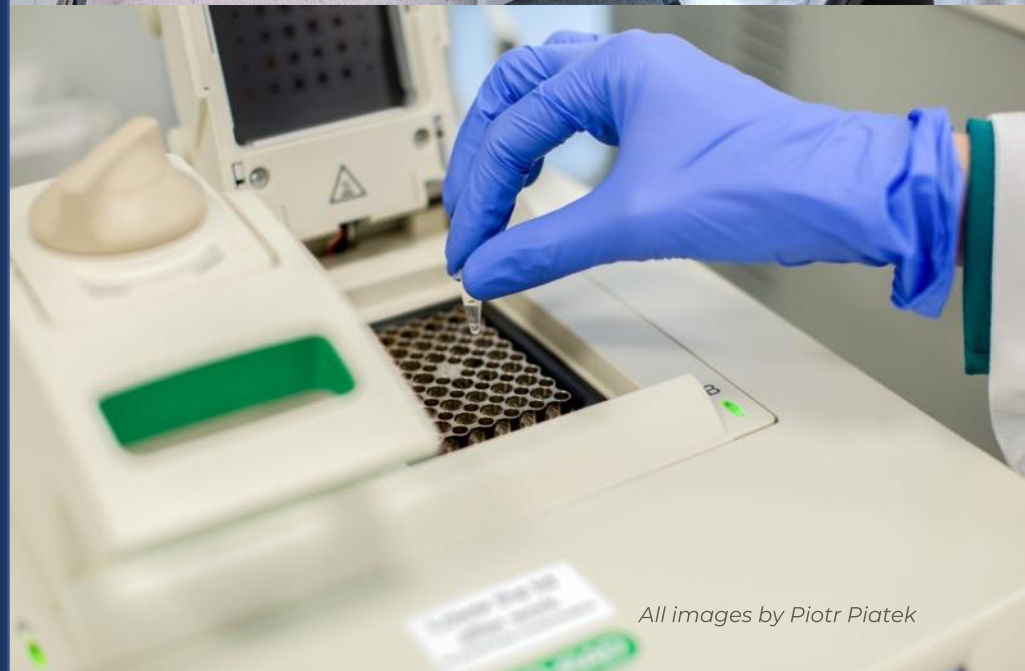




# Captor Strategy Next Steps 2023-2025



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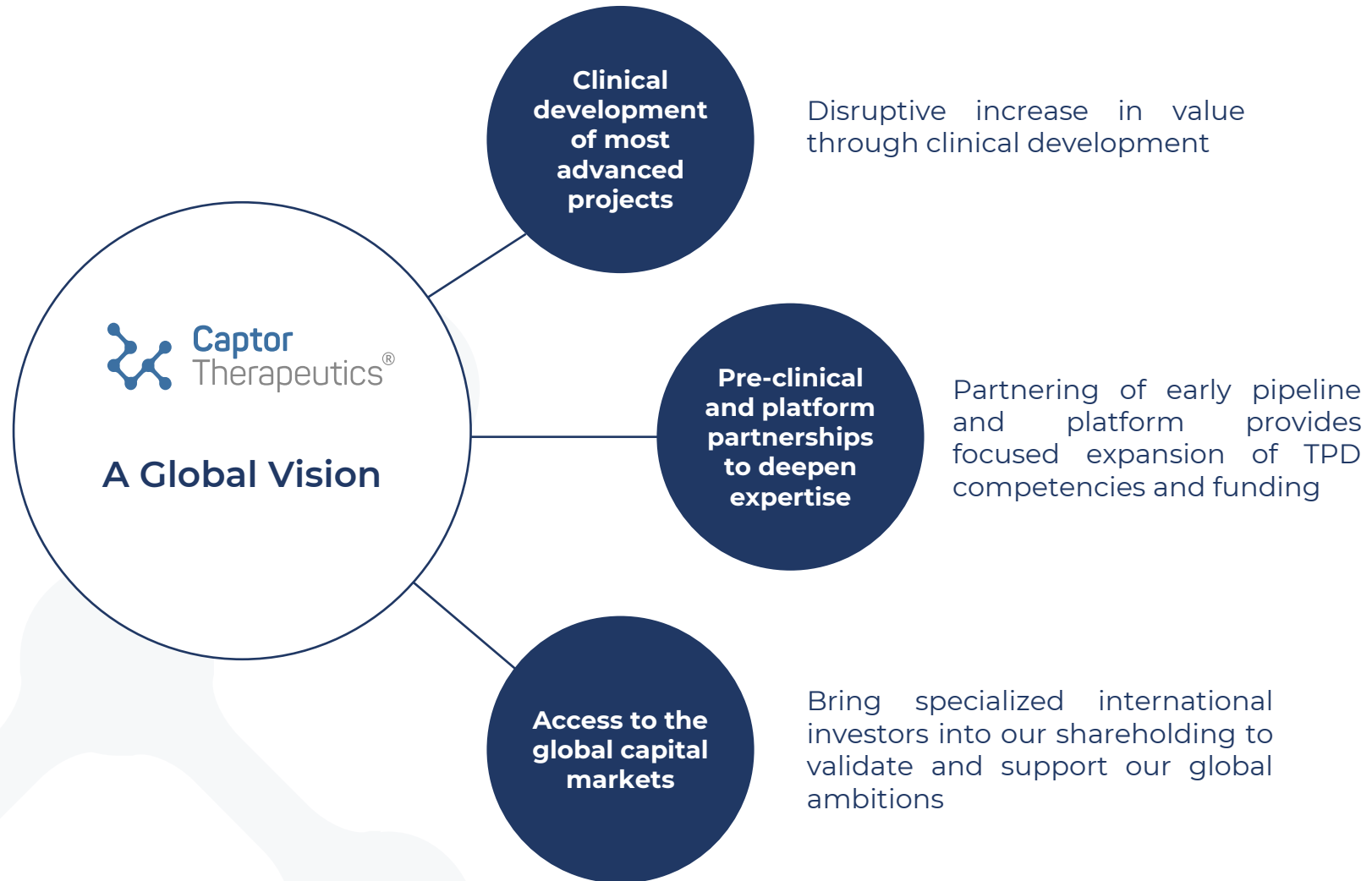
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# Towards a global TPD Company



