

POST-TEST

Investigator Perspectives on the Current and Future Management of Newly Diagnosed Ovarian Cancer

THE CORRECT ANSWER IS INDICATED WITH YELLOW HIGHLIGHTING.

- The ongoing Phase III PAOLA-1 trial is investigating the PARP inhibitor _____ in combination with bevacizumab as maintenance therapy for patients with advanced high-grade serous ovarian cancer after first-line platinum-based chemotherapy and bevacizumab.
 - Niraparib
 - Olaparib
 - Rucaparib
 - Veliparib
- In Phase III trials of rucaparib (ARIEL3) or niraparib (ENGOT-OV16/NOVA) as maintenance therapy after a complete or partial response to platinum-based doublet chemotherapy for recurrent high-grade serous ovarian cancer, no statistically significant improvement in progression-free survival was reported with either PARP inhibitor for patients with HRD-positive disease.
 - True
 - False
- The ongoing Phase III PRIMA trial is investigating maintenance therapy with _____ for patients with Stage III or IV ovarian cancer after a response to front-line platinum-based chemotherapy.
 - Veliparib
 - Olaparib
 - Niraparib
- In the Phase III BOOST trial evaluating the optimal duration of first-line bevacizumab in combination with carboplatin and paclitaxel, patients with primary ovarian cancer will receive bevacizumab for 15 months or _____.
 - 6 months
 - 24 months
 - 30 months
- _____ is a side effect associated with bevacizumab in the front-line treatment of advanced ovarian cancer.
 - Gastrointestinal perforation
 - Hypertension
 - Proteinuria
 - All of the above
 - Both b and c
- The results of the 3-arm Phase III GOG-0218 trial of chemotherapy with or without bevacizumab for patients with newly diagnosed advanced ovarian cancer who had undergone debulking surgery _____ demonstrate a statistically significant improvement in progression-free survival for those who received bevacizumab initiation followed by maintenance placebo in comparison to the control group, who received chemotherapy followed by maintenance placebo.
 - Did
 - Did not
- The results of the Phase III ICON7 trial of carboplatin/paclitaxel with or without bevacizumab for newly diagnosed epithelial ovarian cancer demonstrated a statistically significant improvement in _____ with the addition of bevacizumab in the overall patient population.
 - Overall survival
 - Progression-free survival
 - Both a and b
 - Neither a nor b
- The FDA recently approved bevacizumab in combination with carboplatin and paclitaxel followed by single-agent bevacizumab for patients with Stage III or IV epithelial ovarian cancer after initial surgical resection.
 - True
 - False

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9. _____ is an adverse event associated with PARP inhibitor therapy in the management of ovarian cancer.

- a. Insomnia
- b. Diarrhea
- c. Both a and b
- d. Neither a nor b

10. The results of the Phase III GOG-0252 trial evaluating intravenous versus intraperitoneal chemotherapy, each in combination with bevacizumab, for newly diagnosed ovarian cancer demonstrated a statistically significant improvement in progression-free survival with intraperitoneal chemotherapy.

- a. True
- b. False