

# Basic Principles of Cancer Immunotherapy

Bryon Johnson, PhD
Professor / Dept. Medicine, Medical College of Wisconsin









#### Disclosures

• I will not be discussing non-FDA approved indications during my presentation.









#### The Premise of Cancer Immunotherapy

- Normally, the immune system eliminates mutated and/or damaged cells
- To exist, tumors must evolve mechanisms to locally disable and/or evade the immune system.

The goal of immunotherapy is to restore the capacity of the immune system to recognize and reject cancer.









# Why Does the Immune System Fail to Eliminate Cancer?

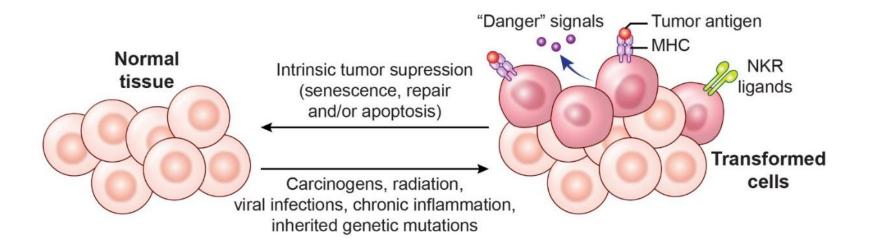
- Cancer cells grow progressively in immunocompetent hosts without evidence of <u>T cell exhaustion</u> or <u>systemic anergy</u>
  - T cell Exhaustion: CD8+ T cells often become dysfunctional, entering a state known as exhaustion, during certain chronic infections or when they enter a suppressive tumor microenvironment
  - Systemic Anergy: A state of immune unresponsiveness. Induced when the T cell's antigen receptor is stimulated, effectively freezing T cell responses pending a "second signal" from the antigen-presenting cell