# Roadmap to our strategic objectives

- Maintain a balanced portfolio of owned clinical and preclinical assets, while sharing development or commercial risks with partners at the optimum time for each asset.
- Develop our scientific capabilities and Company resources to maintain highest quality and deliver highest value to shareholders.



4 active pipeline projects\*

Optigrade™ Platform

Clinical trials in patients of 2 lead pipeline assets – CT-01 and CT-03

Further preclinical work on CT-02 and CT-05, with partnering or licensing at preclinical stage

- 2 new collaborative areas – Novel ligases and ADCs
- Leverage our platform for additional non-dilutive funding and validation
- Source of new early pipeline projects

# Captor's key objectives for 2023-2025

2023

**CT-01:** Phase Ia/Ib initiation in liver cancer patients

2023

**CT-02** and **CT-05:** *In vivo* proof of concept in autoimmunity

2024

**CT-03:** Phase Ia/Ib initiation in haematological cancer patients

2024

**CT-01:** Clinical readouts: safety, pharmacology, & mechanism

2024

First degrader of a new target based on **new E3 ligase**

2025

**CT-01:** Clinical readouts: combination safety, pharmacology & mechanism

2025

Lead compound of a new target based on **new E3 ligase enters pipeline**

2025

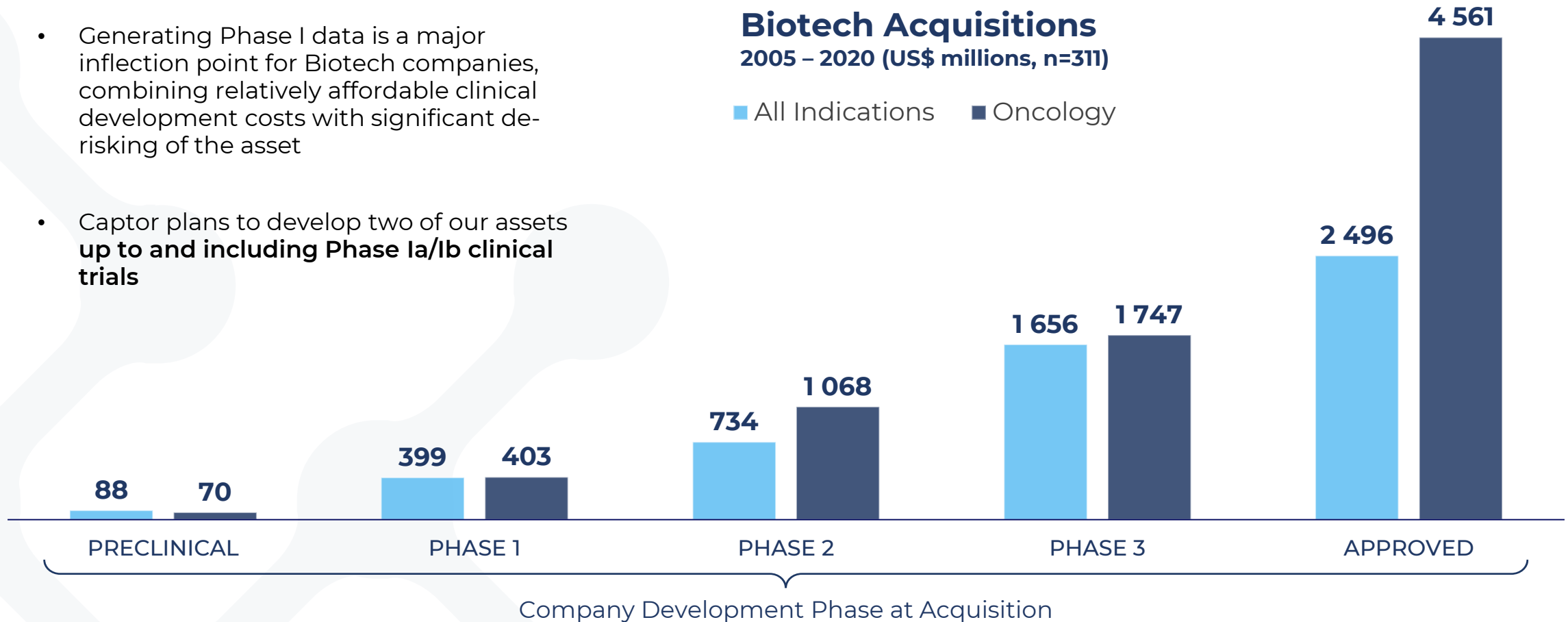
**CT-03:** Clinical readouts: safety, pharmacology & mechanism (monotherapy and combination)

