

SNAP Biosciences

...

Ushering In the **Next Generation** of CAR-Based
Therapeutics

Company Overview
2025

Disclosures and Disclaimers

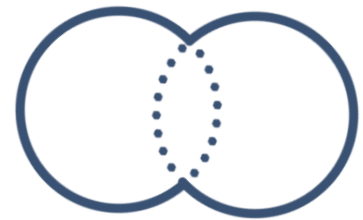
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Company Overview

At SNAP Biosciences, we are developing a “universal” CAR platform called SNAP-CAR that is designed to overcome several limitations of current CAR-based therapeutics. With our SNAP-CAR NK cells we aim to **usher in the next generation of cell therapy.**

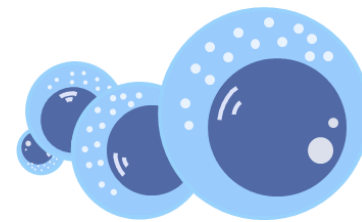
SNAP-CAR Technology



SNAP-CAR is a highly modular CAR-based platform that, instead of binding directly to target antigens, binds to targeting antibodies that direct SNAP-CAR effector cells to target cells.

+

Allogeneic NK Cells



We utilize a proprietary pooled-donor umbilical cord stem cell platform that enables the production of safe, and effective, truly “off-the-shelf” NK cell therapies.

=

Revolutionary Clinical Potential



By combining SNAP-CAR with our truly off-the-shelf NK cells, we can easily and efficiently treat a wide array of diseases including solid and liquid cancers, as well as autoimmune disorders.

The **Future of Cell Therapy** Is Bright

Cell therapies have been proven to **offer curative potential for cancers** and other diseases, often outperforming traditional therapies.

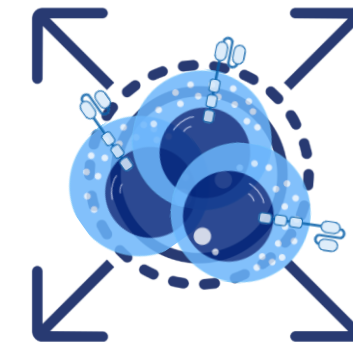
2 key indicators of a strong future for cell therapy

Strong market growth



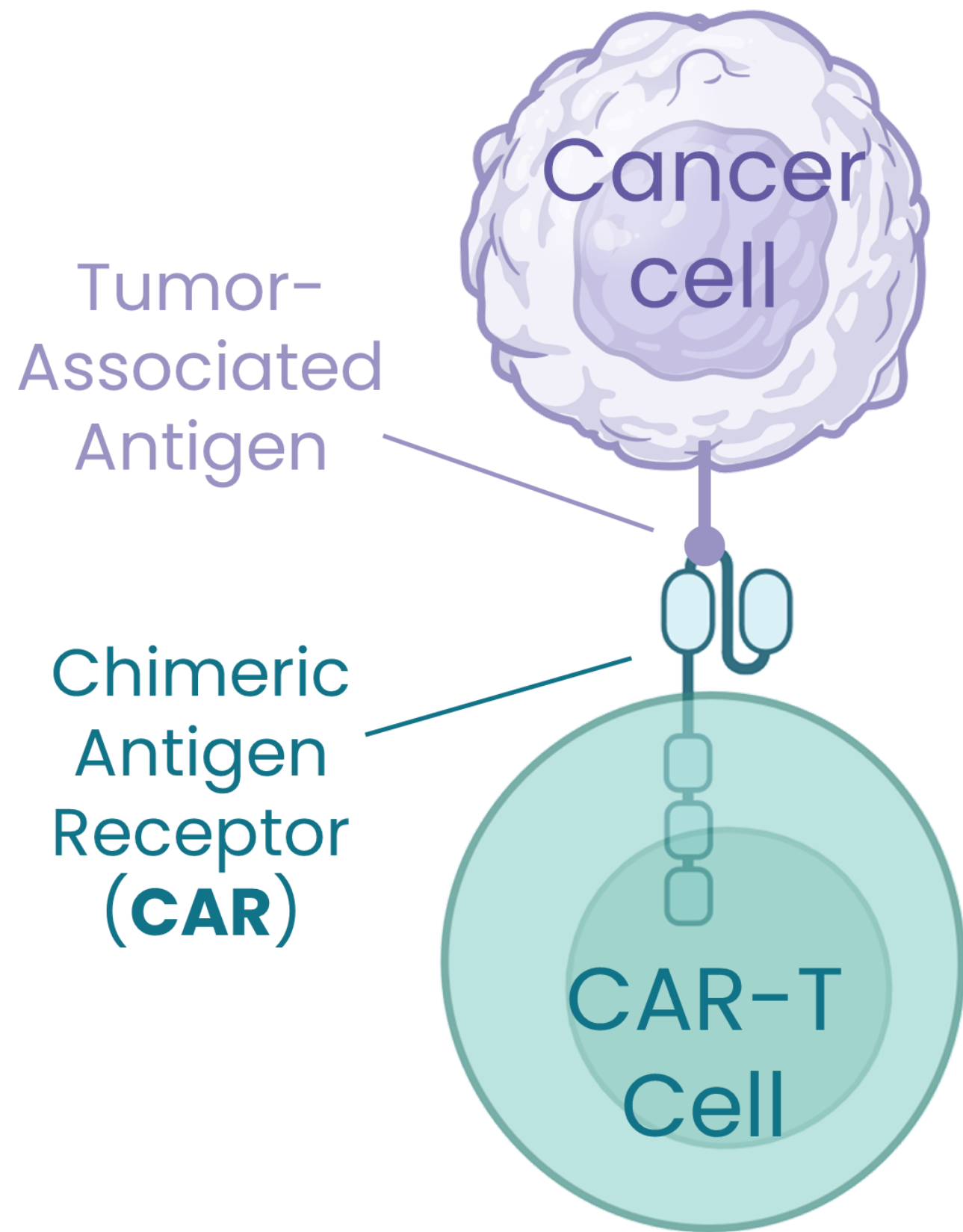
The global cell therapy market is projected to grow at a **CAGR of 23%, reaching over \$37 billion by 2033**, with CAR-based and NK-based therapies playing a prominent role¹.

Rapidly Expanding Indications



Traditionally successful in treating hematologic malignancies, cell therapies are now advancing into **solid tumors and autoimmune diseases**.

Prominent Therapeutic: **CAR-Based Cell Therapies**



CAR-Based therapeutics have a simple mechanism of action:

- 1 Design CAR to recognize a molecule on the surface of cancer cells.
- 2 Express CAR in highly cytotoxic immune cell like a T cell or an NK cell
- 3 CAR directs immune cell to find and kill cancer cells

There are currently 7 FDA approved CAR-T cell products targeting either CD19 (5/7) or BCMA (2/7) for the treatment of B cell malignancies.

Key Limitations that **Restrict CAR-T Market**

Despite their popularity and success, CAR-based therapies suffer from several limitations that restrict their clinical use and therapeutic impact.

①

Rigid Antigen Targeting

Traditional CARs “hard-code” antigen specificity into the receptor, **requiring entirely new designs** to target different antigens.

②

Toxicities

CAR-T cell-based therapies are associated with **severe toxicities** including cytokine release syndrome and neurotoxicity.

