

Your Body. Your Hope. Your Cure.

Retooling Your Immune System In Vivo to Fight Cancer

ASGCT, May 2021

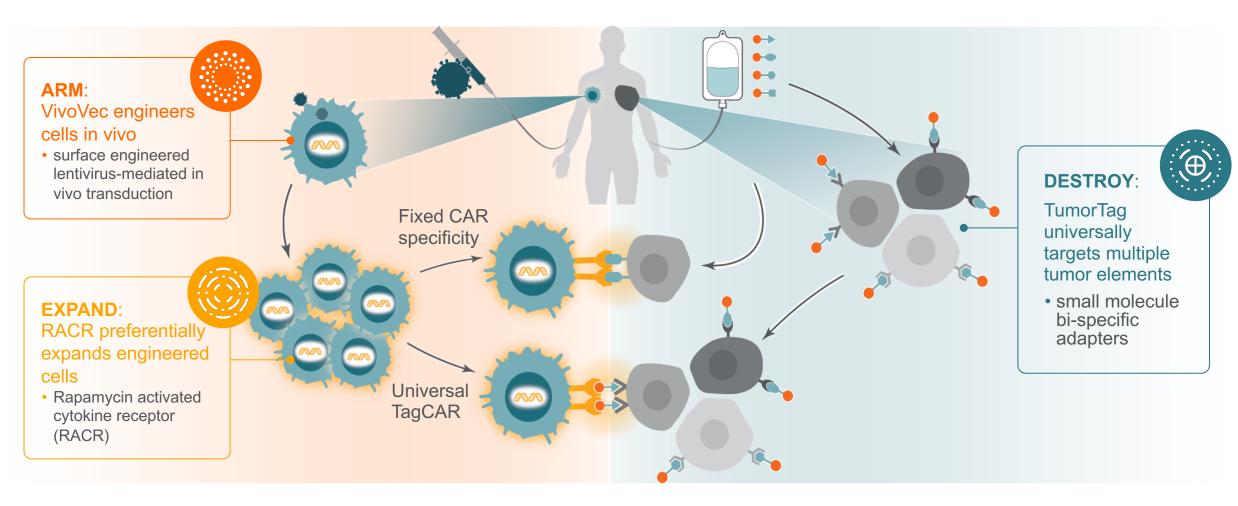


Our mission is to advance and improve access to immunotherapy by retooling the patient's immune system in vivo, freeing them from the burden of cancer in their daily lives



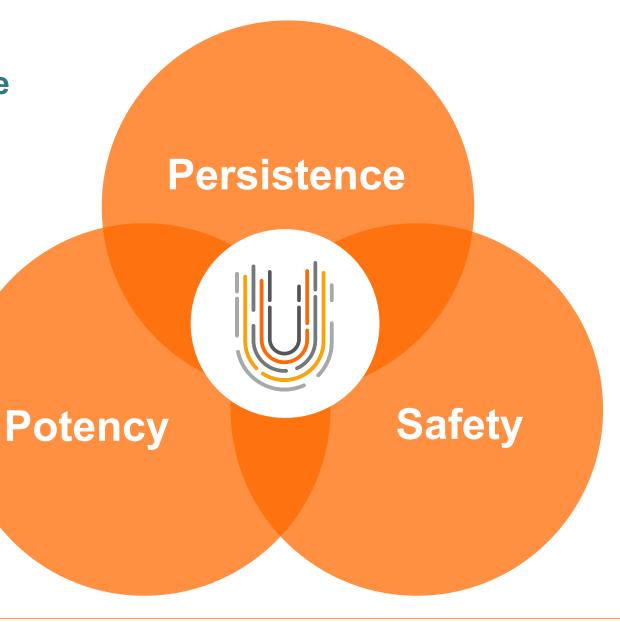


Umoja's integrated immunotherapy platform provides solutions to the challenges in both blood and solid tumor CAR-T therapies



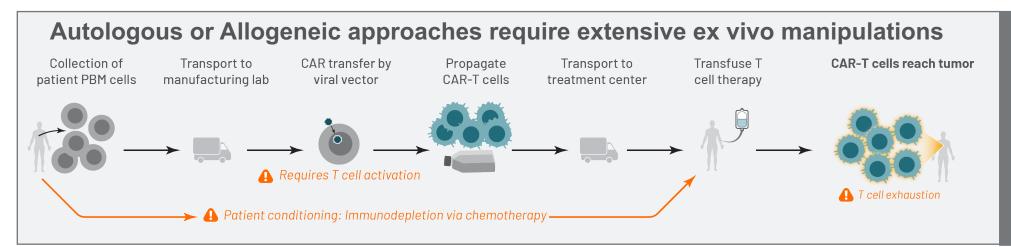


Umoja's platform captures multiple key potency attributes associated with <u>autologous CAR-T cells since</u> it is compatible with the patient's own immune system...