# Average company value and development phase of lead asset

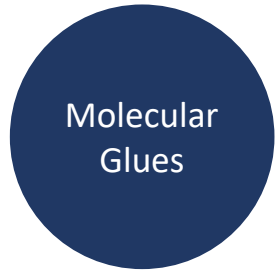
- Generating Phase I data is a major inflection point for Biotech companies, combining relatively affordable clinical development costs with significant de-risking of the asset
- Captor plans to develop two of our assets **up to and including Phase Ia/Ib clinical trials**

## Biotech Acquisitions 2005 – 2020 (US\$ millions, n=311)

■ All Indications ■ Oncology



# Clinical development – CT-01



**Project:**

CT-01

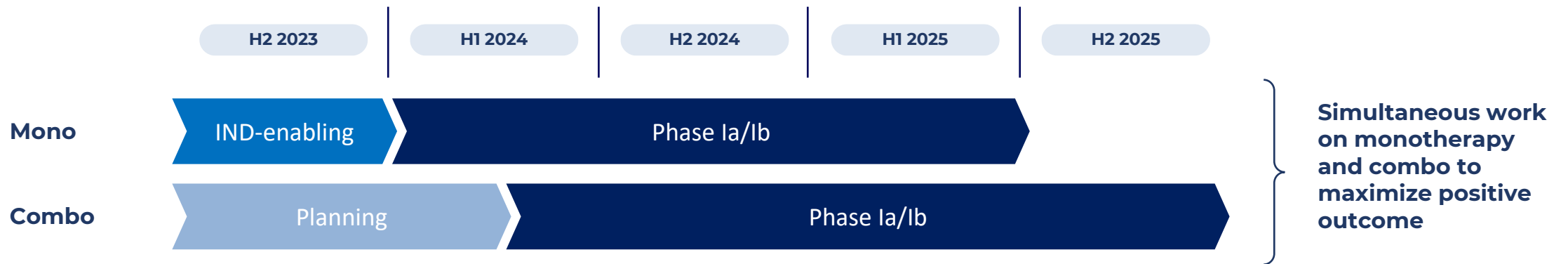
**Main indication:**

hepatocellular carcinoma

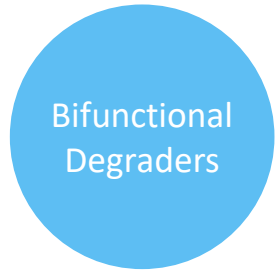
**Expected milestones:**

- IND/CTA approval allowing the initiation of clinical trials in Q3 2023
- Initiation of Phase I clinical trial in Q4 2023
- Phase Ia/Ib top-line data to be reported by the end of 2024
- Combination study data by end 2025

- Anticancer activity in different HCC models *in vitro*
- Excellent *in vivo* efficacy with oral administration as monotherapy
- Full tumour regression observed with doses of 10 and 25mg/kg



# Clinical development – CT-03



**Project:**

CT-03

**Main indications:**

blood cancers

**Expected milestones:**

- IND/CTA approval in Q3 2024
- Initiation of Phase I clinical trial in Q3/Q4 2024
- Phase Ia/Ib top-line data to be reported in 2025

- Anticancer activity in vitro in both liquid and solid tumours
- Potent and sustained MCL-1 degradation *in vivo* after single injection
- Cancer cell killing and tumour shrinkage in vivo





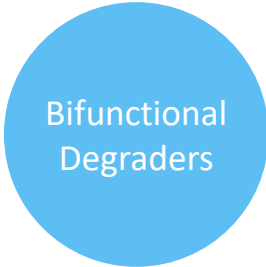
# Targeting undrugged proteins with degradation

- *In vitro* and pharmacokinetic *in vivo* data support high value in the CT-02 and CT-05 projects
- CT-02 pathway remains one of the most attractive in the pharma industry with potential multi-billion dollar deals
- Inhibitors of the CT-05 target have been of high interest but showed inadequate selectivity
- Tuned degradation of the CT-02 and CT-05 targets superior activity, selectivity and safety