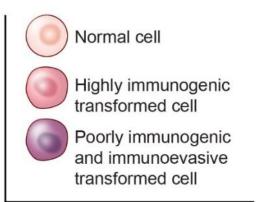










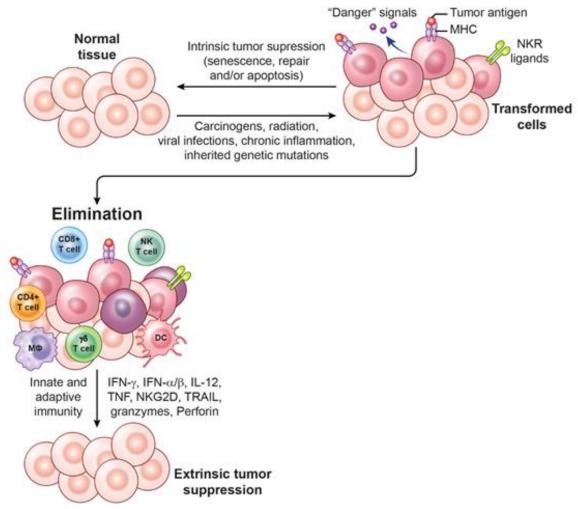


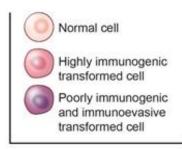










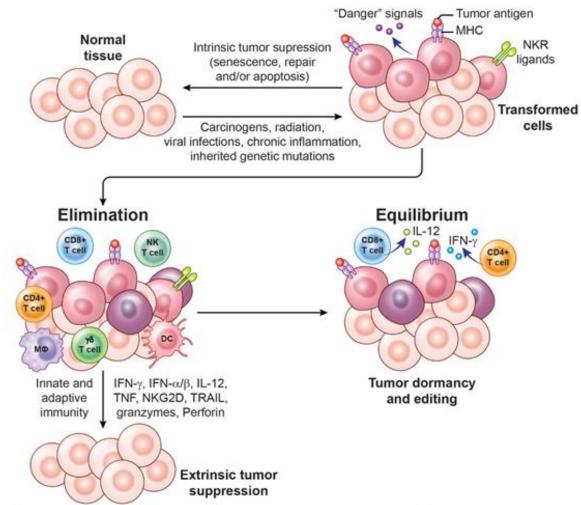


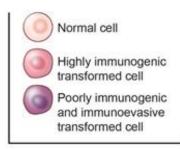










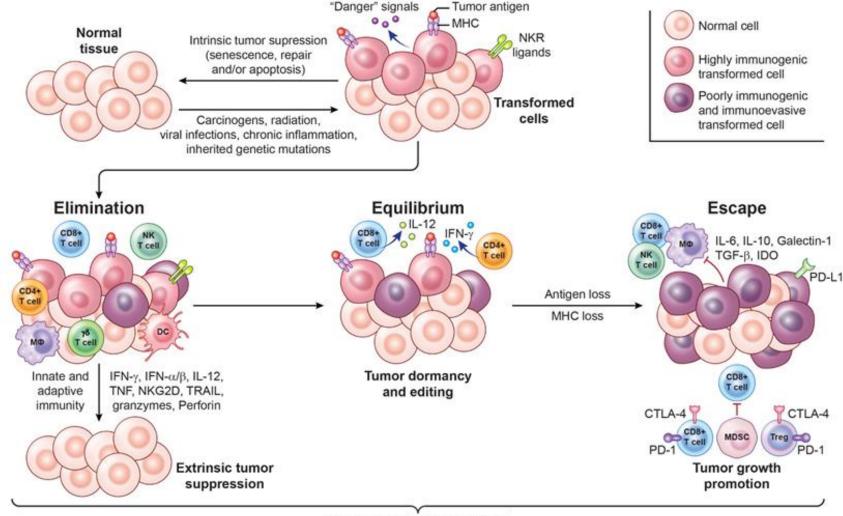














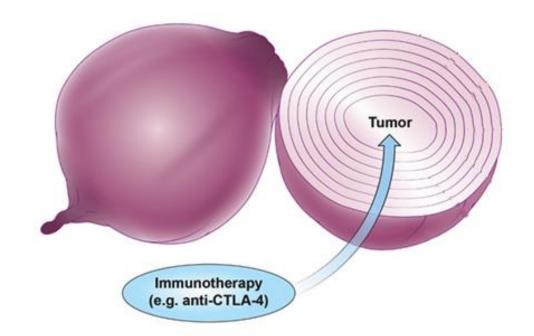






#### Multi-layered Immunosuppression

- Tumors insulate themselves with dense layers of immunosuppressive stroma
- Overcoming the many layers of interconnected and often functionally redundant immune suppressive mechanisms represents a daunting challenge for tumor-specific T cells
- Immunotherapy can "peel back" the layers of local immune suppression, thereby restoring the capacity of T cells to eradicate the tumor



