

THE CORRECT ANSWER IS INDICATED WITH YELLOW HIGHLIGHTING.

- 1. Which of the following monoclonal antibodies recently received FDA approval in combination with lenvatinib for patients with advanced endometrial carcinoma that is not microsatellite instability-high or mismatch repair deficient (dMMR) who have experienced disease progression after prior systemic therapy in any setting and are not candidates for curative surgery or radiation therapy?**
 - a. Pembrolizumab**
 - b. Dostarlimab
 - c. Durvalumab
 - d. Nivolumab
- 2. In the GARNET trial evaluating the efficacy and safety of dostarlimab for advanced solid tumors, dostarlimab elicited durable antitumor activity in which subgroup of patients with recurrent or advanced endometrial cancer?**
 - a. Patients with dMMR disease only
 - b. Patients with mismatch repair-proficient (MMRp) disease only
 - c. Patients with either dMMR or MMRp disease**
 - d. Patients with neither dMMR nor MMRp disease
- 3. Which of the following outcomes was demonstrated with lenvatinib/pembrolizumab as second-line therapy for advanced endometrial cancers?**
 - a. Improvement in only progression-free survival (PFS) for only those patients with MMRp disease
 - b. Improvement in only PFS for all patients
 - c. Improvement in only overall survival (OS) for only those patients with MMRp disease
 - d. Improvement in only OS for all patients
 - e. Improvement in OS and PFS for only those patients with MMRp disease
 - f. Improvement in OS and PFS for all patients**
- 4. Which of the following conditions was the most common immune-mediated adverse event in patients receiving pembrolizumab/lenvatinib in both the KEYNOTE-146 and KEYNOTE-775 trials?**
 - a. Colitis
 - b. Infusion reactions
 - c. Severe skin reactions
 - d. Hypothyroidism**