③

Time & Costs

Treatments are highly personalized taking up to 6 weeks to receive infusions and costing upwards of **\$350,000–\$500,000 per patient**.

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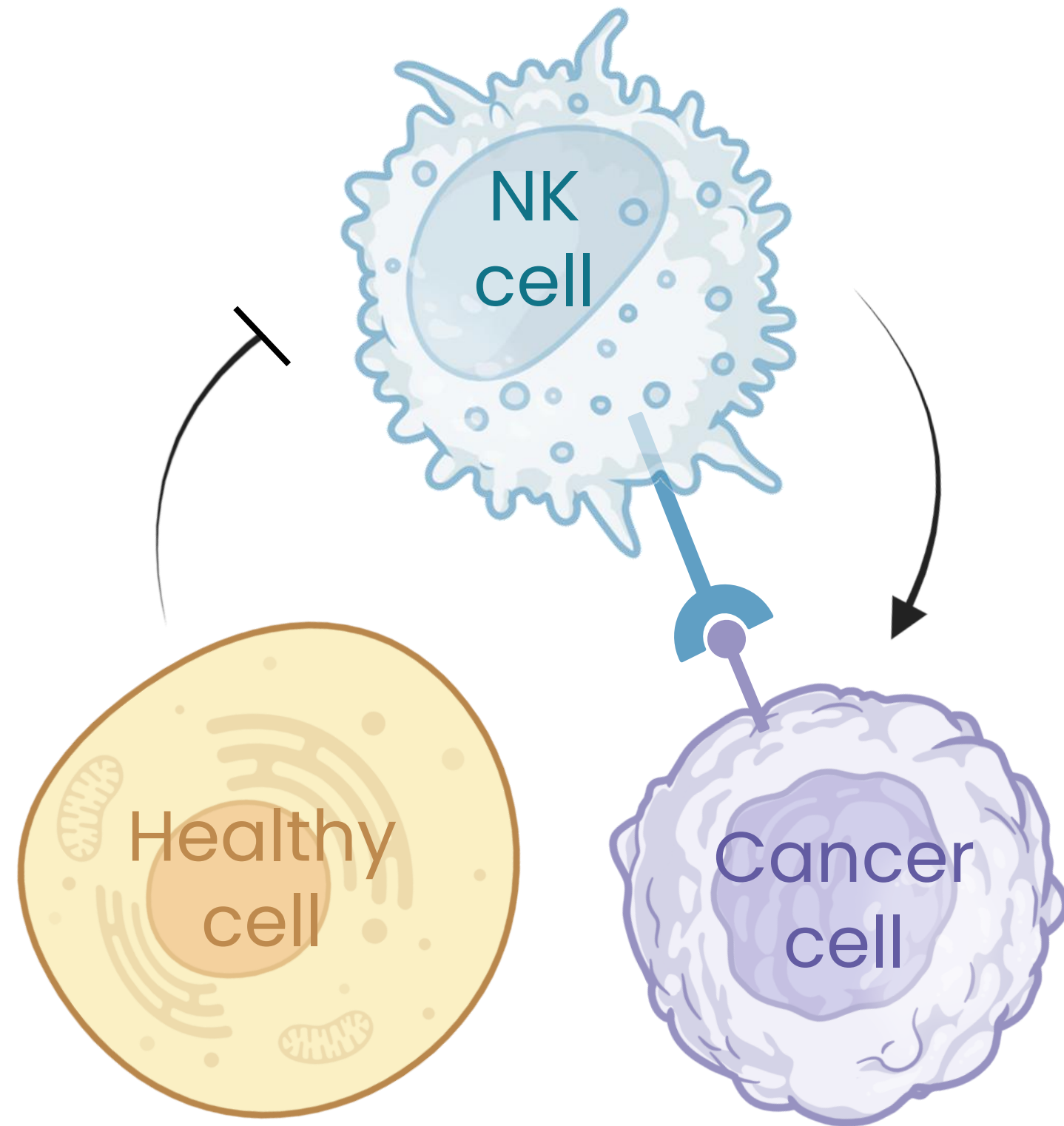
Our SNAP-CAR NK Cell platform
is designed to **combat each of
these limitations**

① Rigid Antigen Targeting
Traditional CARs “hard-code” antigen specificity into the receptor, requiring entirely new designs to target different antigens.

② Toxicities
CAR-T cell-based therapies cause toxicities including cytokine release syndrome and neurotoxicity.

③ Time & Costs
Treatments are highly personalized taking up to 6 weeks to receive infusions and costing upwards of \$350,000–\$500,000 per patient.

NK Cells Offer Distinct Advantages Over T Cells



Cytotoxic innate cells with powerful anti-cancer capabilities

- Recognize cancer cells through a **variety of mechanisms**.
- Release cytotoxic substances and kill target cells efficiently.
- Trigger a broader anti-cancer response, **activating nearby immune cells**.

More convenient “off-the-shelf” cell option

- Do not attack healthy cells unlike T cells – **safe**.
- Can therefore be used without any genetic modification – **quick and cost-effective**.