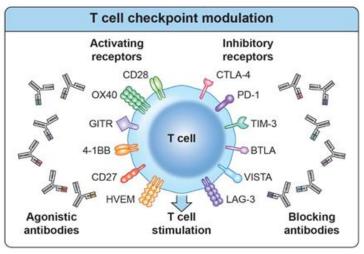


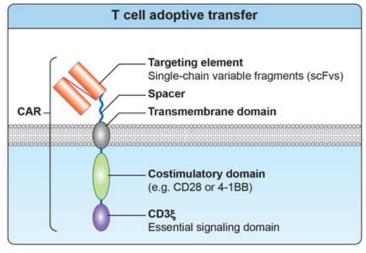


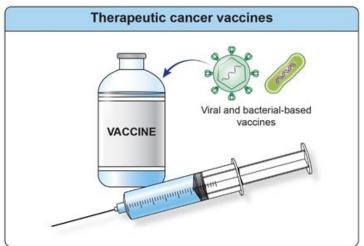


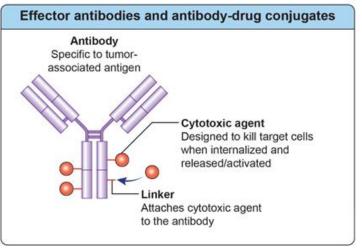


### Types of Immunotherapy









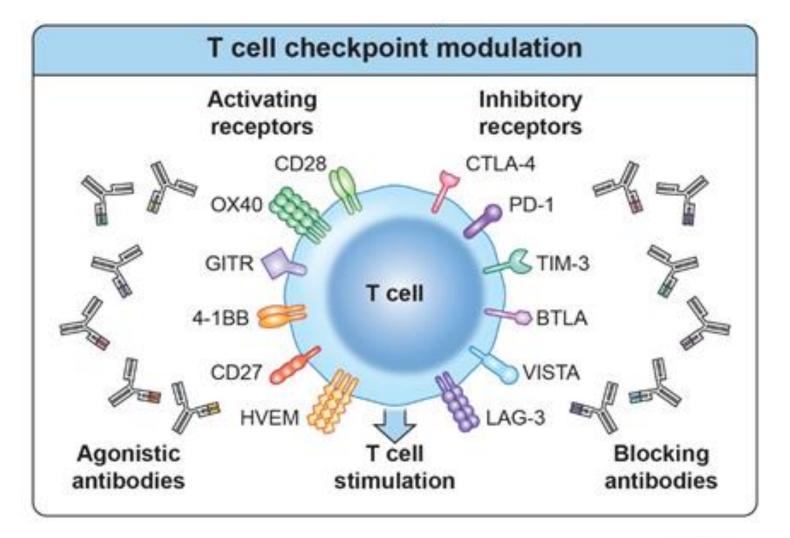








### T cell Checkpoint Modulation



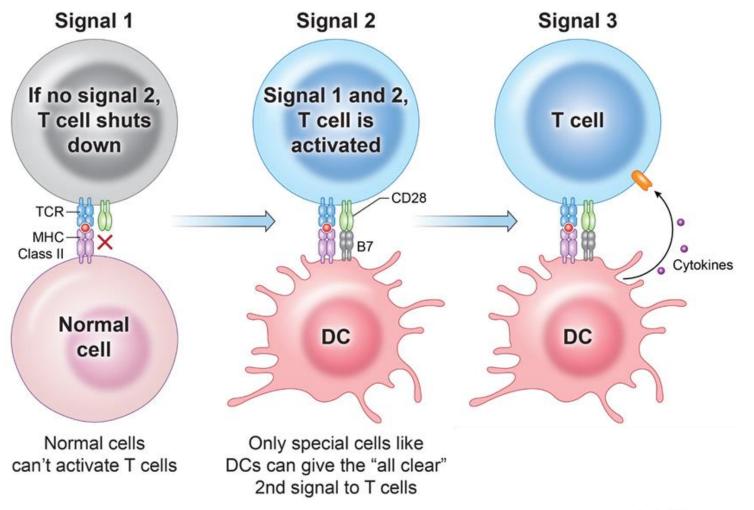








### Antigen-Specific T Cell Activation





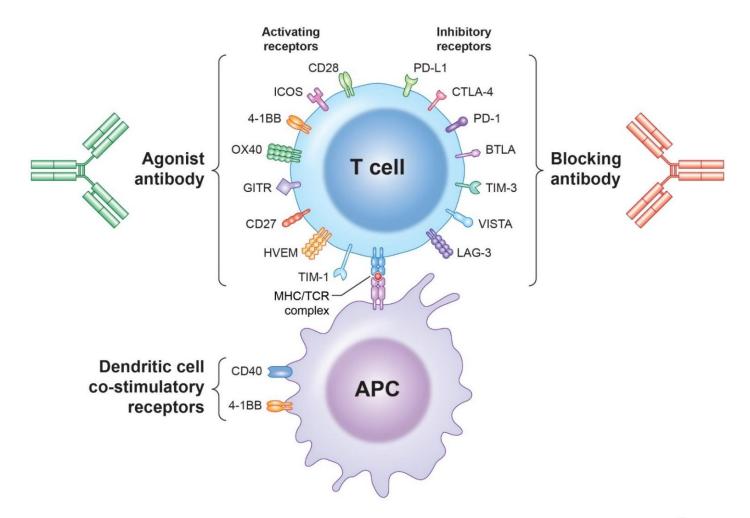






#### T Cell Checkpoint Modulation

- To exist, tumors must evolve mechanisms to locally disable and/or evade the immune system.
- The goal of T cell checkpoint blockade is to make T cell "off-switches" inaccessible to tumor cells, thus restoring tumor-specific immunity.





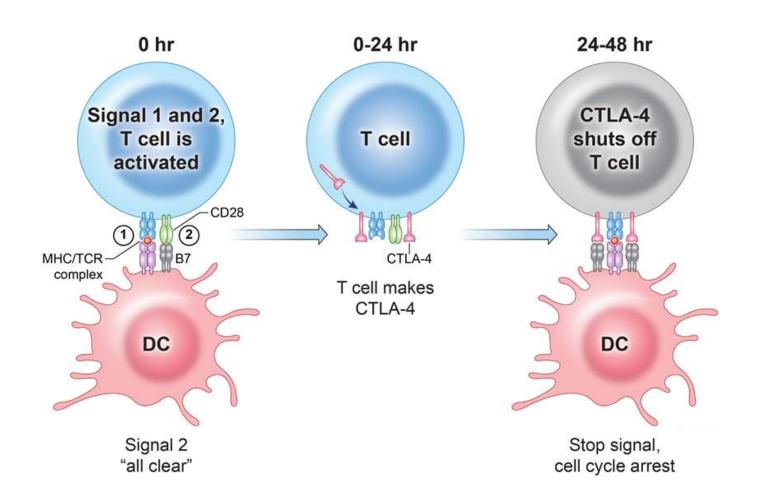






#### The CTLA-4 Checkpoint

- <u>C</u>ytotoxic <u>T</u>-<u>L</u>ymphocyte
   <u>A</u>ssociated Protein <u>4</u>
- Also known as CD152
- Negative regulator of T cell activation



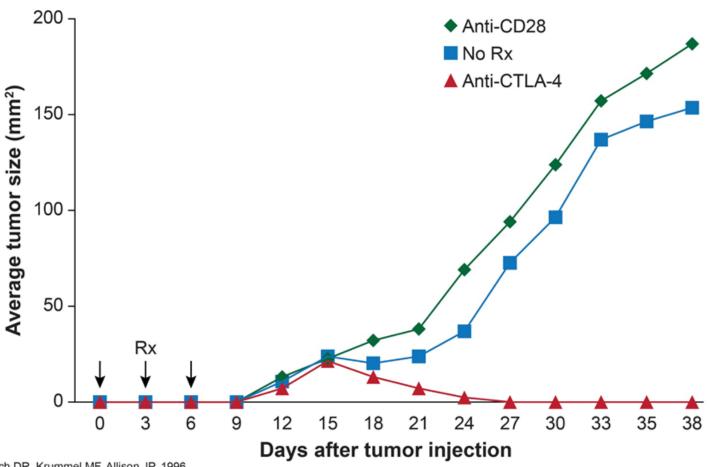








# Anti-CTLA-4 Induces Regression of a Transplantable Colon Carcinoma



Leach DR, Krummel MF, Allison JP. 1996. Enhancement of antitumor immunity by CTLA-4 blockade. Science. 217(5256): 1734-6.



