POST-TEST

Beyond the Guidelines: Clinical Investigators Provide Perspectives on Biomarker-Guided Decision-Making for Patients with Non-Small Cell Lung Cancer

THE CORRECT ANSWER IS INDICATED WITH YELLOW HIGHLIGHTING.

- 1. The anti-PD-1 antibody pembrolizumab is FDA approved for which of the following populations of patients with non-small cell lung cancer (NSCLC)?
 - a. As first-line therapy for patients with metastatic NSCLC whose tumors express PD-L1 with a tumor proportion score (TPS) of 50% or higher
 - As first-line therapy in combination with pemetrexed and carboplatin for patients with metastatic nonsquamous NSCLC irrespective of PD-L1 expression
 - c. As second-line therapy for patients with metastatic NSCLC whose tumors express PD-L1 with a TPS of 1% or higher and who experience disease progression after platinumbased chemotherapy

d. All of the above

- e. All except c
- f. Both b and c
- g. None of the above
- 2. Results from the CheckMate 026 Phase III trial evaluating nivolumab versus investigator's choice of chemotherapy as first-line therapy for Stage IV or recurrent PD-L1-positive NSCLC demonstrated a statistically significant improvement in ______ with nivolumab for patients with a TPS of 5% or higher.
 - a. Overall survival
 - b. Objective response rate
 - c. Both a and b
 - d. Neither a nor b

- 3. ______ is an anti-PD-L1 antibody that is FDA approved for patients with metastatic NSCLC and disease progression during or after platinumbased chemotherapy and for those with disease progression on an appropriate FDA-approved targeted therapy if their tumor harbors a mutation in the EGFR or ALK gene.
 - a. Durvalumab
 - b. Atezolizumab
 - c. Nivolumab
 - d. Avelumab
- 4. Which of the following agents has demonstrated activity in patients with advanced ROS1-rearranged NSCLC?
 - a. Gefitinib
 - b. Cabozantinib
 - c. Crizotinib
 - d. Alectinib
 - e. All of the above
 - f. Both b and c
 - g. Both c and d
- 5. The Phase III MYSTIC trial evaluating the anti-PD-L1 antibody durvalumab with or without the MEK inhibitor tremelimumab versus standard platinumbased chemotherapy for patients with previously untreated metastatic NSCLC demonstrated a statistically significant improvement in progression-free survival (PFS) with durvalumab/tremelimumab compared to standard chemotherapy.

a. True b. False

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- 6. The results from the Phase III AURA3 trial evaluating second-line osimertinib versus platinum-based doublet chemotherapy for patients with locally advanced or metastatic NSCLC demonstrated a statistically significant improvement in ______ with osimertinib.
 - a. PFS in the intention-to-treat population
 - b. PFS for patients with CNS metastases

c. Both a and b

- d. Neither a nor b
- 7. The Phase III ASCEND-4 trial evaluating first-line ceritinib versus platinum-based chemotherapy for patients with advanced ALK-rearranged NSCLC demonstrated a statistically significant improvement in PFS with ceritinib.
 - a. True
 - b. False
- 8. _____ is an ALK inhibitor that has demonstrated antitumor activity in patients with ALK-rearranged NSCLC and CNS metastases.

a. Alectinib

- b. Crizotinib
- c. Both a and b

- 9. Recently the FDA approved the BRAF inhibitor ______ in combination with the MEK inhibitor trametinib for the treatment of metastatic NSCLC in patients harboring BRAF V600E mutations as detected by an FDA-approved test.
 - a. Vemurafenib
 - b. Dabrafenib
 - c. Cabozantinib
 - d. All of the above
- 10. If a tissue biopsy shows no evidence of a T790M mutation, a patient's disease is conclusively T790M mutation-negative and no further testing is required.
 - a. True
 - b. False