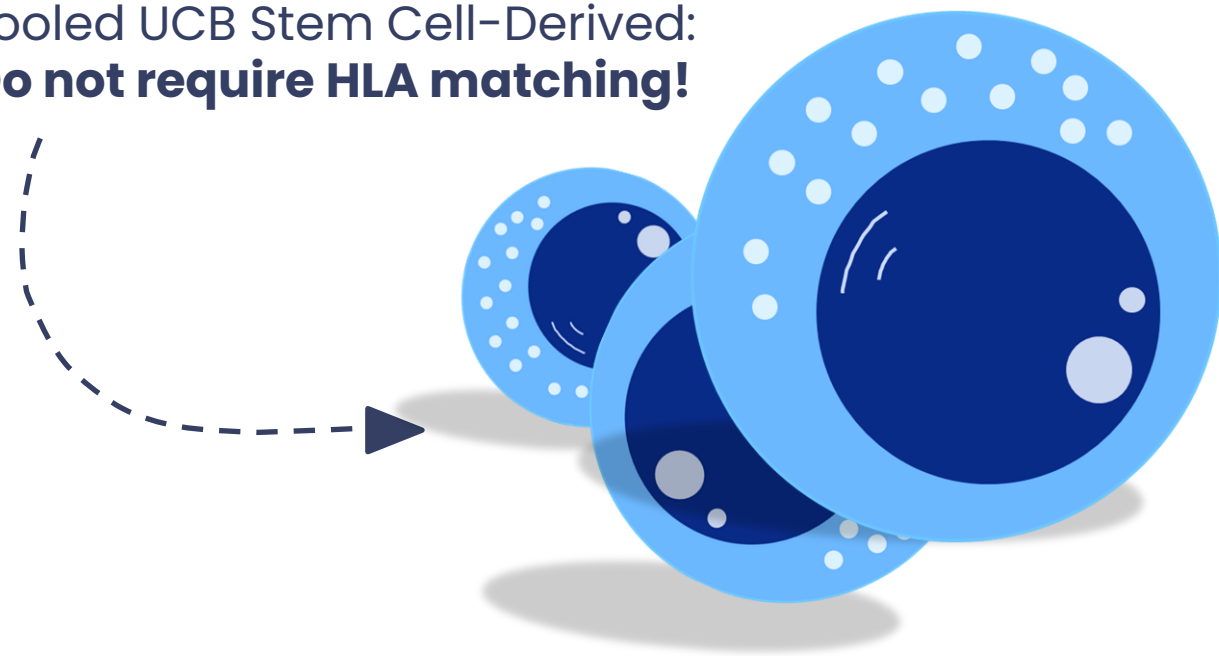
3 Key Components of SNAP-CAR Technology

1

Universal, Truly “Off-The-Shelf” NK Cells

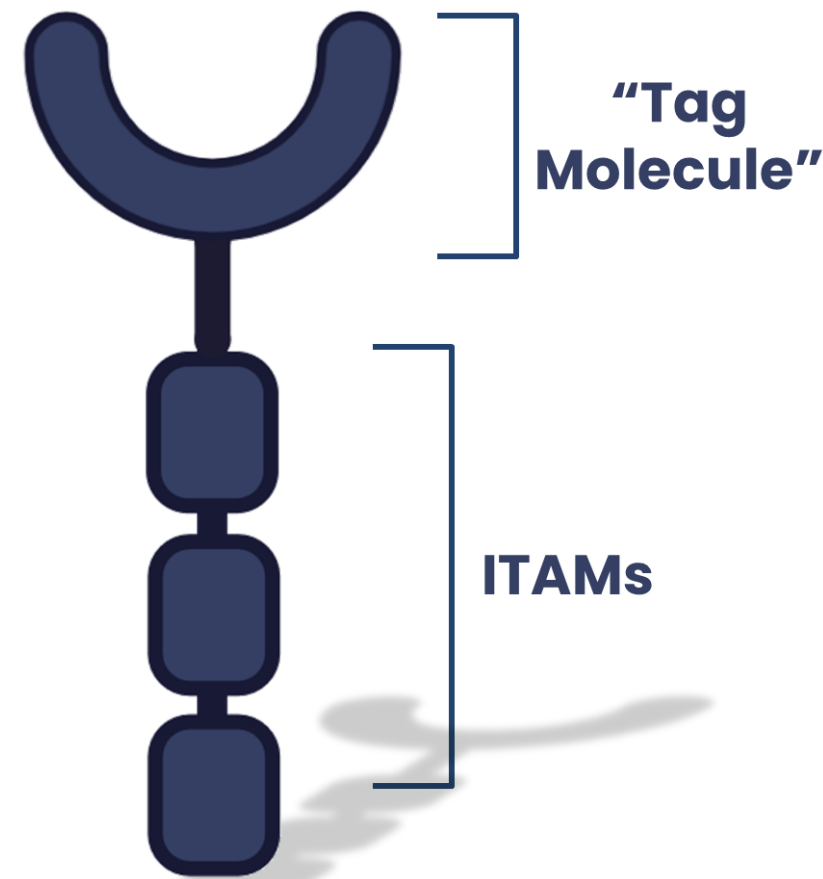
EFFECTIVE STANDALONE THERAPY

Pooled UCB Stem Cell-Derived:
Do not require HLA matching!



