_\_\_\_\_ a statistically significant improvement in median overall survival with midostaurin in combination with standard induction and consolidation chemotherapy compared to standard induction and consolidation chemotherapy alone.

  - Demonstrated
  - Did not demonstrate
- The Phase III randomized INO-VATE study comparing inotuzumab ozogamicin to standard chemotherapy for relapsed/refractory ALL demonstrated a 2-year survival rate of \_\_\_\_\_ with inotuzumab and 10% with chemotherapy.

  - 5%
  - 23%
  - 50%
- The SWOG-S0777 trial evaluating RVD versus Rd for patients with previously untreated MM without an intent for immediate ASCT demonstrated \_\_\_\_\_ with RVD.

  - A significant improvement in PFS
  - No improvement in PFS
- The Phase III randomized CASTOR study evaluating daratumumab/bortezomib/dexamethasone versus bortezomib/dexamethasone did not demonstrate a significant improvement in PFS with the addition of daratumumab for patients with relapsed or refractory MM.

  - True
  - False
- The Phase II CheckMate 205 study evaluating the efficacy of nivolumab for relapsed/refractory classical HL demonstrated a 6-month overall survival rate of approximately \_\_\_\_\_.

  - 50%
  - 75%
  - 100%
- Which of the following is the FDA-approved indication for nivolumab in classical HL?

  - Previously untreated classical HL
  - Classical HL after failure of at least 2 prior multiagent chemotherapy regimens in patients who are not candidates for autologous hematopoietic stem cell transplantation
  - Classical Hodgkin lymphoma that has relapsed or progressed after autologous hematopoietic stem cell transplant and post-transplant brentuximab vedotin
  - Nivolumab is not FDA approved for the treatment of classical HL