

& MOMENTUM

SITC 2016-2018 Strategic Plan Goals

Education

Serve as the leading resource for information and education on cancer immunotherapy

Collaboration

Cultivate meaningful Relationships with key strategic partners

Engagement

Create a network of cancer immunotherapy stakeholders, providing greater Opportunities for engagement and interaction

Global Impact

Advance the science and application of cancer immunotherapy world wide

Scientific Research

Challenge the thinking and seek the best research in the exploration and development of tumor immunology and cancer immunotherapy



Society for Immunotherapy of Cancer

The History of SITC. Established in 1984

SITC is a non-profit organization comprised of influential basic and translational scientists, clinicians, health care professionals, government leaders and industry professionals around the globe. Through an emphasis on high-caliber scientific meetings, education and outreach activities, initiatives of major importance to the field, and commitment to collaborations, SITC aims to make cancer immunotherapy one of the four standards of care and the word "cure" a reality for cancer patients everywhere. Be a part of the future of cancer care, join SITC Today!



Society for Immunotherapy of Cancer

MILESTONES & MOMENTUM

*30 Years of Advancing
Cancer Immunotherapy*

MILESTONES

The History of Immunotherapy

Immunotherapy is a cancer treatment more than 100 years in the making, beginning most notably with Dr. William B. Coley, who worked with patients and doctors to study how cancer tumors reacted to bacterial infections. He treated cancer patients with inoperable tumors by injecting a combination of bacteria, which became known as Coley's Toxins, directly into their tumors. His results showed that this kind of treatment shrank the tumors and sometimes even cured the patient. He believed that the body's increased response to the bacteria also helped fight off the cancer. In the modern era, Dr. Donald Morton was an early proponent of immunotherapy, particularly cancer vaccines. His work with bacillus Calmette-Guerin (BCG) for melanoma led to the use – and eventual approval – of BCG for bladder cancer, the first successful immunotherapy treatment against a human tumor.

1890

Dr. Coley creates the first immunotherapy treatment when he injects bacteria directly into inoperable tumors to stimulate an immune response that fights the cancer

1973

Dendritic cells are identified as "antigen-presenting cells" (APCs)

1984

Society for Immunotherapy of Cancer (formally SBT) is founded

1986

The Extramural IL-2/LAK Working Group is formed with funding from the National Cancer Institute to confirm results of the high-dose IL-2/LAK cell regimen in the treatment of melanoma and renal cell cancer

1990

- Bacillus Calmette-Guerin is approved for bladder cancer
- First treatment with genetically modified TIL

1992

High dose IL-2 is approved to treat metastatic kidney cancer

1999

Denileukin diftitox (Ontak), a fusion of IL-2 and diphtheria toxin, is approved to treat lymphomas

1956

Tumor-specific cell antigens are discovered

1978

- Bacillus Calmette-Guerin (BCG) is first studied as a possible treatment
- Interleukin-2 (IL-2) is discovered
- Tumor-specific monoclonal antibodies (mAbs) are discovered

1985

- Results from IL-2 and lymphokine activated killer cell (LAK) therapy in various tumors are first reported
- Adoptive T cell transfers are studied as a possible cancer treatment

1988

First results for tumor-infiltrating lymphocytes (TILs) therapy are reported

1991

Sargramostim (Leukine), a granulocyte macrophage-colony stimulating factor (GM-CSF), is approved to boost white blood cell counts

1996

Interferon alfa-2b (IntronA) is approved for the adjuvant treatment of high-risk melanoma

1997

The first monoclonal antibody (mAb), rituximab (Rituxan), is approved to treat B cell malignancies

1998

- IL-2 is approved to treat metastatic melanoma
- The mAb drug trastuzumab (Herceptin) is approved to treat breast cancer

2000

The drug gemtuzumab ozogamicin (Mylotarg), which combines a mAb and a toxin from the bacteria *Micromonospora echinospora*, is approved for AML

2011

- Pegylated interferon (PEG-Intron) is approved for adjuvant therapy of selected melanoma patients
- Ipilimumab (Yervoy) is approved to treat advanced melanoma

2014

- Blinatumomab (Blinicyto), a monoclonal antibody, is approved for ALL.
- Nivolumab (Opdivo), a PD-1 inhibitor, is approved for advanced melanoma
- Pembrolizumab (Keytruda) is the first PD-1 inhibitor approved for advanced melanoma
- Ramucirumab (Cyramza), a monoclonal antibody, is approved for advanced gastric and lung cancers

2009

Anti-CD20 monoclonal antibody of atumumab (Arzerra) is approved for CLL

2010

- The first therapeutic cancer vaccine, sipuleucel-T (Provenge), is approved for advanced prostate cancer
- Gemtuzumab ozogamicin is discontinued due to safety concerns and lack of benefit

2012

Several clinical studies of T cell checkpoint inhibitors targeting PD-1 and PD-L1 demonstrate therapeutic activity in many types of cancers

2013

- The first phase III trial of oncolytic virus immunotherapy shows improvement in long-term response rate in melanoma patients
- The combination of agents targeting CTLA-4 and PD-1 checkpoints shows activity against melanoma

2015

- The first biosimilar product, filgrastim-sndz (Zarxio), is approved to treat severe chronic neutropenia
- Nivolumab (Opdivo) is the first checkpoint inhibitor approved for lung cancer
- Ramucirumab (Cyramza) is approved for metastatic colorectal cancer