Molecular  
Glues

**CT-02**  
**Autoimmunity, Oncology,**  
**Metabolism, CNS**



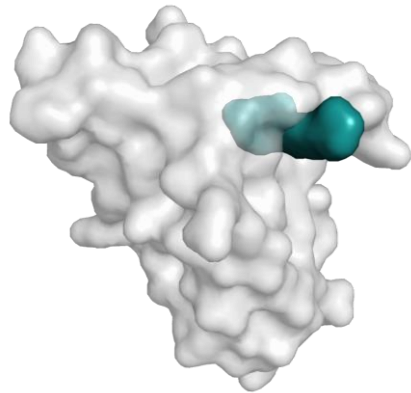
Bifunctional  
Degraders

**CT-05**  
**Autoimmunity, Transplantation,**  
**Oncology, Metabolism**

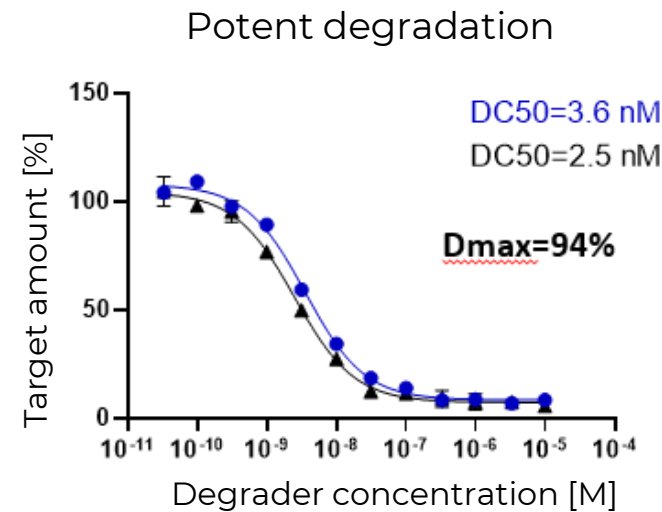
# High therapeutic potential of the CT-02 molecular glues

CT-02 -  
Molecular  
Glue

Autoimmunity, CNS,  
Metabolism, Oncology



Atomic structure of E3 ligase and a potent degrader



## 2023 expected milestones

Demonstration of the *in vivo* efficacy in inflammation model

Assessment of potential to cross blood-brain barrier for the **treatment of neurodegenerative diseases**

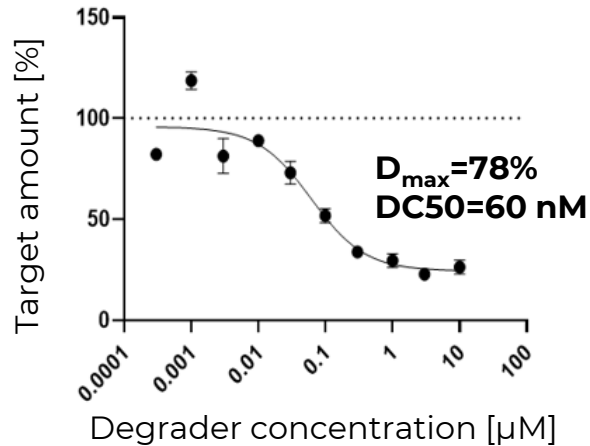
Commercialization of the entire programme or separated by therapeutic area

# Targeting an inadequately drugged protein by degradation

Bifunctional  
Degraders

Autoimmunity, Transplantation,  
Oncology, Metabolism

CT-05 potent degradation  
in immune cells



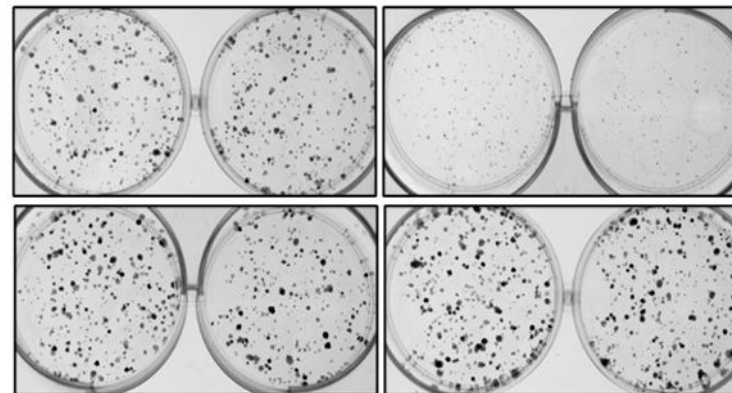
Big pharma compound  
Stopped in clinic;  
Inadequate Selectivity

