

CB-10

**A Clinic-Ready Novel Class IO therapeutic Antibody
Empowers Patients to Fight Multiple Types of Cancer**

Contact: cancurellc@canzxbio.com
www.cancurellc.com

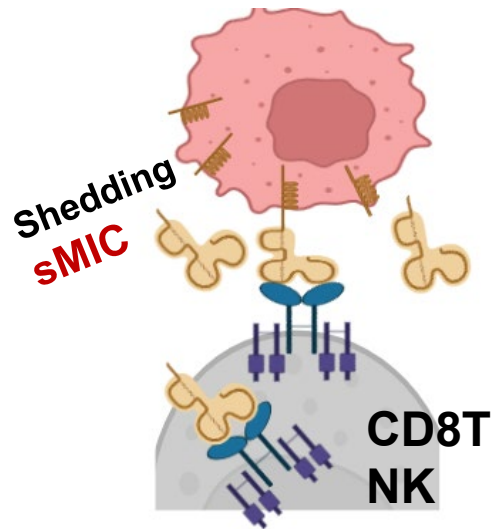
CB-10 Executive Summary



- **The problem:** Tumors overexpress MIC and produce soluble MIC (sMIC) to disable patient's immune system and ability to fight cancer
- **The solution:** CB-10 is a clinic-ready, monoclonal IgG1 antibody which captures sMIC and converts it into an immune reviving molecule to empower patients own immune system to fight off cancer.
- CB-10 has completed preclinical development and is **clinic-ready**
 - MOA, IND-enabling safety studies completed; IND to be filed in Q2 2025
- 2 clinical trials are fully funded through **NON-diluted** Awards and academic centers to treat the 1st patient in Summer 2025
- Drug Manufacturing will be completed by April 2025
- To date, CB-10 has been developed by CanCure LLC solely through **non-dilutive** capital (Founder's and NIH grants)
- CanCure LLC is seeking an investment of > \$1.5 M to help manufacture sufficient CB-10 for expanding clinical testing
- CanCure LLC would consider options for a larger investment to enable parallel company-sponsored trials to expand patient population, speed up the program development, and more disease indications
- CB-10 outperforms its competitor preclinically
- CB-10 is the leading asset of CanCure

CB-10 Novel Mechanism of Action - One molecule, multi-Modal actions

Cancer Cell:
Produce sMIC
to suppress immunity



- Immune dysfunction
- Tumor metastasis
- Poor response to current immunotherapy

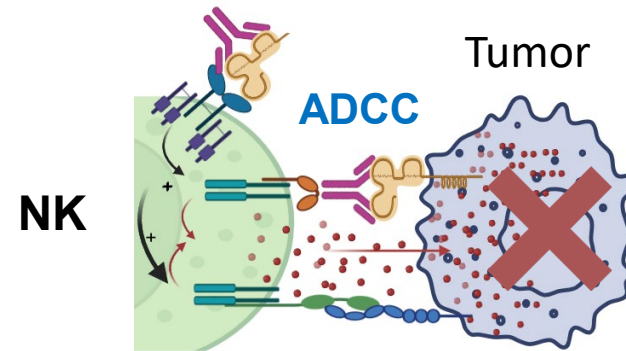
CB-10:
transforms sMIC to
an immune activator

CB-10/sMIC
complex

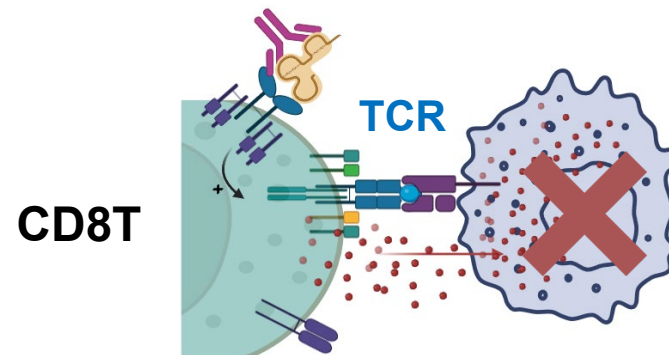


OUTCOME:

Immune Revamping (NK and CD8 T)
Tumor Elimination



- Co-stimulate NK
- Enhance “memory” or stemness
- ADCC



- Co-stimulate CD8T
- Enhance memory
- Enhanced effector function

Tumor elimination

CB-10: humanized B10G5, hIgG1

Social and Economical Impact

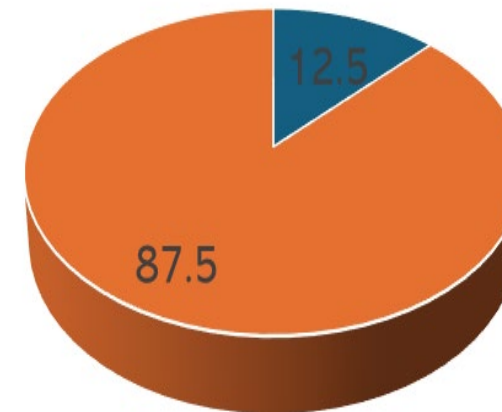
Social Impact

CB-10 could fulfill the large global unmet need for cancer treatment, prolong many millions of lives

Economic Impact

- Counting all cancer patients, only a small population (12.5%) of ALL cancer patients can benefit from current IO drugs; but the Global IO drug market is about \$138.87Bn in 2024, projected \$224.30Bn by 2030, 8.3% CAGR
- CB-10 can not only capture all the IO drug failures but also makes current IO drugs work better.
- CB-10 capture even 25% of the Global Cancer Market, ~\$276Bn annual revenue