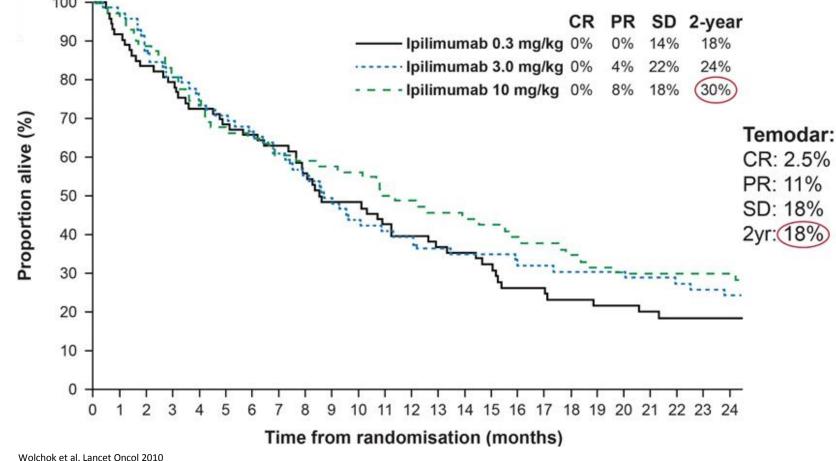






## Ipilimumab (Human anti-CTLA-4)

 Granted FDA approval for treatment of patients with metastatic melanoma in 2010



olchok et al. Lancet Oncol 2010



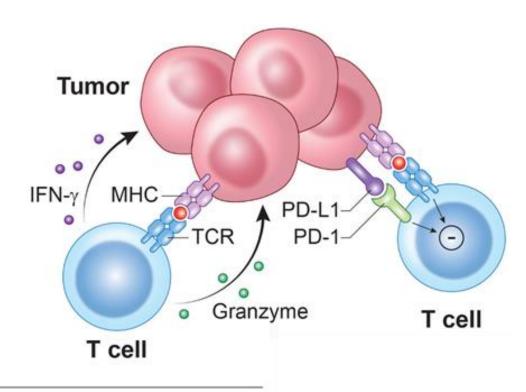






#### The PD-1/PD-L1 Checkpoint

- Promotes T cell tolerization through inhibiting activation signaling
- T cell PD-1 interacts with PD-L1 and PD-L2
- Many cells express PD-L1/PD-L2 and can suppress T cell activation
- Tumors express PD-L1 through two primary mechanisms
  - TIL production of IFN-y
  - Oncogenic signaling pathways



Francisco, L. et al. *Immunol Rev.* 2010. 236: 219. Pardoll, D.M. *Nat Rev Cancer*. 2012. 12: 252.



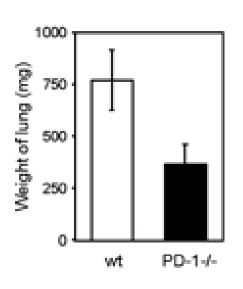




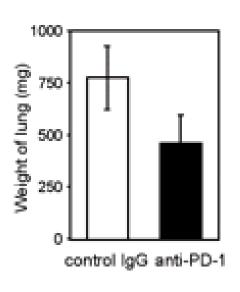


# Anti-PD-1 Slows Tumor Growth in Pre-clinical Models

• PD-1 deletion or inhibition reduced CT26 colon cancer cell growth in BALB/c mice









Iwai et al. Internat. Immunol 2004



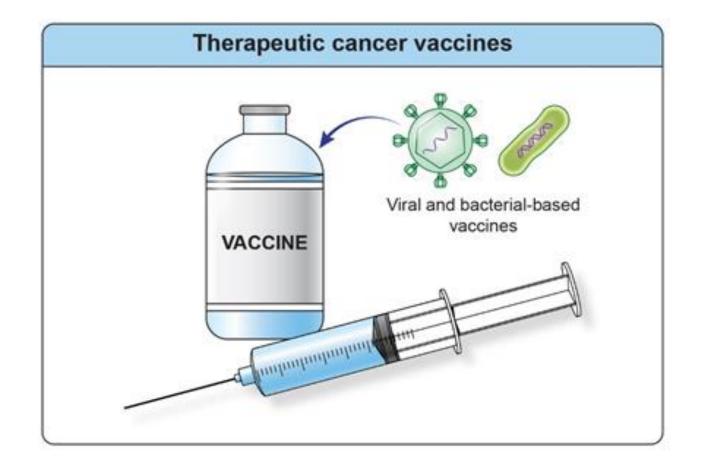






#### Therapeutic Cancer Vaccines

- The goal of therapeutic cancer vaccination is to increase the immunogenicity of tumor antigens in order to generate a high frequency of tumor-specific T cells.
- To date, therapeutic vaccines have not been very effective in the clinic.
   A new emerging area for improving the efficacy of cancer vaccines is using them as cancer-preventive therapeutics (immunoprevention).