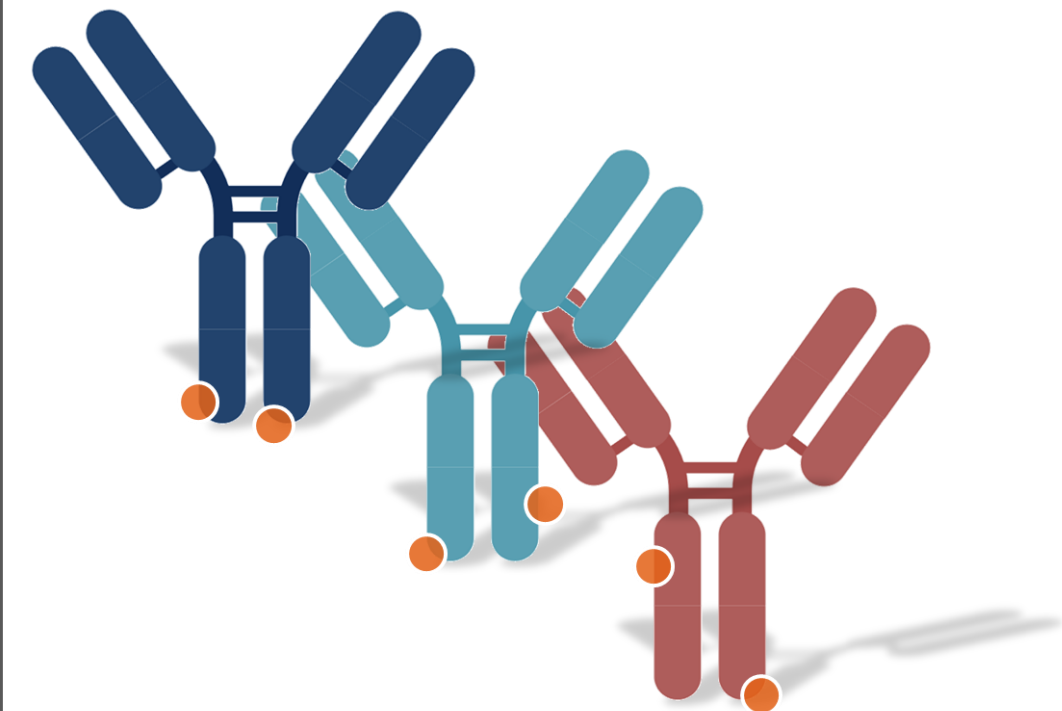
2

SNAP-CAR Receptor



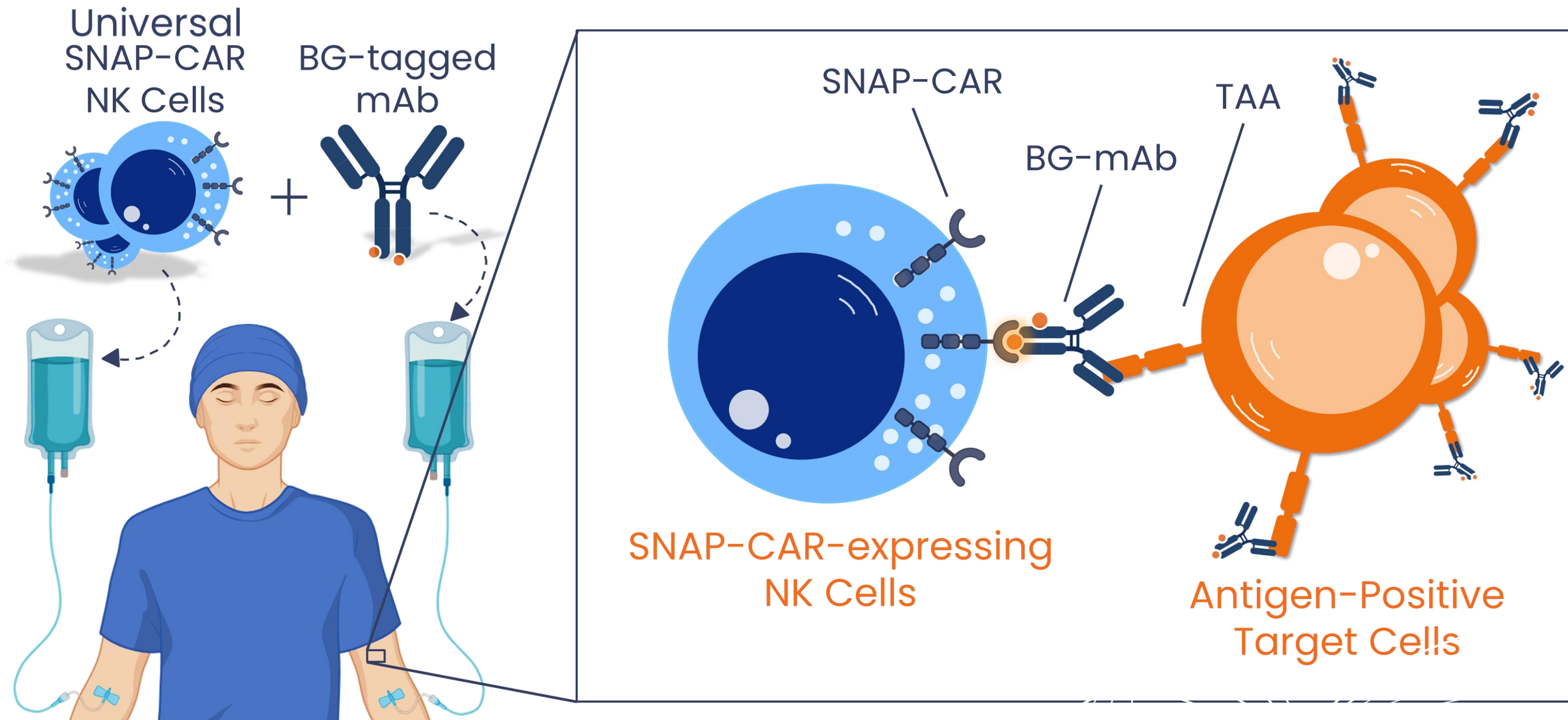
3

BG-Tagged FDA-Approved mAbs



Mechanism of Action

SNAP-CAR-NK Cells are administered in combination with BG-tagged mAbs. The two components cooperate to eliminate target cells.



Our SNAP-CAR NK Platform **Overcomes Prominent Limitations**

Our universal SNAP-CAR NK cell therapy is designed to overcome several of the key challenges facing CAR-based therapy field.

①

More Control Over Toxicities

By combining the intrinsic safety of NK cells with the tunable activity of SNAP-CAR, **our therapy is much safer than traditional CAR-T therapies.**

②

Dynamic Antigen Targeting

SNAP-CAR enables flexible targeting by using interchangeable antibodies, **addressing antigen escape and combatting antigen-negative recurrence.**

③

Quick & Cost-Effective

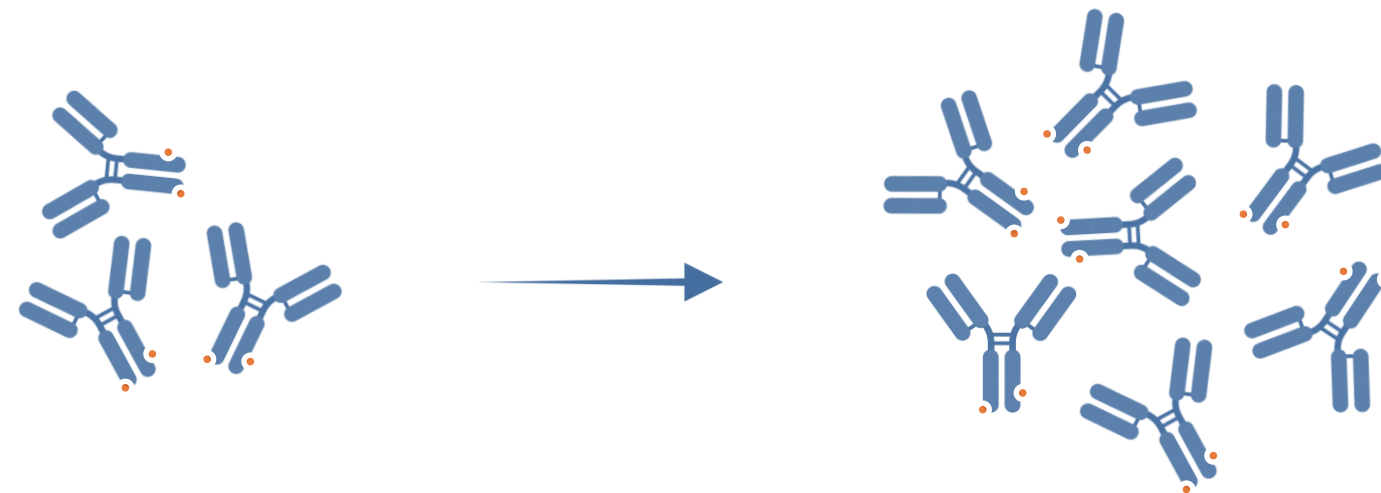
Combining the off-the-shelf nature of our cells with the dynamic nature by which we can change the target antigen of SNAP-CAR, our platform is **very cost-effective and highly scalable.**

SNAP-CAR NK Gives **More Control Over Toxicities**

1

Modular Design of SNAP-CAR gives us more control over toxicities

- Since the SNAP-CAR NK cells are reliant on the presence of the targeting antibody for activity, we can **easily tune their activity** by modulating the level of antibody.