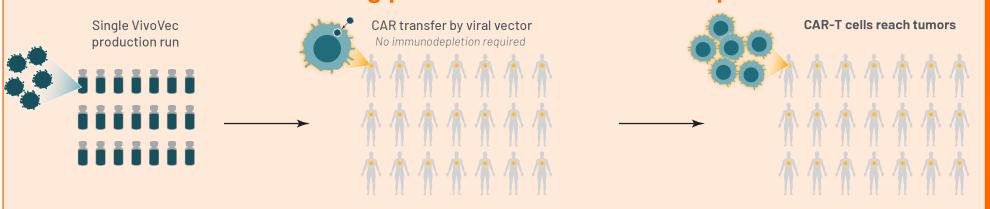


... while expanding convenience and scalability beyond allogeneic products



Logistically complex, expensive, introduces less-than ideal activation and expansion conditions that may lead to unfavorable T cell health

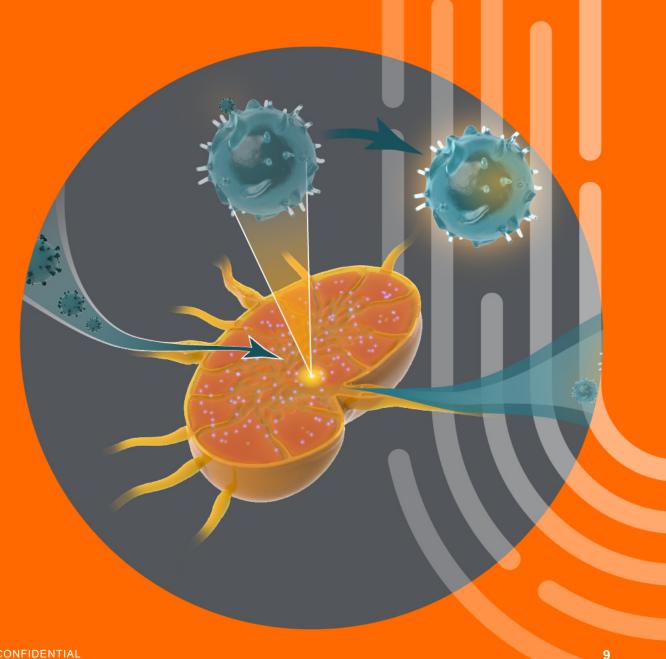




One lot treats many patients who "manufacture" their own CAR-T cells with minimal manipulation and with potentially enhanced potency



VivoVec In vivo CAR T cell generation





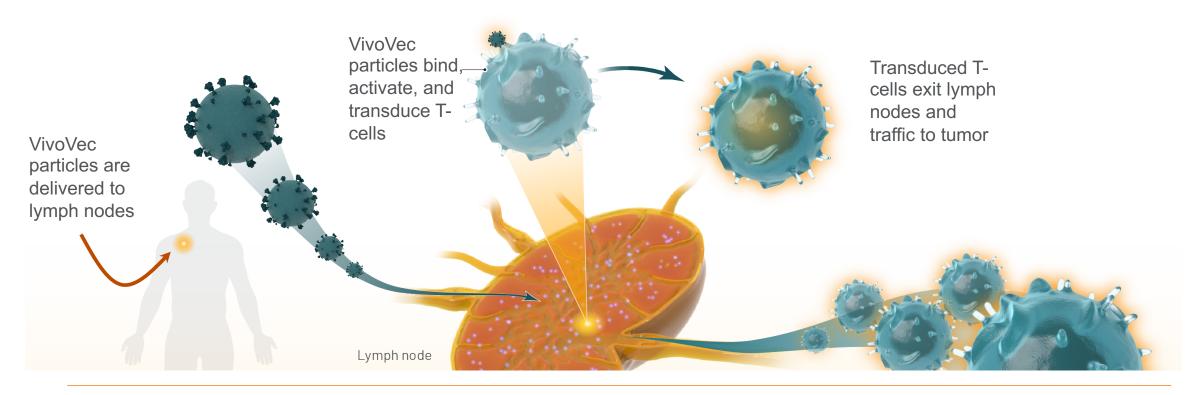
VivoVec platform solves the technical barriers to *in vivo* genetic engineering of T cells

Technical hurdles for in vivo genetic engineering "Condition"/activate T cells for efficient transduction	VivoVec Solutions	
	⊘	Lentivirus surface engineering for efficient T cell activation and transduction in vivo
In vivo expansion of engineered T cells	⊘	Drug-regulated cytokine receptor in the payload enables in vivo stimulation and expansion of transduced cells
Avoid exhaustion during expansion	\odot	"Natural" expansion process in the body maintains high potency
VSV-G enveloped lenti particles are highly immunogenic and rapidly rejected	\odot	Novel glycoprotein reduces potential for immunogenicity (relative to VSV-G)