Benefit (12.5%)
2024 Market: \$138.87Bn
2030 Market: \$224.50Bn

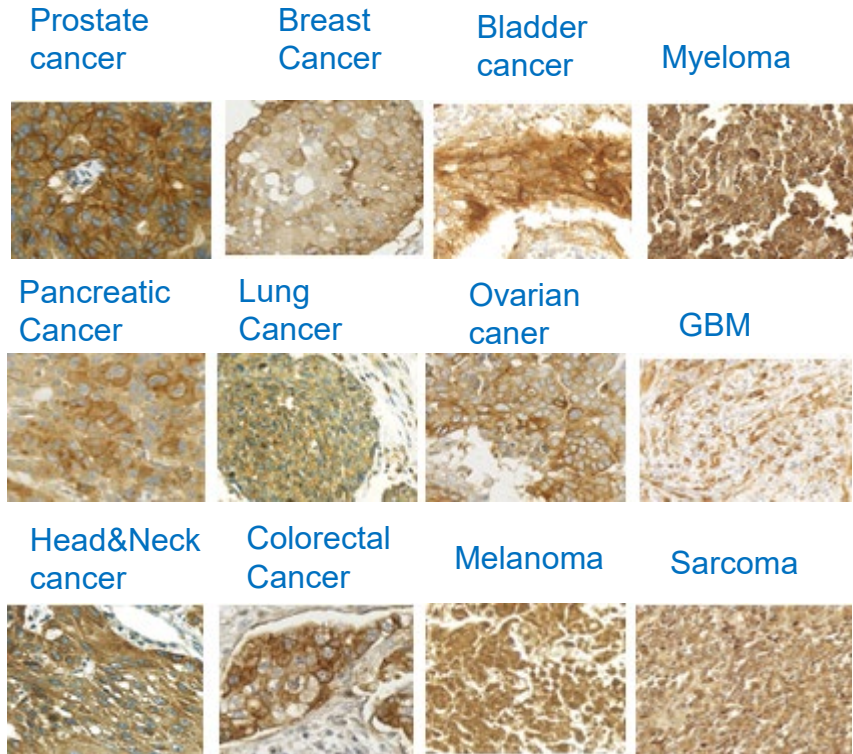


NOT Benefit,
Unmet need
87.5%

All Cancers

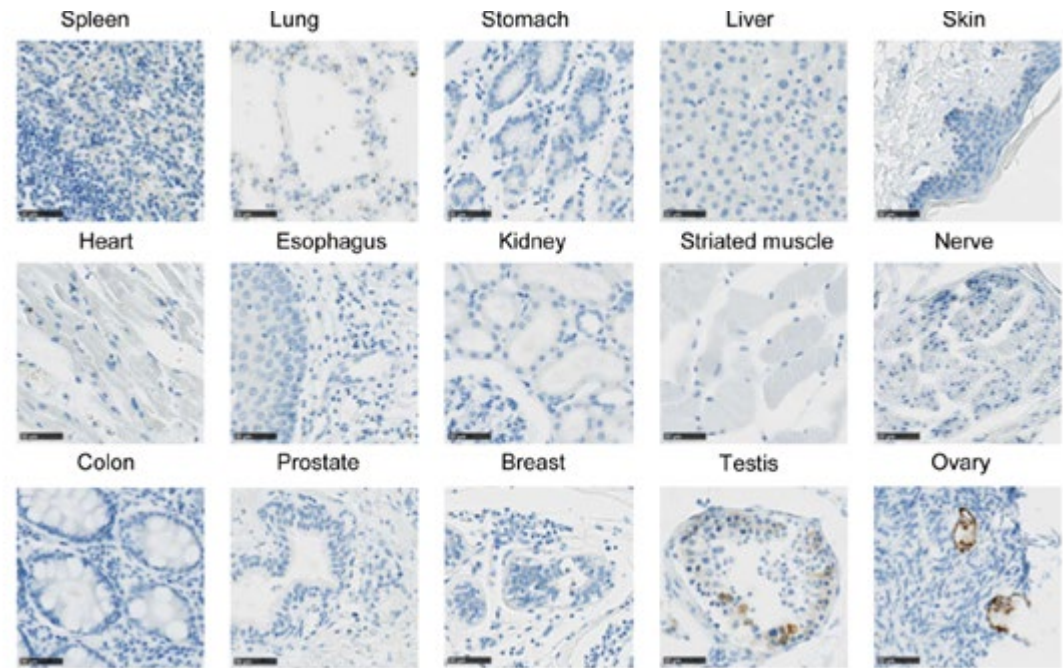
The CB-10 target MIC expression is prevalent in multiple solid tumor types with minimal presence in normal tissues

TUMOR (>90%)



Unpublished (CanCure)

NORMAL TISSUE

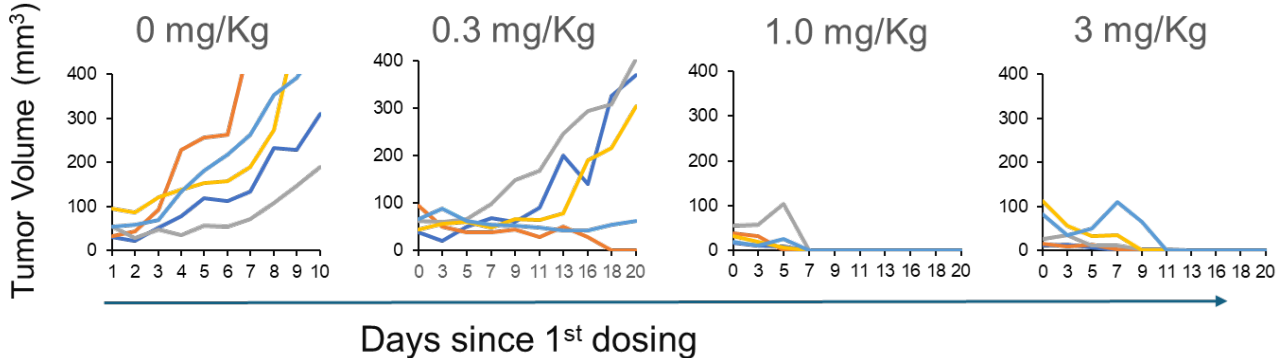


Blery M. Open Res Eur. 2021 Oct 27;1:107. PMID 35967081

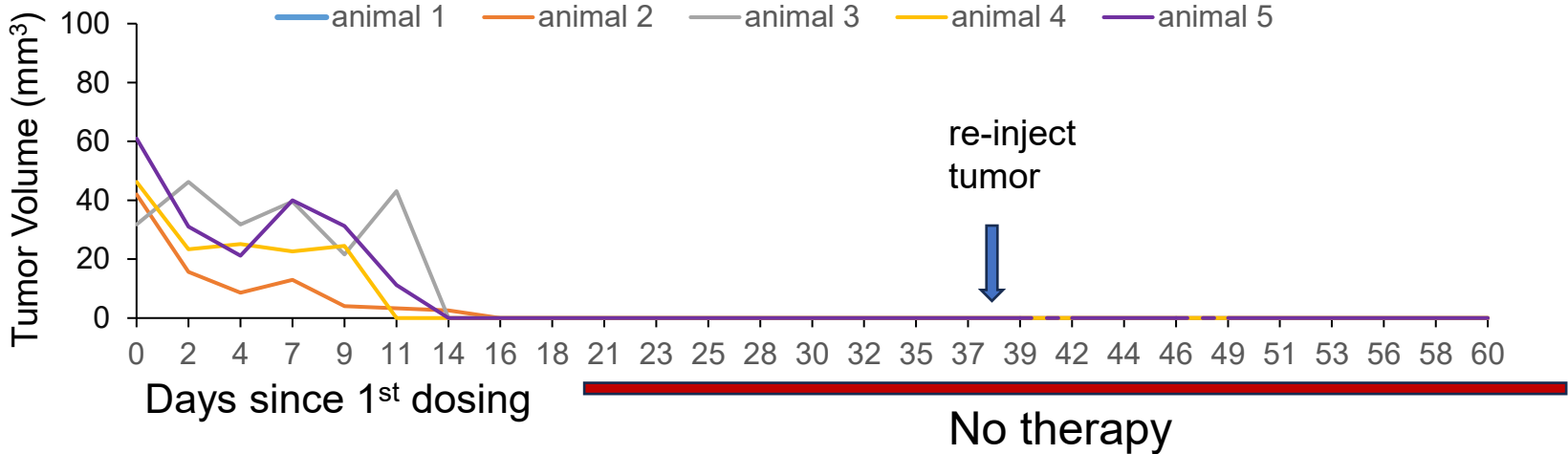
- ❖ Consistent with the lack of MIC expression in normal tissues, GLP toxicity studies demonstrated no significant pathologic findings at the highest dose level tested (100 mg/kg)