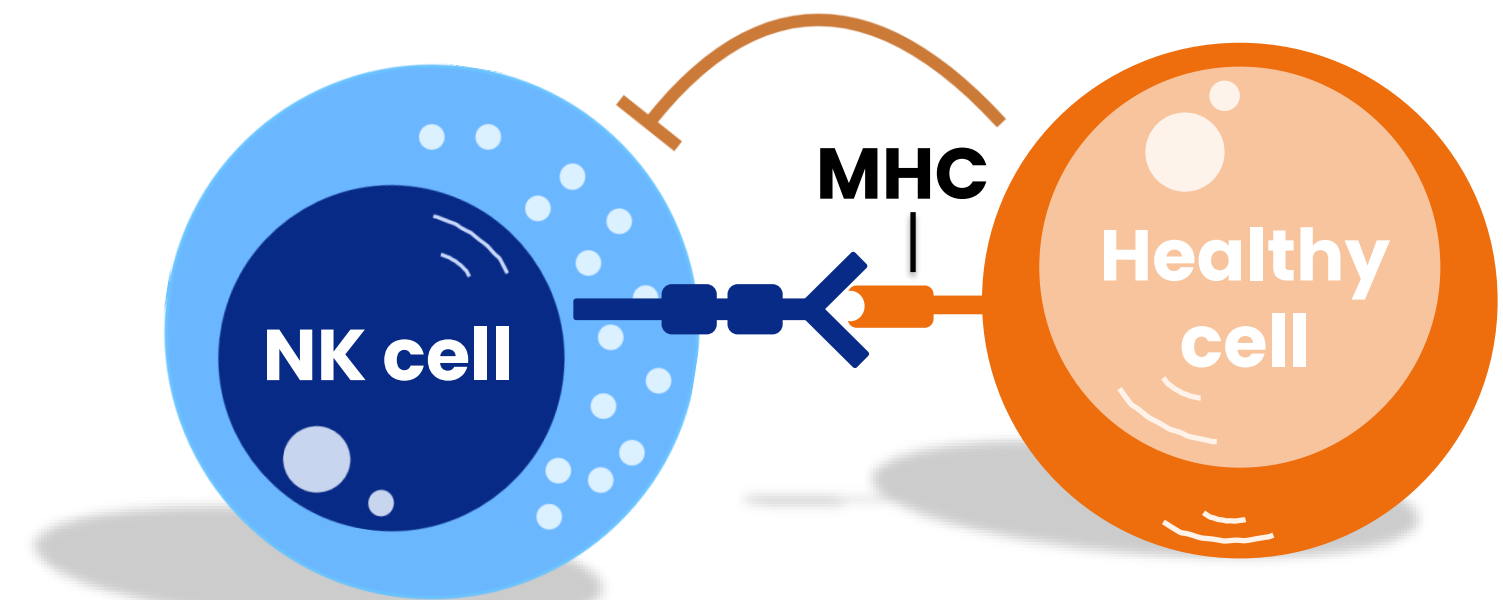
mAb concentration

LOW — **Activity of SNAP-CAR NK** — **HIGH**

2

NK Cells are a safer donor-derived cell type than T cells

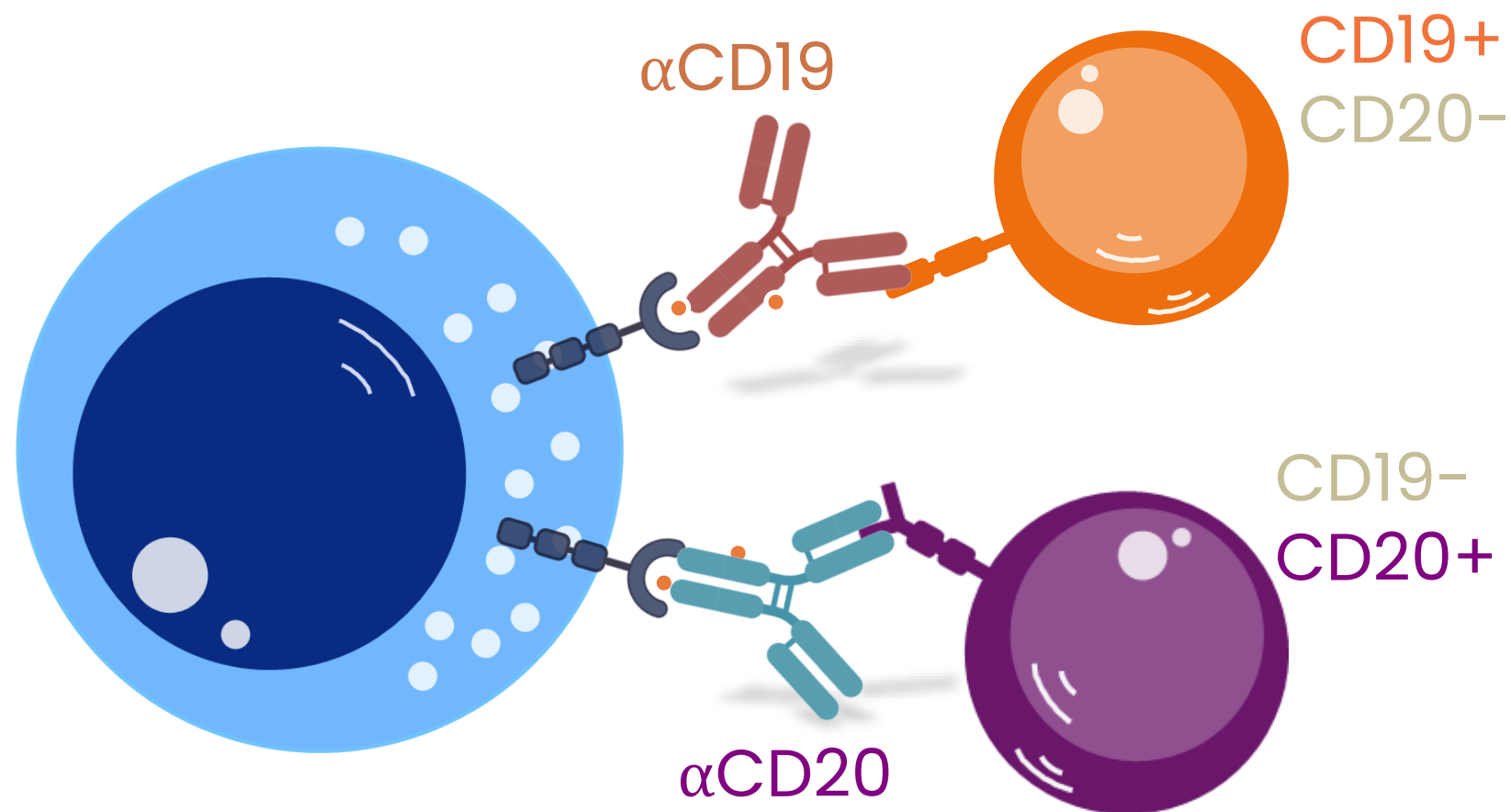
- Unlike T cells, NK cells are inhibited by MHC on healthy, normal cells. So, donor cells will not induce GvHD upon infusion.
- Also, NK cells do not expand as robustly or release the high levels of cytokines associated with T cells, making them much safer for cell therapy.