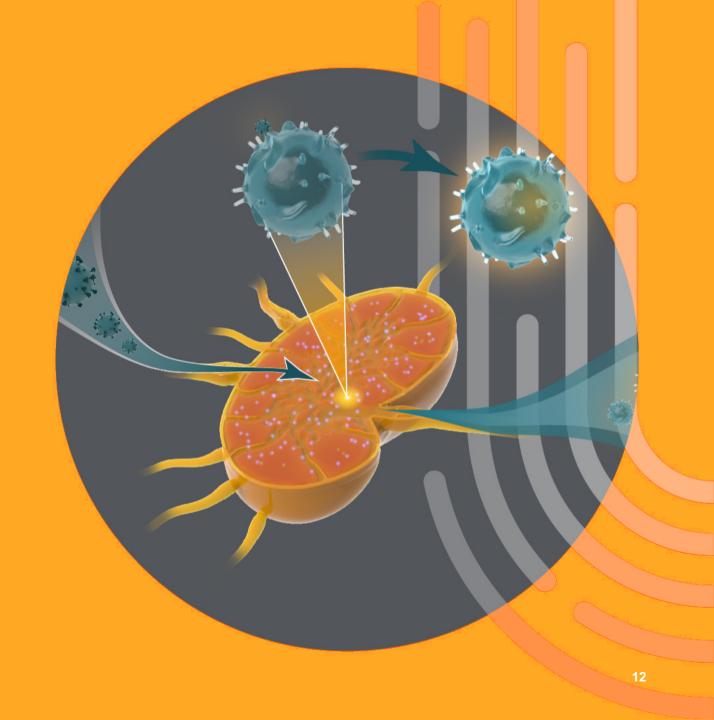
Foundational concept: lymph nodes are nature's optimized T cell "manufactory"

Umoja leverages a deep understanding of the human immune system's physiology for its proprietary approach to *in vivo* T cell engineering