CB-10 demonstrates monotherapy activity in multiple preclinical tumor models

❖ Effective with low dose (1 mg/Kg) and outperforms CLN-619



❖ Long lasting cure and prevent tumor re-growth



CLN-619
1 mg/Kg

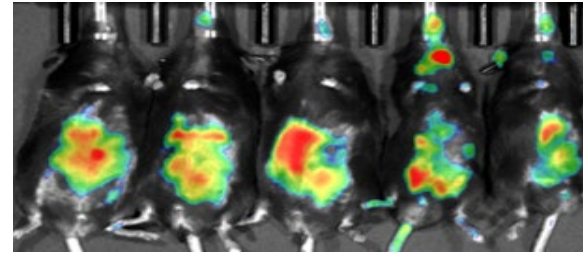
The graph shows tumor volume (mm³) over time (Days since 1st dosing) for CLN-619 at 1 mg/Kg dosage. The y-axis ranges from 0 to 400 mm³, and the x-axis ranges from 0 to 21 days. Tumor volume increases significantly for all animals, indicating that CLN-619 is less effective than CB-10 at this dosage.

CLN-619 is an IgG1 monoclonal antibody which reduces MIC shedding. The main MOA of CLN-619 is ADCC.

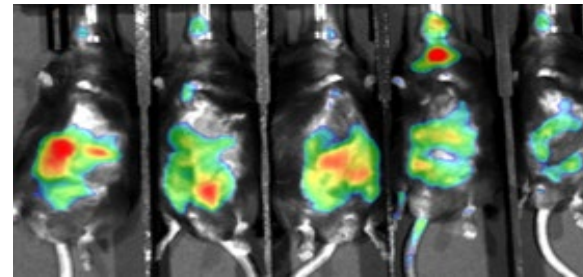
Model system: Syngeneic TRAMP-C2-MICB tumor, s.c. (Unpublished)

CB-10 monotherapy leads to complete regressions in a carcinogen-induced metastatic bladder tumor model

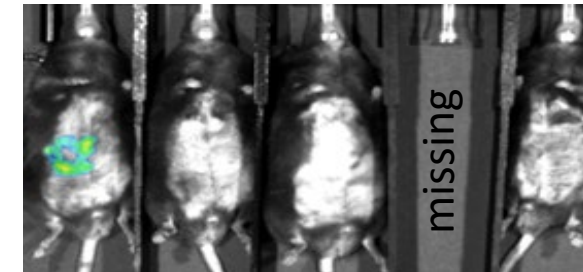
Mice (engineered*) were fed with bladder-specific carcinogen (BBN)



2 wks CB-10 therapy (i.p. twice weekly)



4 wks CB-10 therapy (i.p. twice weekly)