SNAP-CAR Provides 2 Mechanisms to **Overcome Antigen Escape**

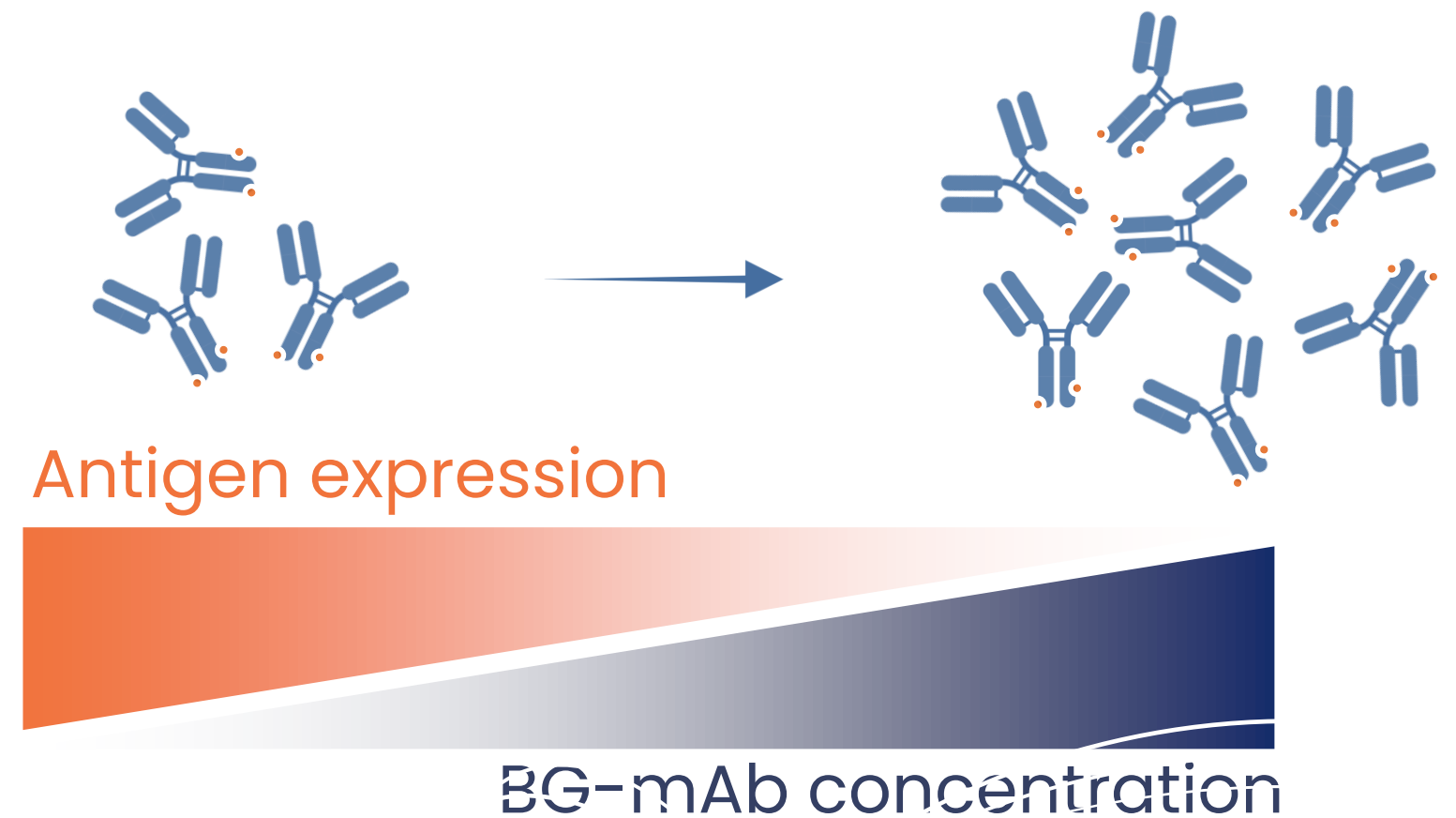
①

Multi-antigen targeting can be achieved by simply administering multiple BG-tagged mAbs, providing a simple way to overcome antigen escape.



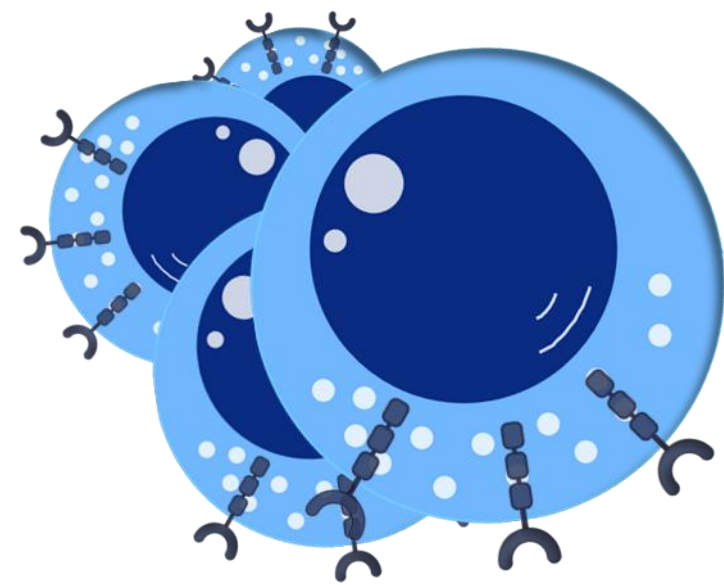
②

By administering higher concentrations of BG-tagged antibody, we can easily lower antigen threshold to eliminate antigen-low cells.



“Unlimited” Therapeutic Potential of SNAP-CAR Cell Therapy

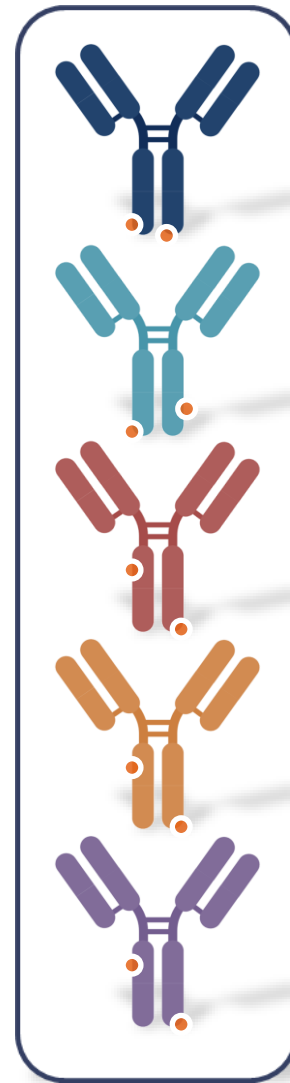
One population of
SNAP-CAR-NK cells



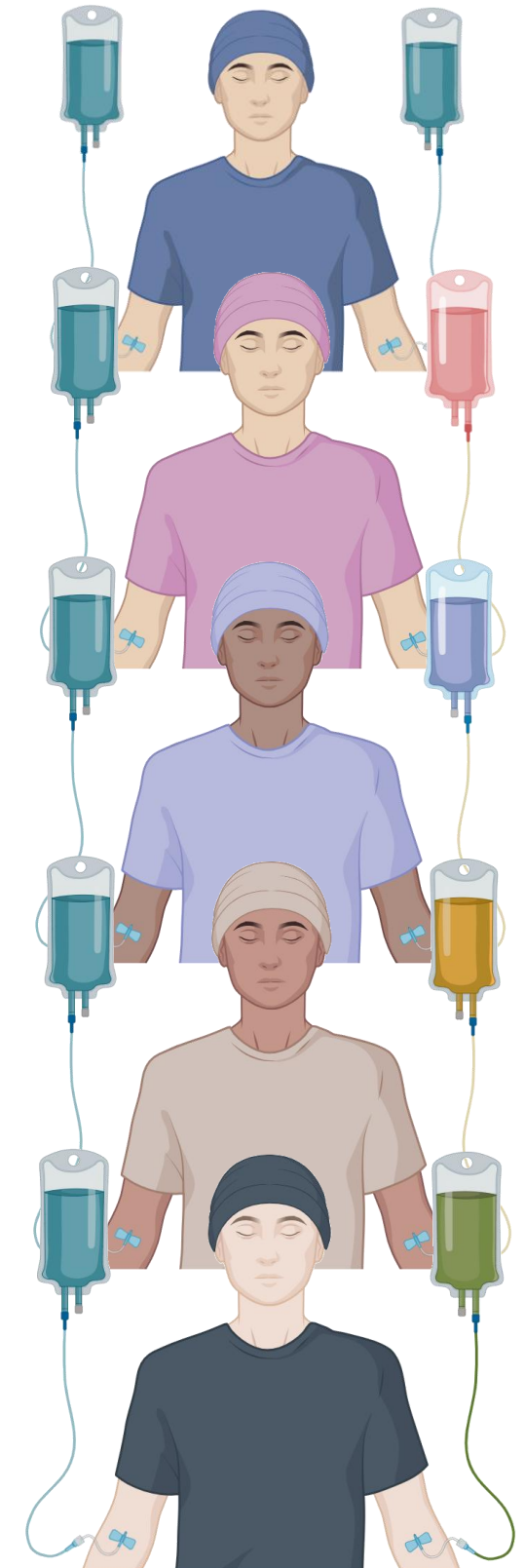
Universal NK cells:
Do not require HLA matching!

