

TMUNITY



UNCANCER THE WORLD™

RELENTLESSLY ENGINEERING THE BEST T CELL
MEDICINES FOR SOLID TUMOR PATIENTS

WHY TMUNITY?

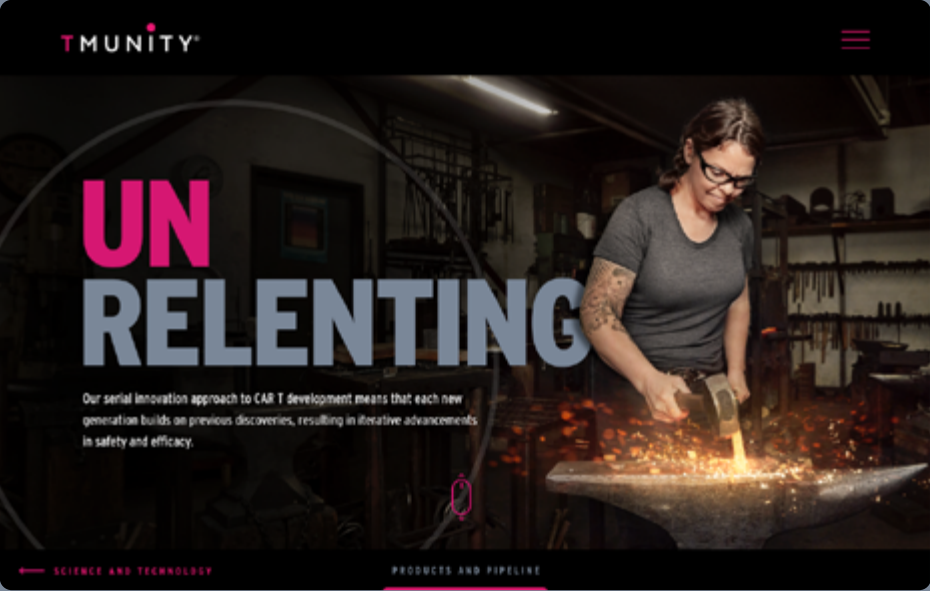
