

THE CORRECT ANSWER IS INDICATED WITH YELLOW HIGHLIGHTING.

- 1. The Phase III EMBARK trial evaluated enzalutamide or placebo with leuprolide acetate and enzalutamide monotherapy in which clinical setting?**
 - a. Patients with metastatic castration-resistant prostate cancer (mCRPC)
 - b. Patients with metastatic hormone-sensitive prostate cancer
 - c. Patients with high-risk biochemically recurrent prostate cancer**
 - d. Patients with low-risk biochemically recurrent prostate cancer
- 2. Androgen deprivation therapy intensification with which of the following androgen receptor pathway inhibitors prolonged PSA progression-free survival among patients with high-risk biochemically relapsed prostate cancer in the Phase III PRESTO trial?**
 - a. Abiraterone
 - b. Apalutamide**
 - c. Enzalutamide
 - d. Darolutamide
- 3. The FDA label for the combination of olaparib with abiraterone for prostate cancer specifies its use for which patient population?**
 - a. Patients with mCRPC and a mutation in a homologous recombination repair gene
 - b. Patients with mCRPC and a BRCA mutation**
 - c. All patients with mCRPC
- 4. Which of the following strategies has been recommended to help manage xerostomia associated with Lutetium Lu 177 vipivotide tetraxetan?**
 - a. Icing salivary glands during treatment
 - b. Use of salivary supplements
 - c. Brushing teeth more frequently
 - d. Icing salivary glands during treatment and use of salivary supplements only
 - e. All of the above**
- 5. The Phase III CONTACT-02 trial evaluated which of the agents below in combination with cabozantinib for patients with mCRPC?**
 - a. Olaparib
 - b. Lutetium Lu 177 vipivotide tetraxetan
 - c. Atezolizumab**
 - d. Abemaciclib