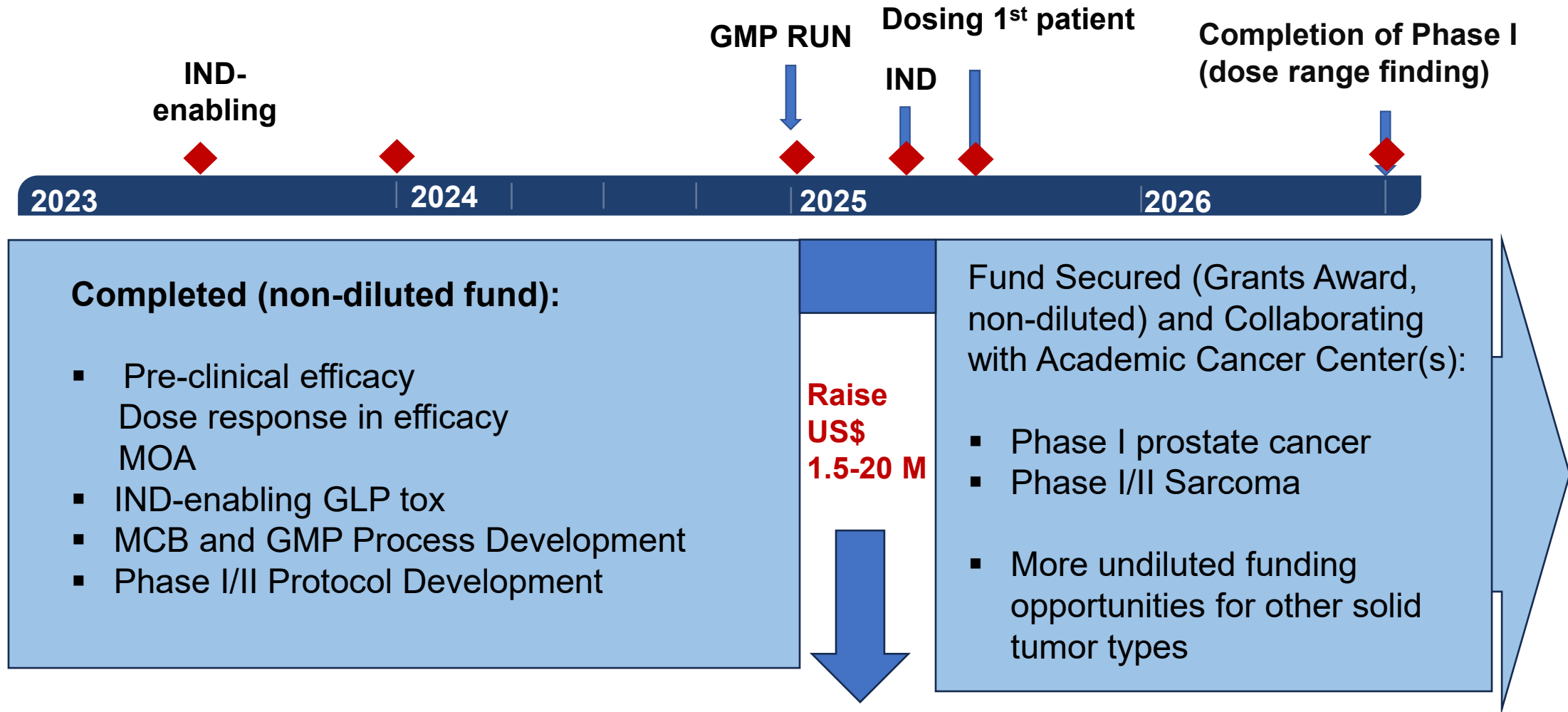
- Primary tumors were induced in the bladder
- Tumor metastasized to liver, lung, LN, and kidney

Burdens of primary tumor and metastatic tumor were reduced

Primary tumor and metastatic tumors were eradicated

***Note: MIC genes do not naturally exist in mice. Mice were engineered to have a MIC gene integrated in the genome.**

CB-10 Clinical Development Plan



Manufacturing Sufficient Clinical Drug to Treat More Patients and Expand the technology Platform and Indications

Impact and Revenue Forecast

Social Impact

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Economic Impact

- Approved IO drugs with ONLY a small population (8%) of patients can benefit, but the Global IO market is about \$138.87Bn in 2024, projected \$224.30Bn by 2030, 8.3% CAGR
- CB-10 can not only capture all the IO drug failures but also makes current IO drugs work better.
- CB-10 captures even 10% of the Global Cancer Market, ~\$140Bn annual revenue

Early-stage revenue stream: Technology out-licensing or partnership

Historical Deal Size (total in \$Bn, based on 2023 data)

- At Phase I clinical stage out-licensing: > \$2 Bn
- At Phase II clinical stage out-licensing \$2.8-22 Bn

(source: www.nature.com/biopharmdeal | December 2023 | B45)

Ask for US\$1.5 - 20M to speed up producing sufficient drugs to expand the technology platform and clinical indications