Captor degrader  
High Selectivity

CT-05: no effect in non-immune cells

100 nM

500 nM



2023 expected  
milestones

Demonstration of the  
*in vivo* efficacy in  
inflammation model

High value target remains undrugged due to selectivity problems of inhibitors

Superior selectivity and prolonged efficacy thanks to degradation

# Captor's Optigrade™ platform

## Molecular Glues

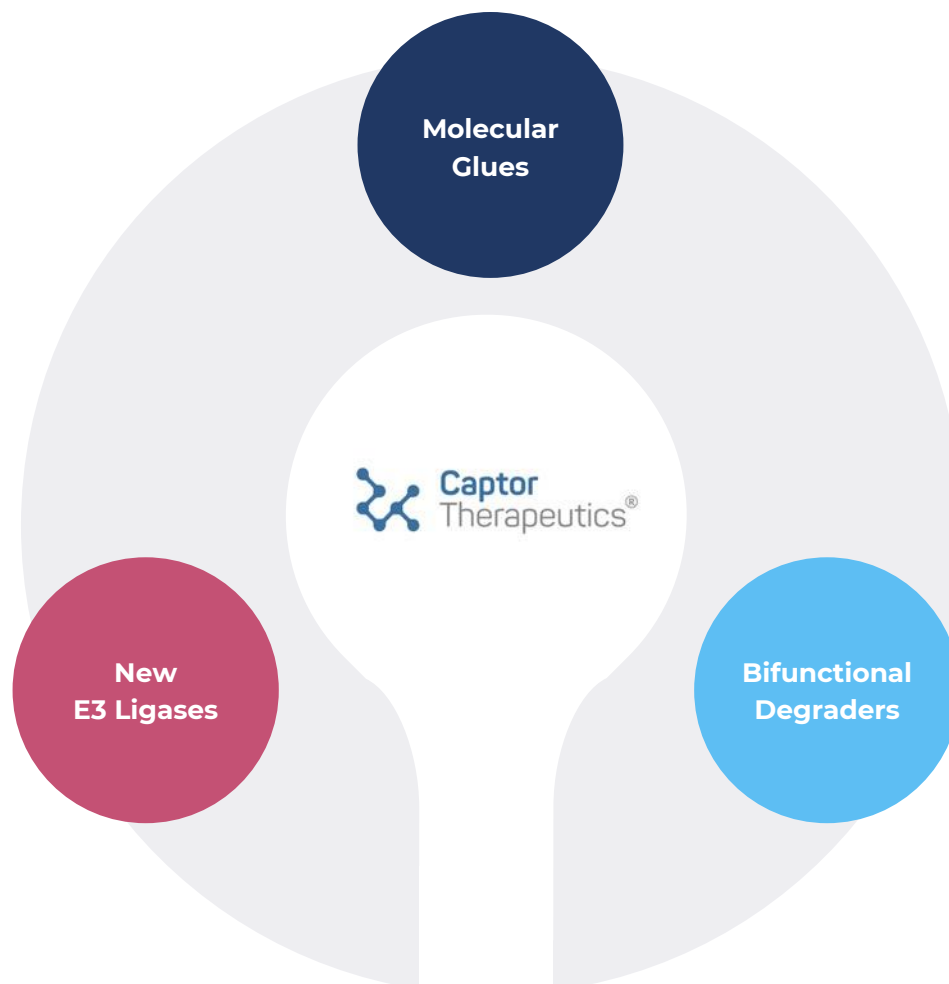
Small molecules with good drug properties

- Rational screening paradigm
- Library of proprietary molecular glues
- Selective degradation and novel efficacy profiles

## Evolving LiLis™ Platform

New generation degraders exploiting novel E3 ligases

- Leading library of E3 Ligase proteins
- Library of novel ligands
- Potential next generation degrader drugs



## Platform differentiation

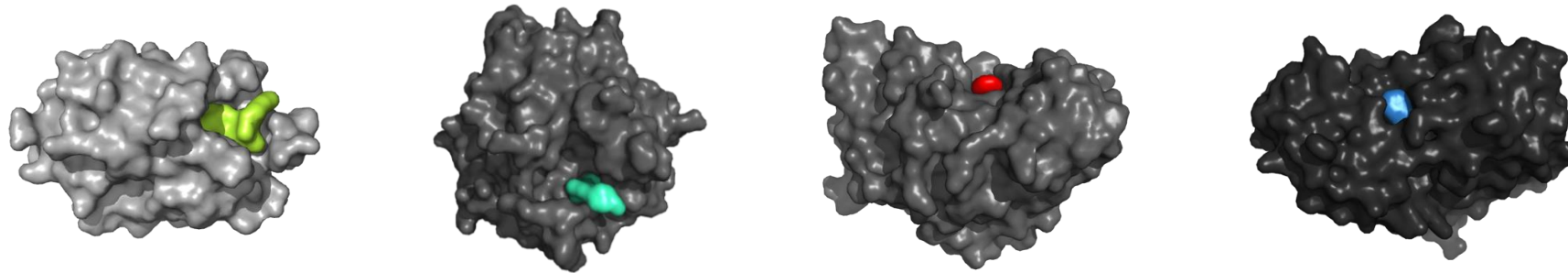
- Both molecular glues and bifunctional degraders
- Structure-based hit finding and lead optimization
- Novel and proprietary chemistry

## Bifunctional Degraders

A modular approach to degrader discovery

- Captor's ligands have improved selectivity
- Degraders against previously undrugged targets