+

Library of BG-
tagged mAbs

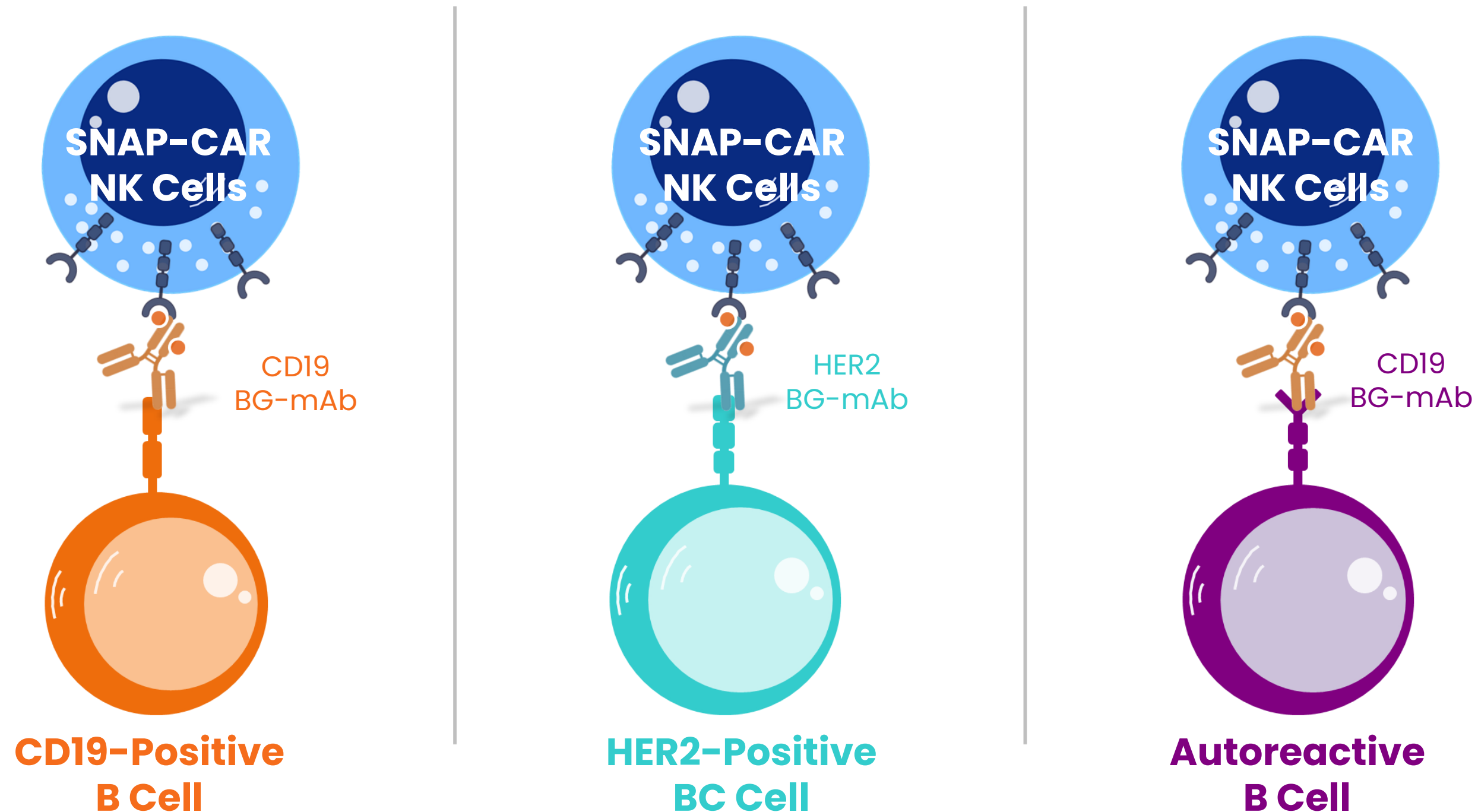


Vast array
of patients



“Unlimited” Therapeutic Potential of SNAP-CAR Cell Therapy

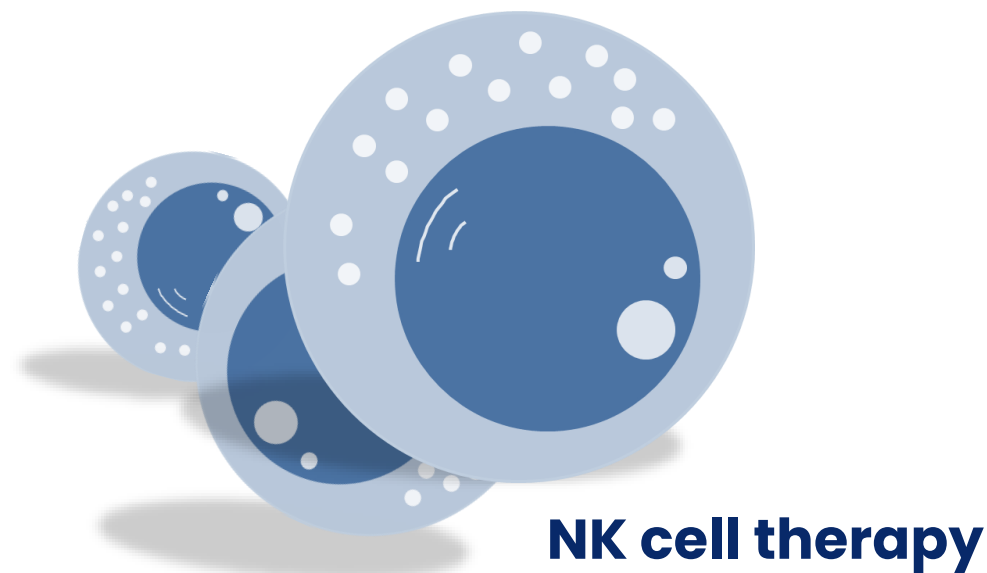
SNAP-CAR NK cells can be used to treat **both solid and liquid cancers**, as well as **autoimmunity**.