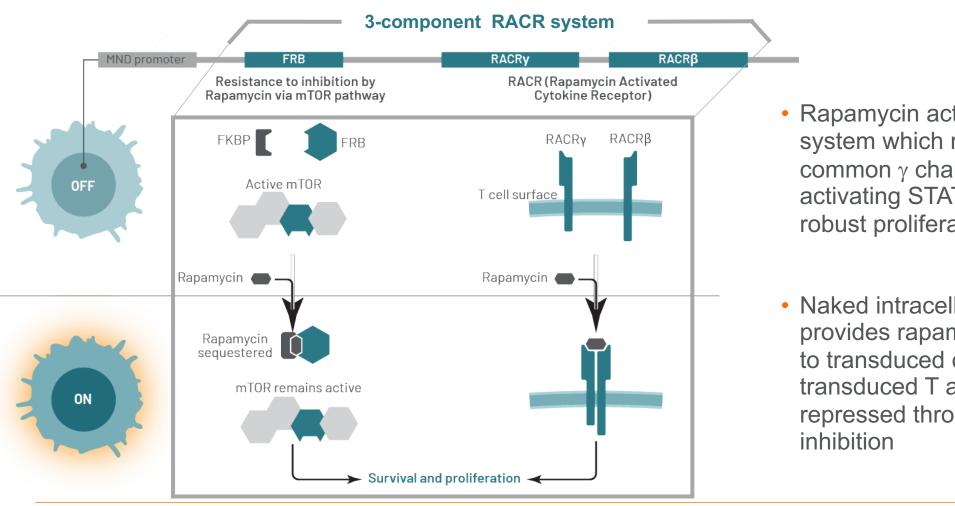




RACR
In vivo CAR T
cell expansion



RACR: Rapamycin Activated Cytokine Receptor provides control over expansion

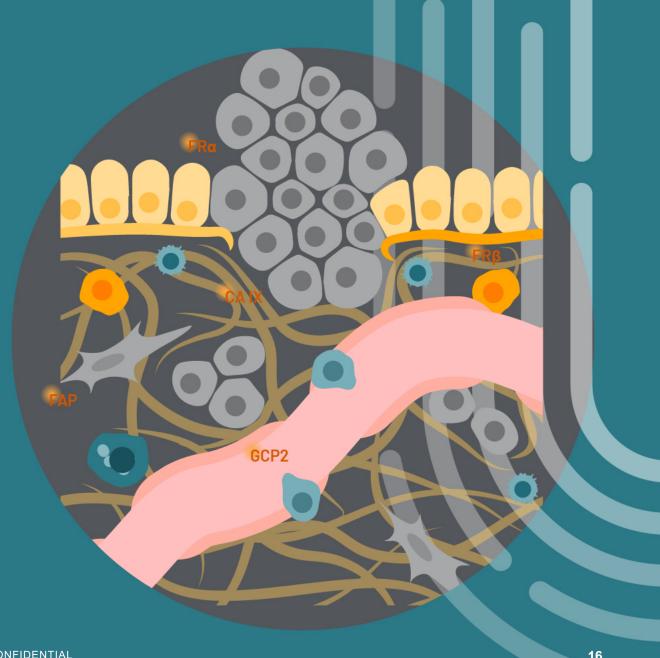


- Rapamycin activates the RACR system which replicates common γ chain cytokine activating STAT5 signaling for robust proliferation and survival
- Naked intracellular FRB domain provides rapamycin resistance to transduced cells while non transduced T and B cells are repressed through mTOR inhibition



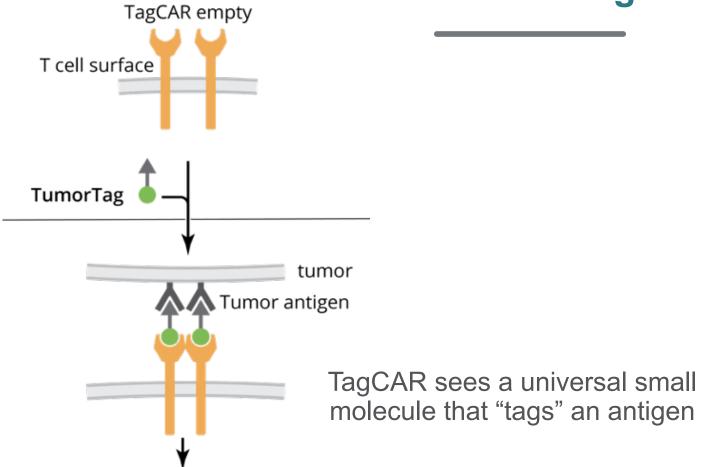
TumorTag

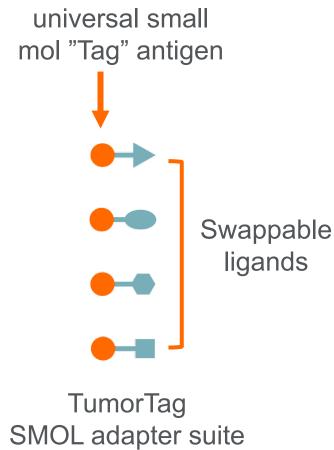
Controllable Combinatorial **Targeting of Tumor and Stroma**





Targeting with adapters for broad tumor and stroma recognition









Proliferation and activation

UB-VV100

In vivo CD19 CAR T generation for the treatment of B cell malignancies

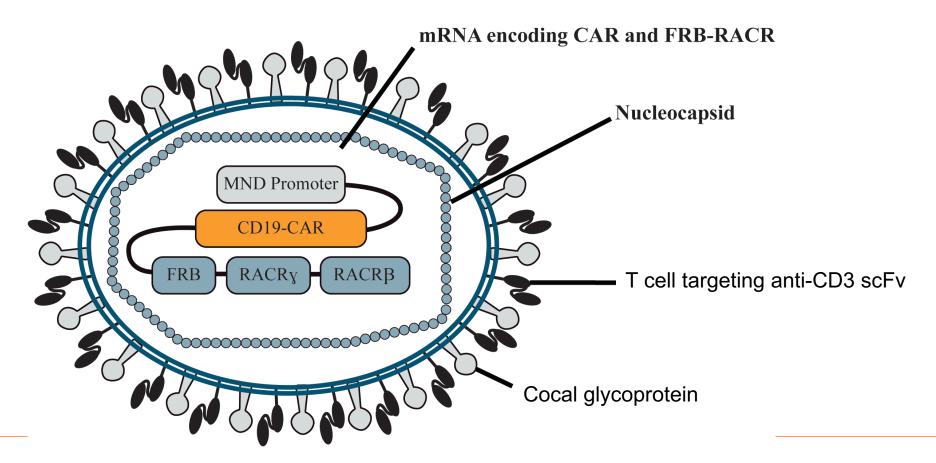




UB-VV100 drug product:

A 3rd generation, self-inactivating, replication-incompetent lentivirus designed for direct injection into patients to target T cells and deliver a payload consisting of a 2rd gen anti-CD19 CAR and a rapamycin-activated cytokine receptor (RACR) system for the treatment of B cell malignancies.

Abstract 637 Session 519 Poster #1210







Thank you

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