## Optigrade™ yields chemistry for four novel E3 ubiquitin ligases



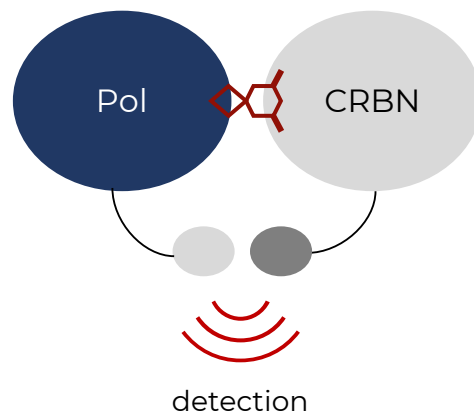
- Chemical handles as starting points for degraders discovered
- High resolution atomic structures obtained in-house
- Opportunity for molecular glue and bi-functional degrader development

**Aim of establishing degradation capability for the first E3 ligase in 2023**

# Opportunistic developments

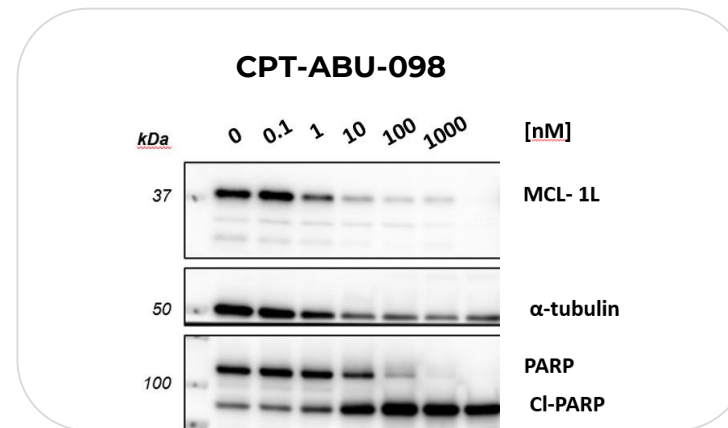
## Exploring novel target space

- Captor MG discovery engine identified novel CRBN targets
- In-house MG library for phenotypic drug discovery
- New pharmacology with novel E3-based degraders



## Antibody Drug Conjugates (ADCs)

- Ultrapotent degraders of GSPT1 and MCL-1



- ADCs offer tissue specific delivery and simple combination with other drugs
- High interest in industry to develop ADC with degrader payloads

## Analytical laboratory

- Expanding our platform to a strengthen competitive position
- Mass spectrometers– accelerate DMPK and PK/PD results
- Faster and deeper profiling of degraders across cells – proteomic mass spectrometer

