Webinar 1 – Cancer Immunology

Learning Objectives:

- A. Describe the role of the immune system in cancer prevention
- B. Understand how immunoediting leads to tumor escape mechanisms.
- C. Describe the three immune phenotypes that lead to cancer growth.

Questions:

- 1. At each step of the cancer cycle, mutations occur that either increase cell death or limit cell proliferation
 - a. True
 - b. False
- 2. Immune deficient mice were shown to be more prone to the development of neoplasms.
 - a. True
 - b. False
- 3. The immune system has all of the following roles in preventing cancer except for
 - a. Protect the host against viral infection
 - b. Prevents establishment of an inflammatory environment
 - c. Destruction of host CD8+ T cells
 - d. Eliminates tumor cells
- 4. In which phase does the adaptive and innate immune system work together to destroy tumor cells long before they become clinically apparent?
 - a. Elimination
 - b. Equilibrium
 - c. Escape
 - d. Entrance
- 5. How can tumor cells inflamed tumors affect MHC class I expression?
 - a. Increase
 - b. No change
 - c. Reduce
 - d. Elimination

Webinar 2 – Immune Checkpoint Inhibition

Learning Objectives:

- A. Describe immune checkpoints
- B. Understand how checkpoints are leveraged by cancer
- C. Describe therapies that target immune checkpoints

- 1. The main inherent purpose of immune checkpoint is to rapidly multiply and propagate the immune response systemically
 - a. True

- b. False
- 2. Each of the following is a well-studied inhibitory receptor in antibody therapy except
 - a. CTLA-4
 - b. PD-1
 - c. TIM-3
 - d. CD4
- 3. Immune checkpoint inhibition molecules are thought to primarily effect what kind of cells function?
 - a. B Cells
 - b. NK Cells
 - c. Macrophages
 - d. T Cells
- 4. What kind of compound is an immune checkpoint inhibitor?
 - a. Nanoparticle
 - b. Monoclonal antibody
 - c. Steroid
 - d. Alkylating agent
- 5. Generally speaking, what kind of host environment is most likely to allow for the success of a checkpoint inhibitor?
 - a. Strong endogenous antitumor immune response
 - b. No endogenous antitumor immune response
 - c. Immunocompromised patient
 - d. Weak endogenous antitumor immune response

Webinar 3 - T Cell Mediated Immunity

Learning Objectives:

- A. Describe the T cell subtypes
- B. Understand how T cell responses are initiated
- C. Describe how cytotoxic T cells function

- 1. How do lymphocytes traffic to lymph nodes for activation?
 - a. Adhesion and chemokines
 - b. Brownian motion
 - c. Cooperatively with red blood cell circulation
 - d. They are unable to specifically reach lymph nodes
- 2. What is the primary function of a dendritic cell?
 - a. Direct destruction of tumor cells
 - b. Oxygenation of peripheral tissues
 - c. Antigen presentation
 - d. Proliferation of bacterial infections
- 3. Which cytokine drives the majority of T cell proliferation and differentiation
 - a. IL-1

- b. IL-6
- c. TNF-a
- d. IL-2
- 4. CD8 T cells primarily cause cell death through targeted and specific apoptosis based on MHCI antigen presentation
 - a. True
 - b. False
- 5. Which of the following types of T cell based immunotherapy utilizes fresh tumor digests to culture T cells for expansion and administration into patients?
 - a. Chimeric antigen receptor therapy
 - b. Genetically altered lymphocyte therapy
 - c. Tumor infiltrating lymphocyte therapy
 - d. Total tumor RNA lymphocyte therapy

Webinar 4 – Surgical Considerations for Immunotherapy in the Treatment of Malignant Glioma

Learning Objectives:

- A. Describe the goals of surgery for gliomas
- B. Understand the different surgical options in the treatment of gliomas
- C. Describe how surgery may be combined with immunotherapy

- Gross total resection confers a survival benefit in malignant glioma over subtotal resection
 - a. True
 - b. False
- 2. Which type of fluorescent compound results in the tumor specific metabolism into PPIX
 - a. Fluorescein
 - b. GFP
 - c. YFP
 - d. 5-ALA
- 3. What type of treatment modality may be best for patients with deep seated focal malignant lesions that are poor surgical candidates?
 - a. Awake craniotomy
 - b. Laser interstitial thermal therapy
 - c. Standard craniotomy
 - d. Whole brain radiation
- 4. What was shown to be the effect of neoadjuvant anti-PD1 therapy in patients with recurrent GBM who underwent surgical therapy
 - a. Improved survival outcomes
 - b. Decreased length of stay
 - c. No effect on mortality

- d. Increased rate of death
- 5. What is the effect of laser interstitial thermal therapy on the blood brain barrier?
 - a. Complete irreversible destruction of the blood brain barrier
 - b. No effect
 - c. Temporary induction of focal permeability
 - d. Reduction of blood brain barrier permeability

Webinar 5 – Immunotherapy for GBM

Learning Objectives:

- A. Describe immunotherapy strategies for GBM
- B. Understand which of immunotherapies have been tested in patients with GBM
- C. Describe how immunomodulation can be leveraged in the treatment of GBM

- 1. Younger age is associated with better survival in patients with glioblastoma (GBM)
 - a. True
 - b. False
- 2. 1p/q9q co deletion is associated with a diagnosis of
 - a. Astrocytoma
 - b. Ependymoma
 - c. Medulloblastoma
 - d. Oligodendroglioma
- 3. Immune checkpoint inhibitors are usually antibodies
 - a. True
 - b. False
- 4. Chimeric antigen receptor (CAR) T cells have not been tested in patients with GBM.
 - a. True
 - b. False
- 5. Vaccine platforms that have been tested in patients with GBM include
 - a. Neural stem cells
 - b. Fibroblasts
 - c. Dendritic cells
 - d. Red blood cells