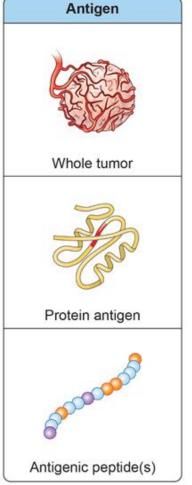


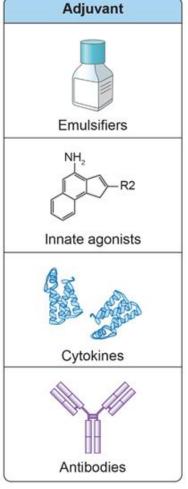


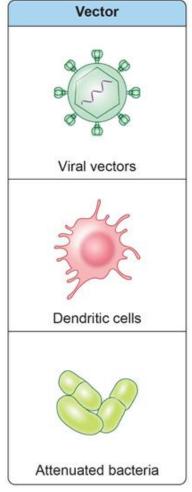


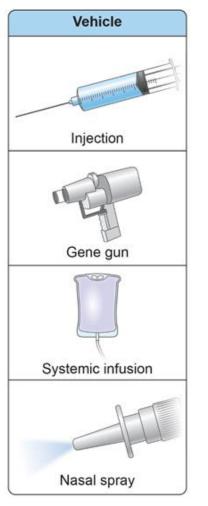


### Components of a Cancer Vaccine









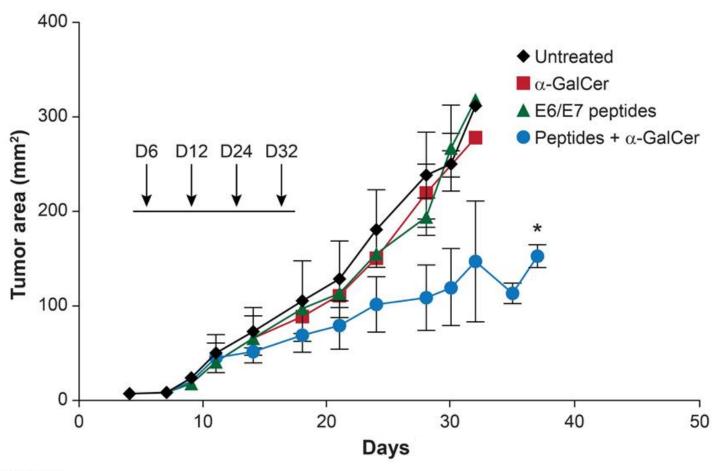








# An Intra-Nasal HPV E6/E7: α-GalCer Vaccine Slows Growth of TC-1 Tumors







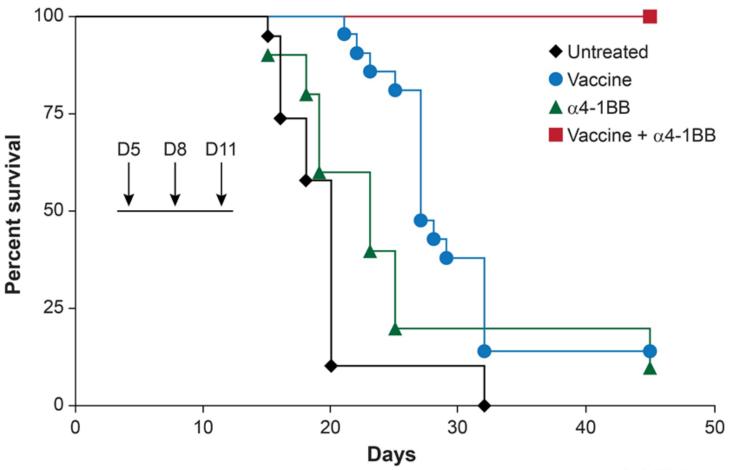






# 4-1BB Agonist Antibody and HPV E6/E7 Vaccine Synergize in Curing TC-1 Tumors

(combining with other agents to improve vaccine efficacy)