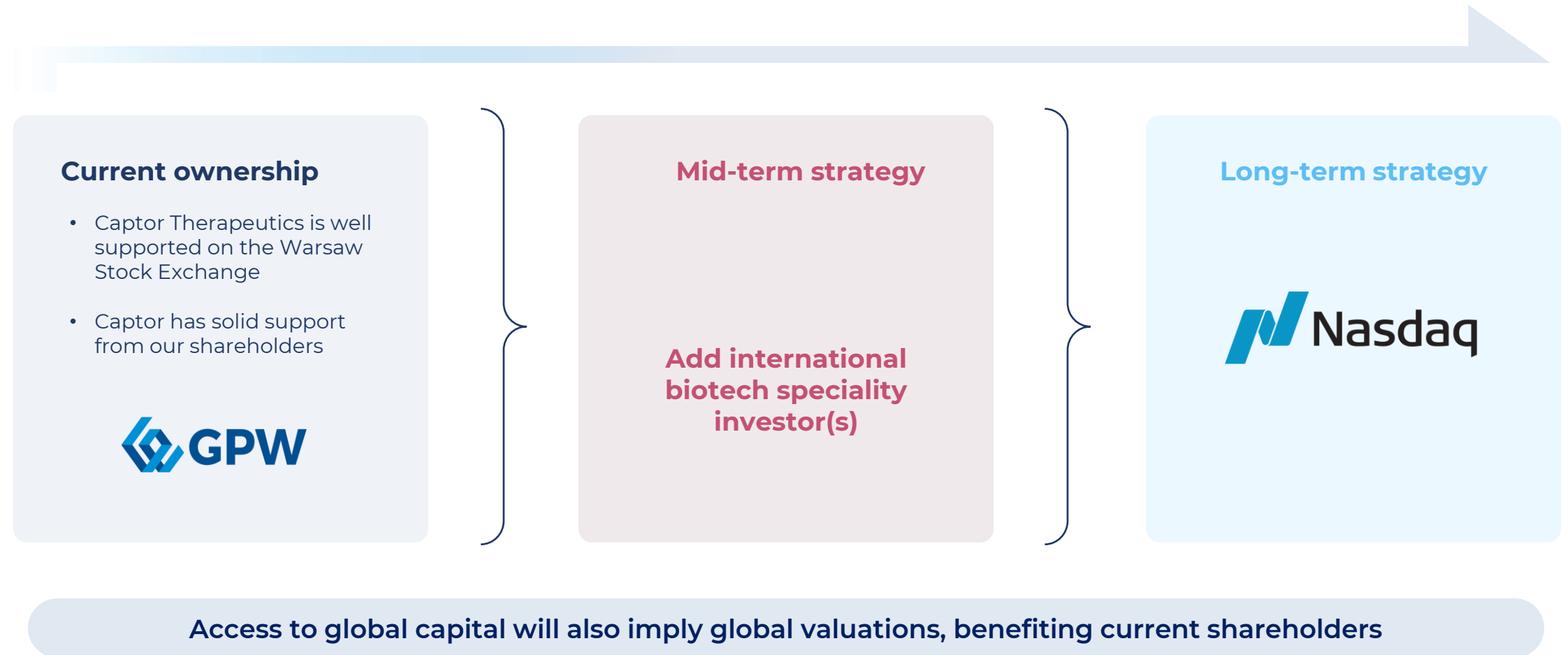
# Company pipeline projects

#	Indications	Modality	Discovery	Preclinical*	IND Filing	Phase Ia / Ib	Phase II	Next Milestone
<b>CT-01</b>	Hepatocellular carcinoma <b>GSPTI, SALL4 + undisclosed target</b>	<b>MG</b>						<ul style="list-style-type: none"> <li>✓ IND/CTA – 3Q 2023</li> <li>✓ Entering Phase I clinical in 4Q 2023</li> </ul>
<b>CT-02</b>	Autoimmunity, CNS, Metabolism, Oncology	<b>MG</b>						
<b>CT-03</b>	Liquid & solid tumours <b>MCL-1</b>	<b>BID</b>						<ul style="list-style-type: none"> <li>✓ IND/CTA – 3Q 2024</li> <li>✓ Entering Phase I clinical in 3/4Q 2024</li> </ul>
<b>CT-05</b>	Autoimmunity, Oncology, Transplantation, Metabolism	<b>BID</b>						
<b>New target projects</b>	Autoimmunity, Cancer	<b>MG</b> <b>BID</b>						
<b>New ligase degraders</b>	Autoimmunity, Cancer	<b>MG</b> <b>BID</b>						

\*Preclinical stage include IND-enabling studies, \*\*First in Human; at least 2 projects expected to enter Phase I by 2023, **BID** – Bi-functional Degradar; **MG** – Molecular Glue