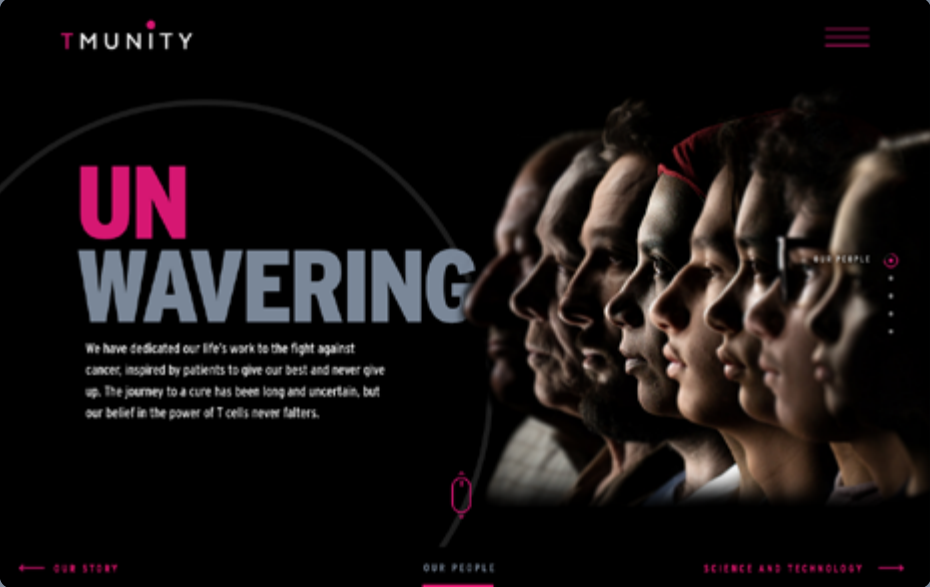
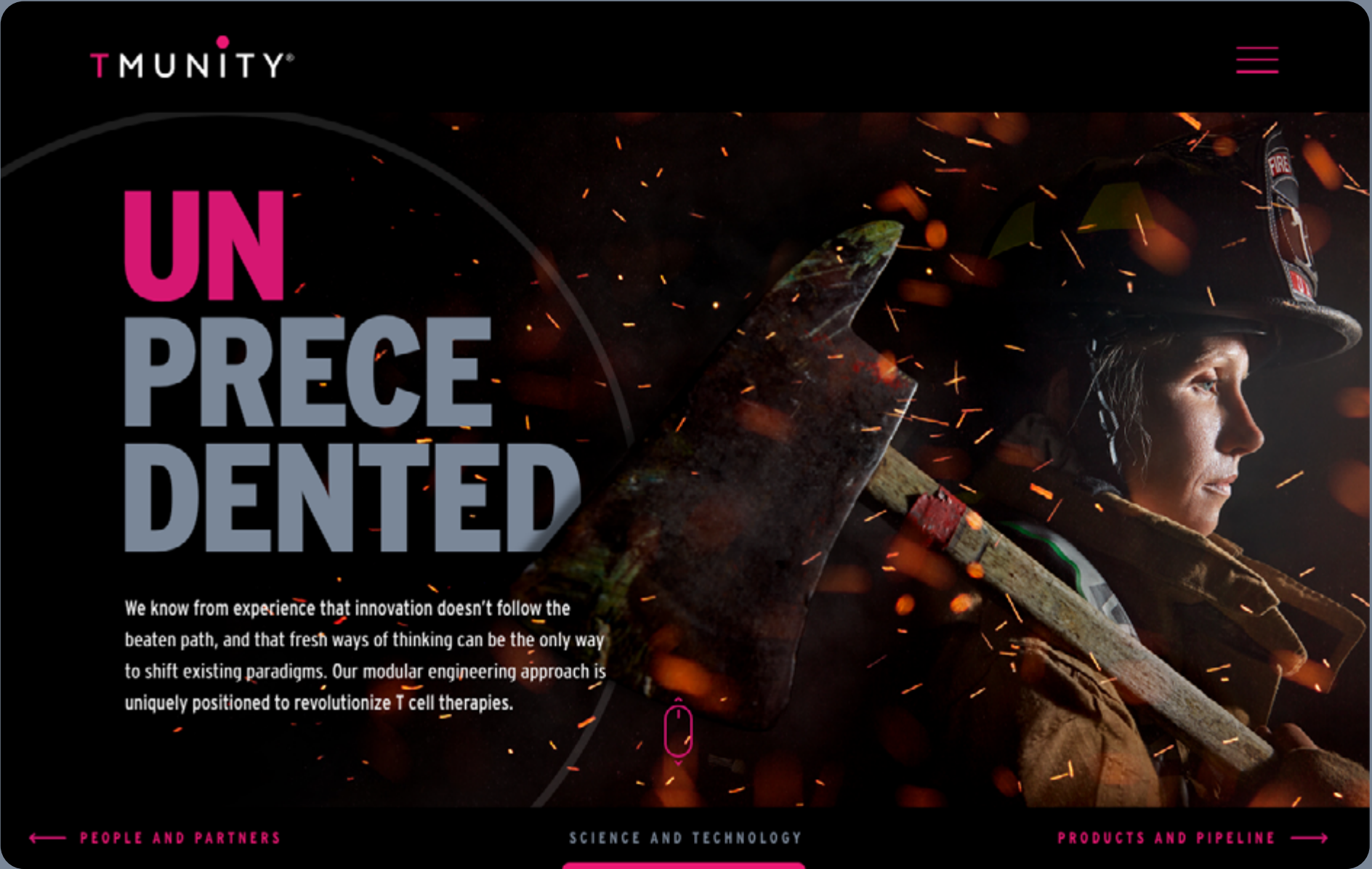
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MEDICINES FOR SOLID TUMOR PATIENTS