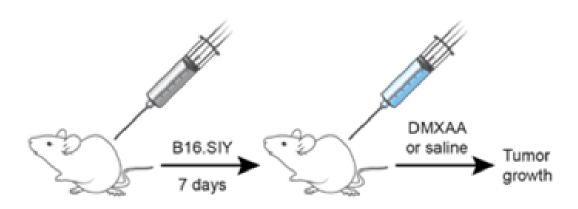


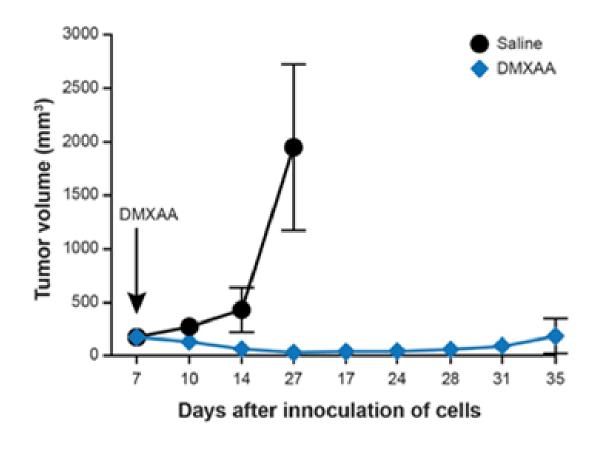




# Intratumoral Injection of Innate Immune Agonists: *Direct Vaccination Approach*

 Intratumoral DMXAA (mouse STING agonist) triggers rejection of B16 melanoma







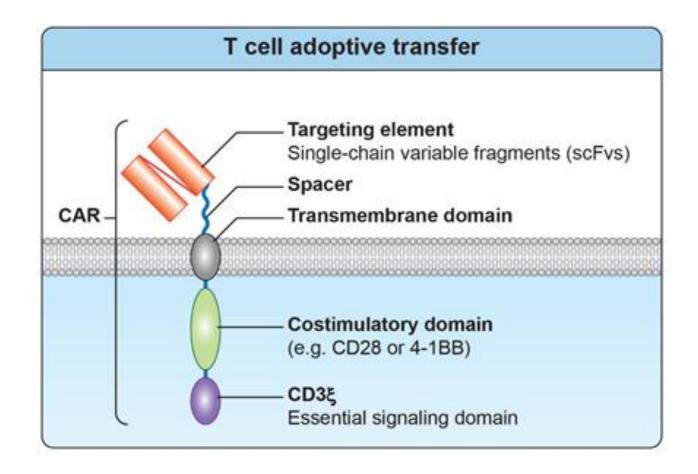






#### Adoptive Cell Transfer

 The goal of adoptive cell transfer is to overwhelm the tumor with a higher frequency of tumorspecific immune cells and/or engineer immune cells to target cancer



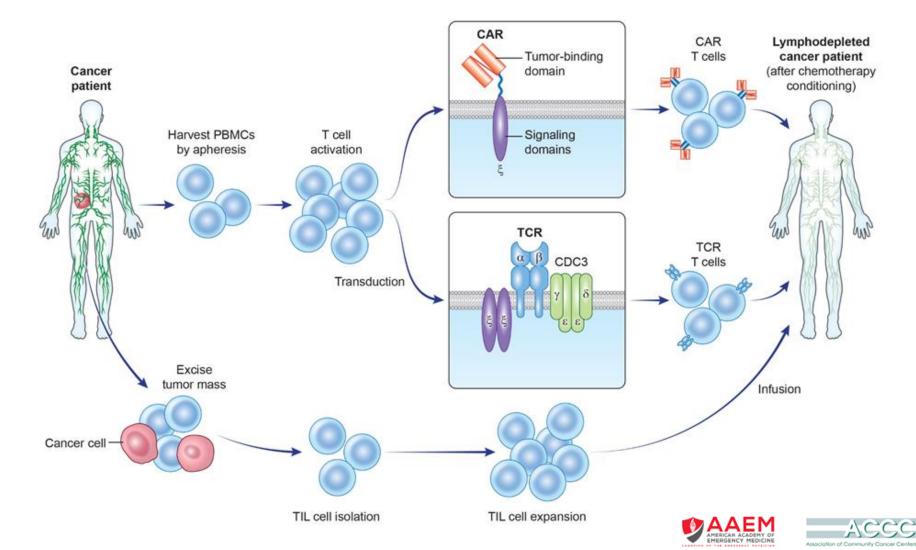






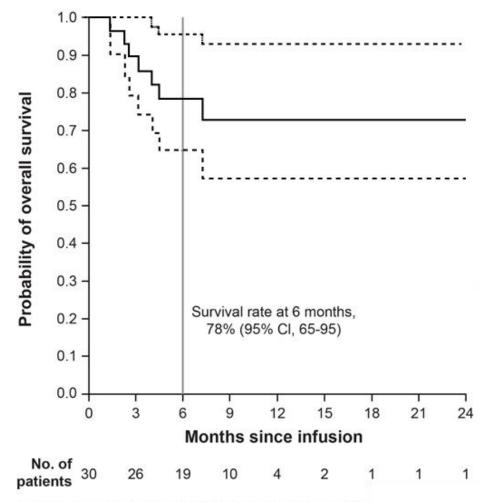


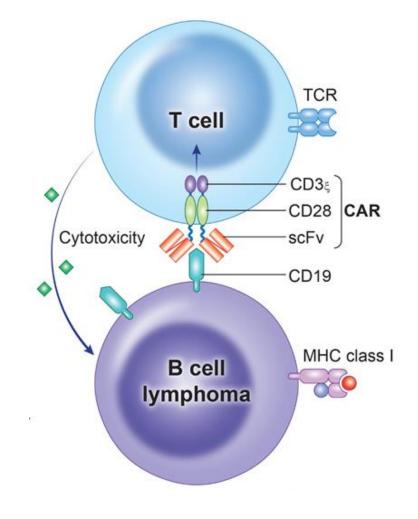
### **Adoptive Cell Therapy Process**

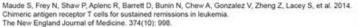




# CD19 CAR T Cell Therapy for Relapsed B Cell ALL









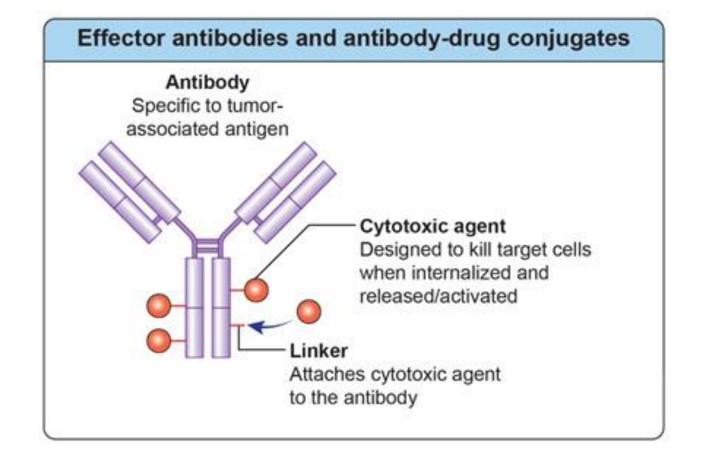






### Effector Antibodies and Antibody-Drug Conjugates (ADCs)

 The goal of effector antibodies is to specifically target and kill tumor cells using innate mechanisms which are difficult to evade of suppress and/or through delivery of cytotoxic agents











