

Clinical Trials Program

Phase Ib trial of pembrolizumab administered following adoptive cell therapy- A multiple cohort study; The ACTIVATE (Adoptive Cell Therapy InVigorated to Augment Tumor **Eradication**) Trial.

April 23rd, 2018 to March 31st, 2020

Highlights

- This therapeutic approach uses immune checkpoint inhibitors, a novel class of anti-cancer drugs and adoptive cell transfer (ACT) to eliminate cancer cells
- •A number of preclinical studies suggest that a combination of checkpoint inhibitors and ACT could result in improved tumor control
- Pembrolizumab is a potent and highly selective humanized monoclonal antibody designed to directly block the interaction between PD-1 and its ligands PD-L1 and PD-L2
- Results from this study will determine the safety and effectiveness of the combination therapy and enable further development of cell therapies for Canadians suffering from different forms of cancer

trials of TILs.



Immune checkpoint inhibitors are a new class of anti-cancer drug. Immune checkpoints are the "brakes" of the immune system, and sometimes cancer cells use these checkpoints to prevent immune cells from attacking the cancer. These drugs block this process, allowing immune cells to eliminate cancer cells. However, for most cancers, tumor shrinkage occurs in only a minority of patients after treatment. As with ACT, improvements are needed to benefit more patients. It has been observed in the laboratory and in some patients that combining checkpoint inhibitors with ACT shows the potential for improved responses. Given these promising results, we are performing a clinical study of the combination of TILs and the immune checkpoint inhibitor pembrolizumab. In this study, we are investigating the feasibility, safety and potential clinical benefit of this novel approach in cancer patients who are left with few therapeutic options.

Clinical trial sites and investigators

Key

Milestónes

Toronto

Princess Margaret Cancer Centre Dr. Marcus Butler

Clinical Team Members Dr. Marcus Butler Dr. Anna Spreafico Dr. Aaron Hansen

Scientific Team Dr. Pamela Ohashi Dr. Naoto Hirano Dr. Trevor Pugh Melania Pintille

Cell Production Team Dr. Linh Nguyen

Partner contributions

Merck Canada Inc. \$1,055,424 (cash & in-kind) Princess Margaret Cancer Foundation \$875,000 (cash) Trial Sponsor.

YEAR 1

- TIL products manufactured for 10 patients by Q4 end
- Enrollment of 10 patients on Cohort 1 of the ACTIVATE study by Q4; 4 patients have already been enrolled and treated with TIL by time of this application submission
- Correlative testing initiated on all treated patients by Q4 end

YEAR 2

- Completion of all TIL manufacture by $\mathsf{Q2}$
- Enrollment of 2 additional patients in Cohort 1 by Q2
- Last patient treated by Q2
- Clinical data collected, QC monitored, and finalized by beginning of Q4 (January 2020)
- Initial response and correlative data evaluation during Q4
- First draft of manuscript by Q4 end (March 2020)

The power to kill cancer lies within us. Let's tell our bodies how.

Biotherapeutics for Cancer Treatment Biothérapies pour le traitement du cancer