  Assumed stage at the end of 2025

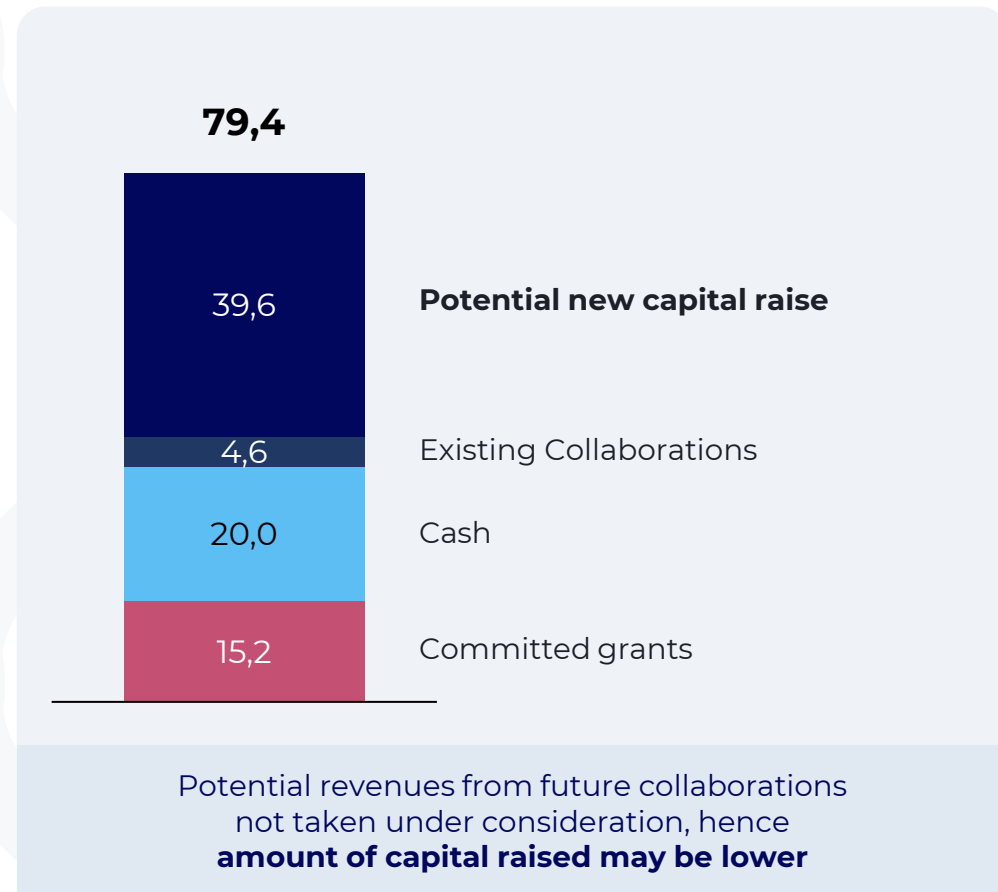
# Accessing the global capital market



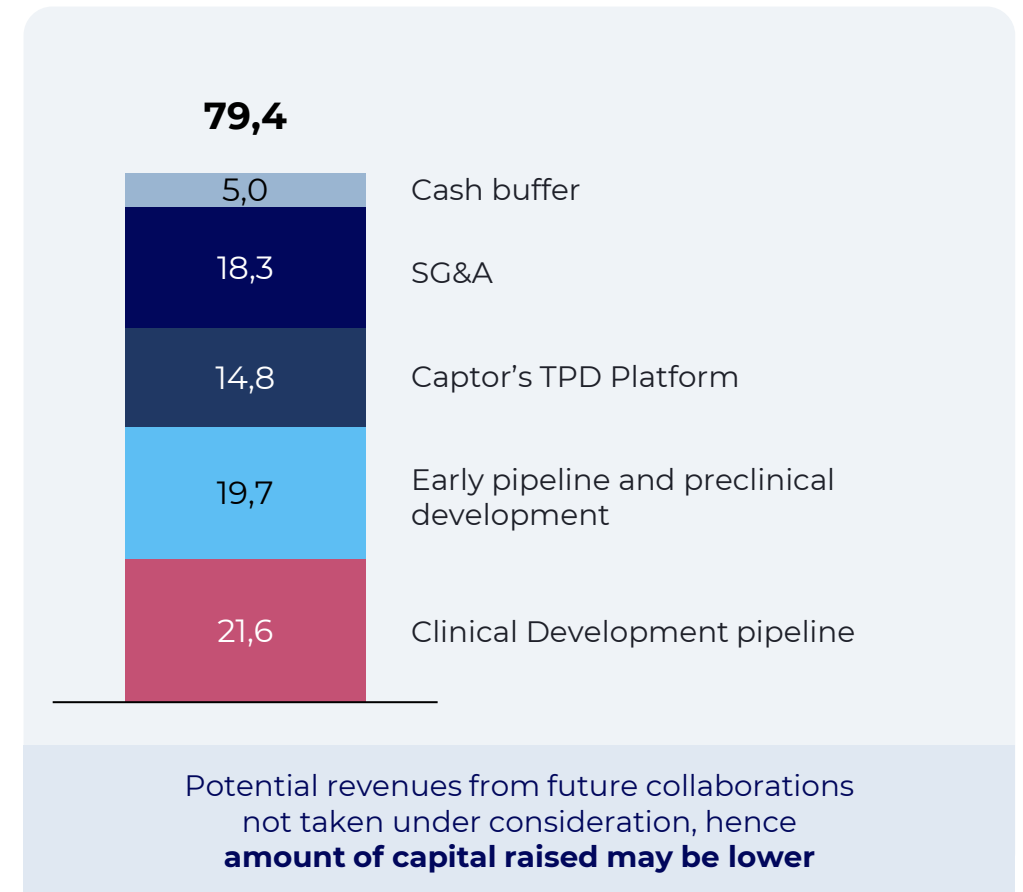


# Budget to realise our growth strategy 2023-2025

## Funding sources (US\$m)



## Application of funds (US\$m)



**By the end of the 2023-2025 period, Captor aims to build a global clinical stage company with:**

1. Two fully-owned clinical assets, with proof of mechanism demonstrated in patients
2. At least one partnered pipeline asset in development
3. Two additional fully owned early-stage assets in our pipeline
4. At least one additional platform collaboration
5. An international investor base

**To achieve these strategic objectives, the key development goals for 2023-2025 are:**

1. Investment and execution of clinical development plans for CT-01 and CT-03
2. In-vivo proof of concept for CT-02 / CT-05
3. Demonstration of target degradation exploiting novel ligase ligands
4. Expansion of Business Development capability and establishing new partnering agreements, both for the early pipeline and the platform
5. Expand our shareholder base to include international biotech specialists, as part of our long-term global capital strategy

## Projects are co-financed by the European Regional Development Fund:

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Discovery and development of a new clinical drug candidate for the eradication of cancer stem cell in the treatment of hepatocellular carcinoma, through degradation of oncofetal transcription factor  
(POIR.01.01.01-00-0740/19)

Discovery and development of non-toxic ligase ligands and their application in the treatment of autoimmunological diseases  
(POIR.01.01.01-00-0741/19-00)

Inducing apoptosis with small molecules as therapeutic intervention in multiple severe malignancies  
(POIR.01.01.01-00-0956/17-01)

Discovery and development of first-in-class of small molecule degrader as a drug candidate for the treatment of colorectal cancer  
(POIR.01.02.00-00-0073/18-00)

Development of an integrated technology platform in the field of targeted protein degradation and its implementation to the pharmaceutical market (POIR.01.01.01-00-0931/19-00)