### Key ADC/Antibody Principles

• **Specificity:** The more tumor specific the target antigen is, the higher the agent can be dosed without limiting toxicity

 Internalization: The target tumor surface protein must internalize to deliver the toxin – it should do so frequently and to a suitable endosomal compartment

• **Stability:** The toxin must remain inert and tethered to the antibody until it is delivered to its target cell



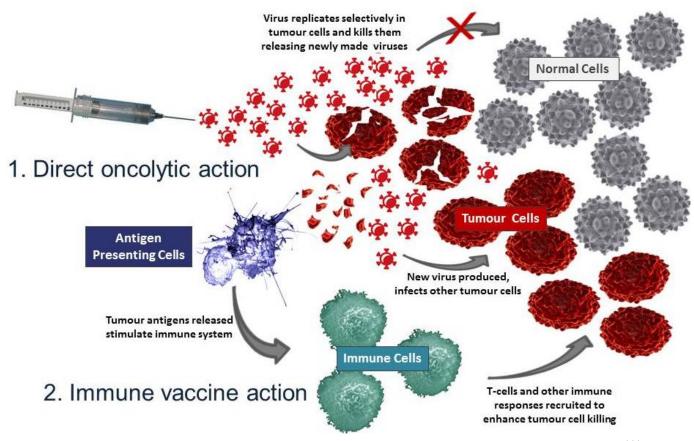






#### **Oncolytic Viruses**

 The goal of an oncolytic virus is to specifically target and kill tumor cells through viral replication