The Future of **Multi-Specific CAR-NK** Cell Therapy Is Bright

Global NK Market is growing rapidly

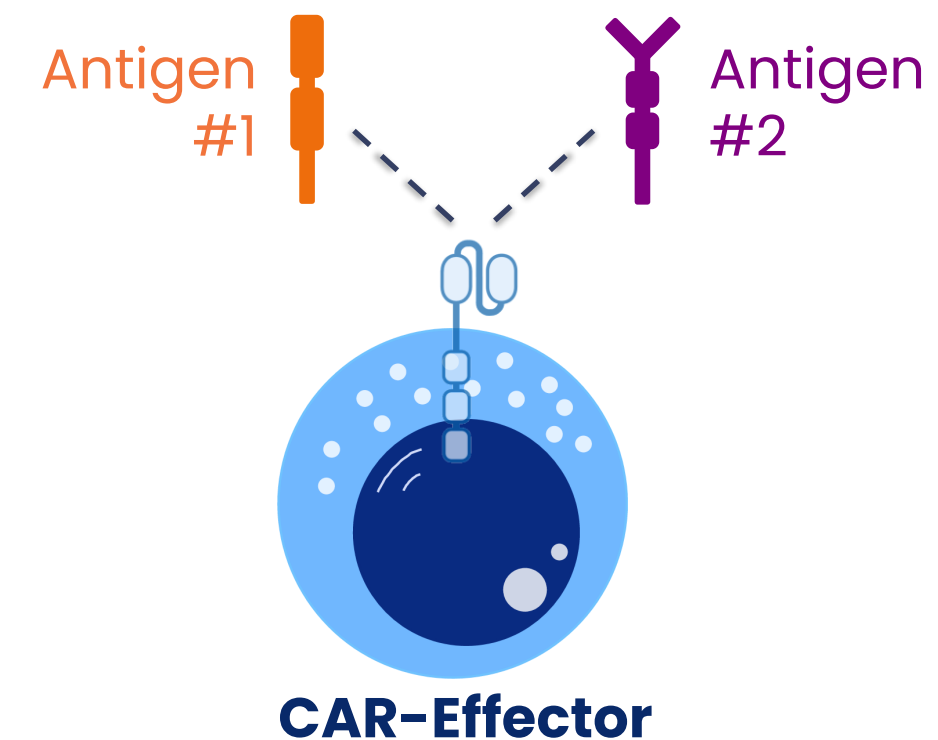
- Global NK market is projected to grow at an outstanding **44.3% CAGR** from **2022–2031** to an expected value of **\$3.16B⁺**.



⁺according to Straits research – 2024

Multi-Specific CAR platforms are gaining prominence

- Growing number of CAR trials now involve **multispecific or modular CAR platforms**, like SNAP-CAR



SNAP-CAR NK is a Highly Differentiated Approach

SNAP-CAR NK is Differentiated in this space

- Coeptis is the only universal CAR platform that has optimized its technology for use in **donor-derived NK cells**.
- SNAP-CAR receptor forms a **strong, covalent bond with the targeting antibody**, ensuring efficient transduction of signaling upon ligand binding.

Universal CAR-T Platforms		
Developer	Platform	Cell Type and Source
	SNAP-CAR	Donor-Derived NK
	ARC-T	Autologous T
	UniCAR	Donor-Derived T
	Click CAR	Unspecified
	BRIDGECAR	Autologous T
	sCAR-T	Autologous T
	TumorTag	<i>In Vivo</i> Delivery to T

Potential Clinical Opportunity: B Cell Malignancies

B Cell Malignancies

Target antigens:
CD19, CD20, CD22, CD72

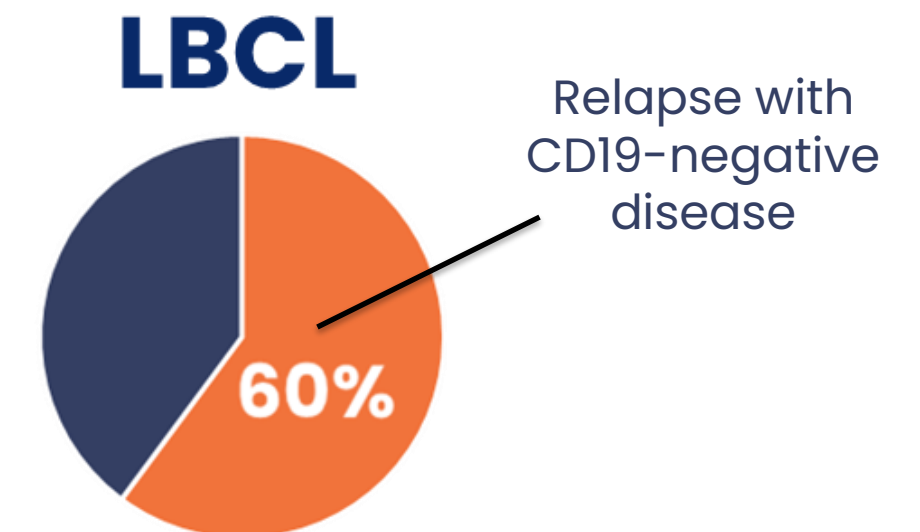
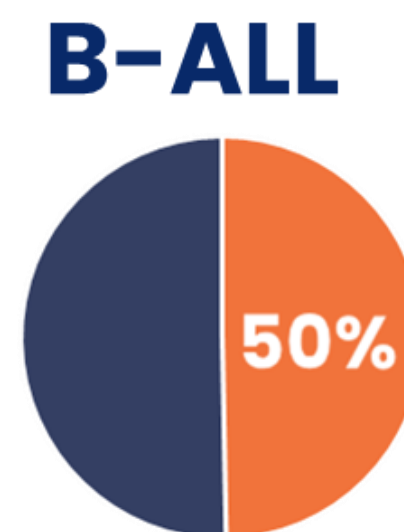
Over 80,000 Americans are diagnosed with B-cell malignancies annually*, primarily including B cell Non-Hodgkin's Lymphoma, and B-ALL

All approved CAR-T products for B cell malignancies target CD19

Multi-Antigen Approach

Targeting Strategy:
CD19 + CD20 + CD22 + CD72

Many patients relapse with antigen-negative disease. By co-administering multiple targeting antibodies, we can provide deeper, longer remissions.



*According to statistics provided by the American Cancer Society – 2024.

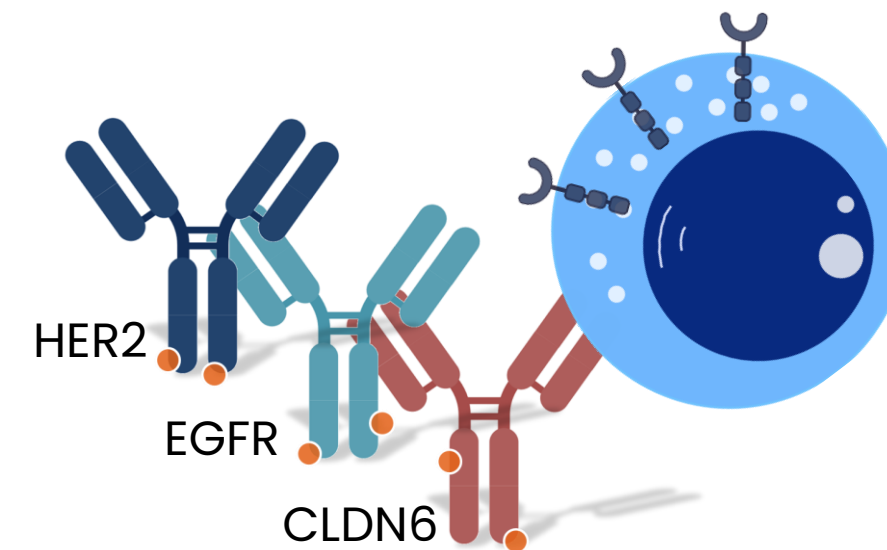
Potential Clinical Opportunity: Solid Tumors

Solid Tumors

Target antigens:
HER2, EGFR, CLDN6

Antibody therapeutics are commonly used to target multiple solid tumor indications, and several CAR-based approaches are being tested.

**no approved CAR-T products in the US for solid tumors*



- This is the ultimate goal, unlocking the full potential of SNAP-CAR NK cells.
- Massive transformative impact across oncology and beyond.

Thank You!

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