WHY TMUNITY?



TMUNITY



UN

WAVERING

The fight against cancer is our life's work.
We're united by a shared purpose and fueled by
our determination to maximize the therapeutic
potential of T cells for solid tumors.



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UN

RELENTING

Our serial innovation approach to CAR T
development means that each new generation
builds on previous discoveries, resulting in
iterative advancements in safety and efficacy.



TMUNITY



UN

PRECE
DENTED

We know from experience that innovation
doesn't follow the beaten path, and that
fresh ways of thinking can be the only way
to shift existing paradigms. Our modular
engineering approach is uniquely
positioned to revolutionize T cell therapies.



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CUSTOMIZABLE CAR T VECTORS

Sed ut perspiciatis unde omnis iste natus error sit voluptatem accusantium doloremque

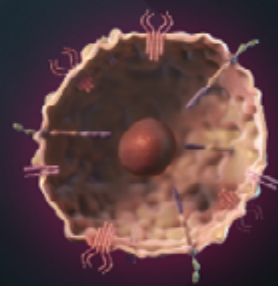
EXPLORE



ENGINEERED T CELLS

Sed ut perspiciatis unde omnis iste natus error sit voluptatem accusantium

EXPLORE



ANTIGEN RECOGNITION

Trinity's proprietary binders recognize a range of targets on solid tumors. Multiple binders can be deployed within a cell to deal with variable solid tumor antigen expression.

PSMA, PSMA	Prostate
TnMUC-1	Pancreatic, non-small cell lung, triple-negative breast, ovarian
MESO	Lung, ovarian, pancreatic, mesothelioma
FRa	Ovarian, lung, breast, cervix, endometrial, kidney, bladder
GPC2	Neuroblastoma, small cell lung, neuroendocrine tumors
EGFR	Glioblastoma
IL13Ra2	Glioblastoma, pancreatic
FAP	Prostate, lung, pancreatic, breast, ovarian, oral, gastric, colon, glioblastoma, esophageal, mesothelioma

1

2

CO-STIMULATORY DOMAINS

3

ARMORS, ENHANCERS, SWITCHES

ANTIGEN RECOGNITION



CO-STIMULATORY DOMAINS

Modules dial up or down the "engine" of T cell activation, with the potential to create CAR Ts with the optimal profile for efficacy, safety, and persistence needed to eradicate a variety of solid tumors.

CD2
4-1BB
ICOS
CD27

1

ANTIGEN RECOGNITION

2

3

ARMORS, ENHANCERS, SWITCHES

CO-STIMULATORY DOMAINS



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SERIAL INNOVATION FEEDS OUR PIPELINE

Iterative learnings from our clinical programs inform the creation of our next generation products.

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SCIENTIFIC FOUNDERS



CARL H. JUNE, MD
SCIENTIFIC FOUNDER

VIEW BIO

VIEW MANAGEMENT

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OUR CURRENT PIPELINE

A diversified portfolio of validated and novel targets in cross clinical development addressing the needs of a range of solid tumors.

1 PSMA
TmPSMA-01: PSMA 41bbz + dnTGF-βRII

TUMOR TYPE:
Prostate Cancer

PRECLINICAL PHASE 1 PHASE 2/3

EXPLORE

2 TnMUC1 Positive
TmTnMUC1-01: TnMUC1 CD2z

TUMOR TYPE:
Lung, Pancreatic, Triple-Negative Breast, and Ovarian Cancer

PRECLINICAL PHASE 1 PHASE 2/3

EXPLORE

3 Mesothelin Positive
TmMSLN-01: MSLN 41bbz

TUMOR TYPE:
Epithelial Mesothelioma, Lung and Ovarian Cancer

PRECLINICAL PHASE 1 PHASE 2/3

EXPLORE

ARMOR / SWITCH RECEPTOR

ARMOR

CAR

T CELL RECEPTOR

MICROENVIRONMENT MODULATION

To counter the immunosuppressive tumor environment, T cells can be engineered to enhance or block cytokines and express switch receptors to modulate checkpoint signaling.

dnTGFβ-R2:

Improves CAR T proliferation and expansion potential by neutralizing TGFβ signaling.

TGFβ/IL-12R switch receptor:

Improves CAR T proliferation and expansion potential by switching the inhibitory effects of TGF-β signaling into a stimulatory IL-12R signal.

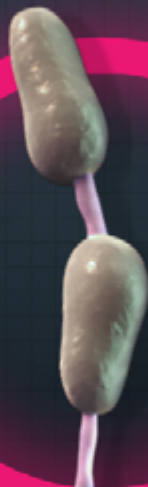
PD1/CD28 switch receptor:

Improve CAR T antitumor activity and reduce tumor-induced inhibition by leveraging the overexpression PD-L1 on tumor cells and switching its inhibitory effects into a stimulatory CD28 signal.

PROTOTYPING

[View Prototype >](#)

ANTIGEN
RECOGNITION



PSMA
0001

ARMOR,
ENHANCERS,
SWITCHES



ONTGF-BRII
0003

CO-STIMULATORY
DOMAIN



CD2
0002