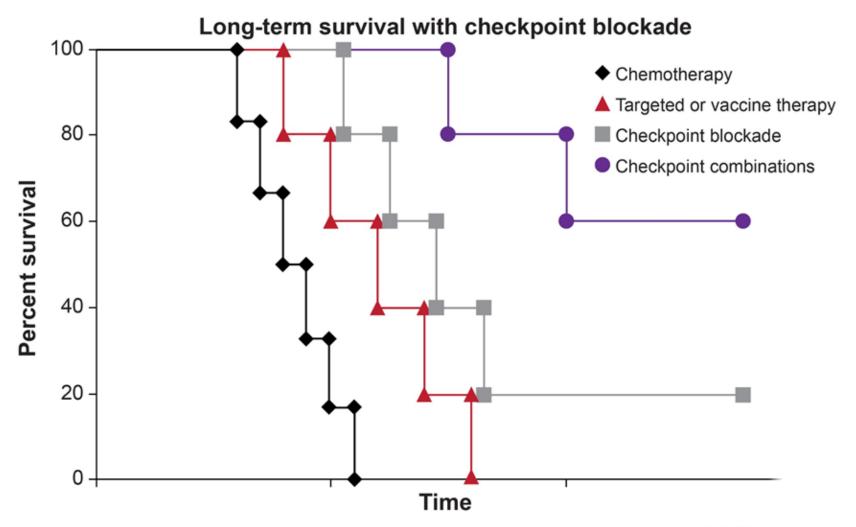
dddmag.com











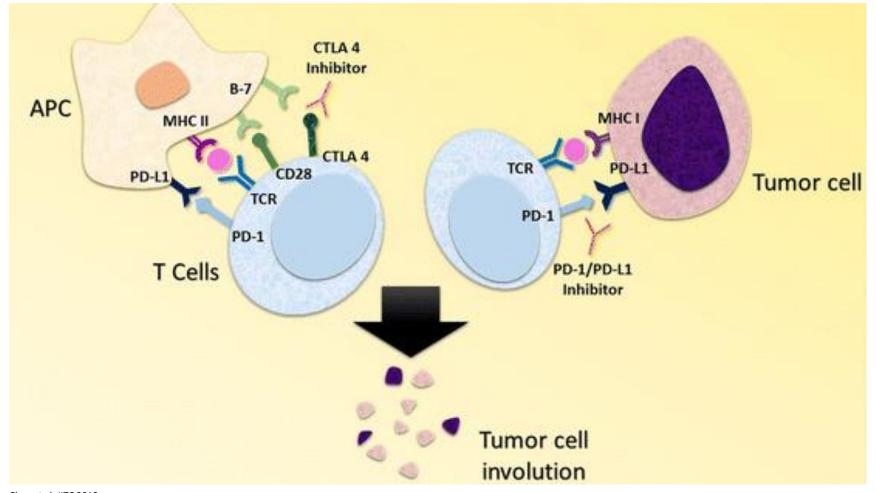








Dual CTLA-4 and PD-1 inhibition



Chae et al. JITC 2018

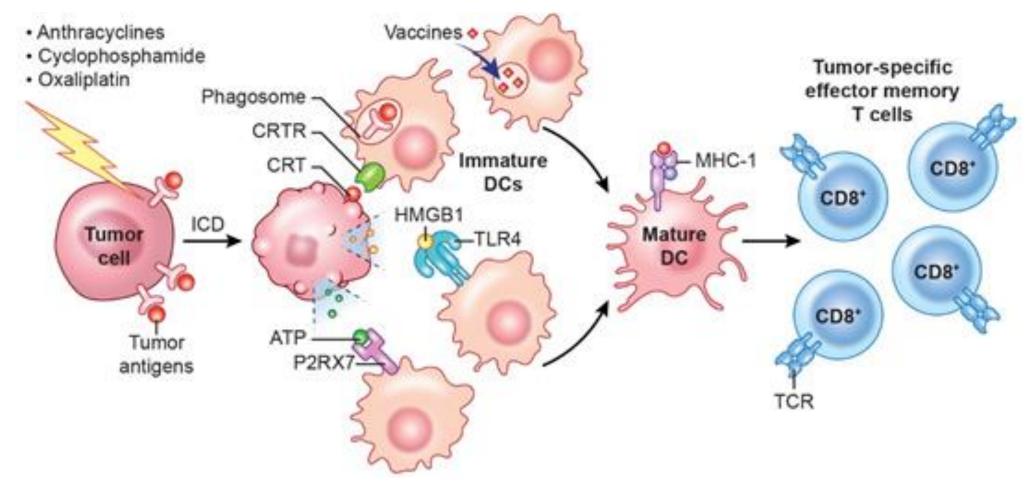








(Chemotherapy can induce an immune response)



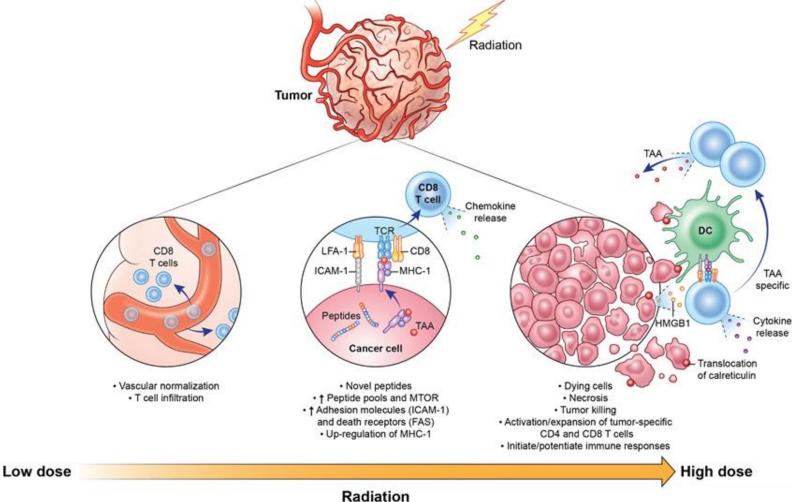








(Radiotherapy can induce an immune response)



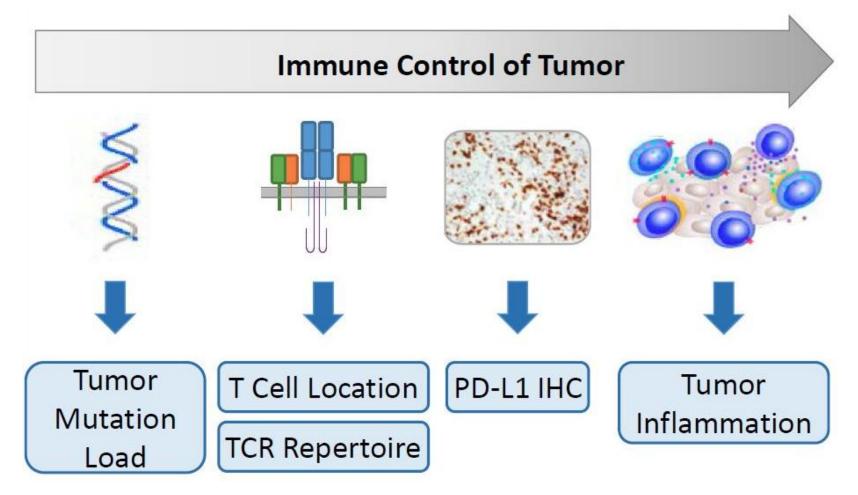








# **Immunotherapy Biomarkers**



Cesano et al. Biomedicines 2018

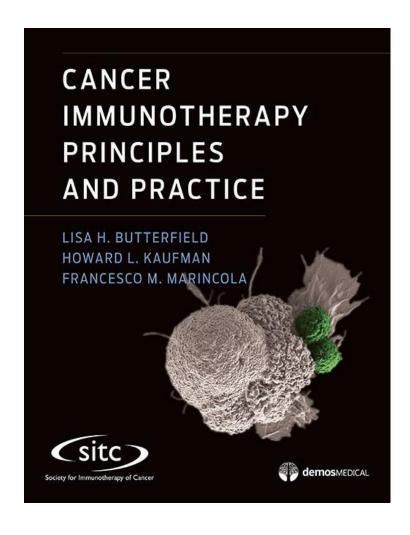








#### **Further Resources**



#### **SOCIETY FOR IMMUNOTHERAPY OF CANCER**







