

智耀和铂 汇创星辰

INNOVATION HARBOUR, SHINING GLOBAL

和铂医药全球研发日

HARBOUR BIOMED GLOBAL R&D DAY

2025

10.28 中国·上海
October 28 Shanghai, China



和铂医药2025全球研发日

探界拓新，永无止境

王劲松 博士

和铂医药创始人、董事会主席及首席执行官

和铂医药2025：全球领先的可持续创新范式

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诺纳生物：世界顶尖的 生物技术创新引擎

全球稀缺的全人源抗体平台，结合
领先的“抗体+”技术模块

技术边界持续拓展，延伸至：体内
CAR-T、小核酸、核药等新领域

AI+全球最大的全人源仅重链抗体
数据库，打造独家算法



全球创新平台和产品出海： 面向世界的出海王者

2025年对外授权交易总额突破**60亿**
美金

与**阿斯利康**达成长期深度战略合作，
与多家跨国药企及顶尖Biotech开展合作

创新孵化成果覆盖多治疗领域



和铂制药：免疫与肿瘤 产品开发第一梯队

20+ 差异化产品管线，具备全球
BIC/FIC竞争力

聚焦免疫、肿瘤**2**大主战场。致力
于解决重大未满足临床需求

和铂医药的创新能力、合作成果及业绩表现广受关注

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媒体报道



代表奖项

诺纳生物进入2025年
盖伦奖
Best Startup候选名单

2025年行业引领
Biotech公司
2025中国医药决策者峰会

2024年中国医药创新
企业双抗赛道TOP5
E药经理人

和铂医药2028：全球领先的平台型制药集团

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产品开发及商业化平台

5+款商业化及准商业化产品*,
全球潜在销售峰值超**200**亿元

3+款以上产品完成全球概念验证,
全球潜在销售峰值超**50**亿美元

每年**3+**产品进入IND阶段, 未来3年
累计递交**10+**IND申请

* 通过合作伙伴、联合开发、自建商业化团队等方式, 高效推动管线建设和商业化



AI+抗体工程技术集群

营收保持**50-80%**高速增长

实现**10亿**人民币常规化收入**

构建装载**人工智能**内核的抗
体工程技术集群

** 常规化收入指非BD产生的一次性收入



全球创新赋能平台

年均达成**2**笔以上规模化产品授权,
总金额 **>15**亿美元

以与跨国药企长期合作为基石, 实
现可持续的固定合作模式。平台
出海模式将拓展至更多跨国药企

利用平台优势和资源禀赋, 至少孵
化**3+**“和铂系”创新药企

未来三年，和铂医药将进入跨越式发展时期

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🎯 2025

📊 2028 愿景

疾病领域

肿瘤、自免

- ✓ 全面升级为以临床需求为导向的平台型公司

拓展至代谢、中枢神经、抗衰 ...

技术平台
& 治疗模式

抗体+

- ✓ 分子类型不断丰富
- ✓ 以内部拓展、投资等路径探索拓展技术边界

拓展至 体内Car-T、siRNA、mRNA、小分子等领域

即将商业化
及全球概念验证产品

3+款

- ✓ 每年3+项目进入临床阶段
- ✓ 撬动中国临床优势和效率，高效推进管线
- ✓ 积极探索外部产品引进机遇

8+款

收入性质

授权合作收入为主

- ✓ 构建多元化业务，整合合作伙伴网络、联合开发与自营团队等渠道，实现投资回报最优化

授权合作+产品销售收入
双引擎

持续挖掘“抗体+”复杂分子在多种疾病领域的治疗潜力

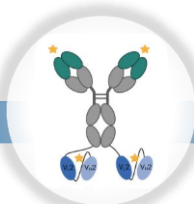
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全面升级为以产品为核心的平台型公司，从技术导向转向临床需求导向

分子类型



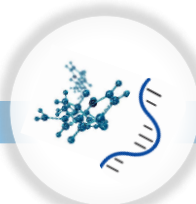
双抗



多抗



XDC



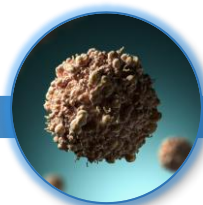
小分子等多样化
创新分子



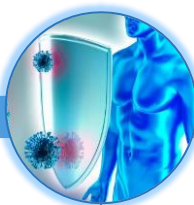
药物递送

疾病领域

肿瘤



免疫疾病



代谢疾病



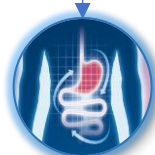
内分泌疾病



心血管疾病



神经退行性疾病



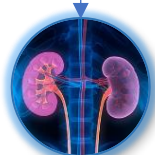
炎症性肠炎



呼吸科



皮肤科



肾病

五大创收源头驱动多元增长，重塑平台型制药集团独特价值

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未来三年收入增长示意图



2028年目标

- 实现至少**5款**商业化及准商业化的自免、肿瘤领域产品矩阵
- 潜在全球销售峰值 > **200亿人民币**
- 年均达成**2笔**以上规模化产品授权，总金额 > **15亿美元**
- 年均转化**数千万美元**高可见度里程碑收入
- 诺纳生物保持**50-80%+**高速增长
- 实现目标**10亿人民币**常规化营收规模
- 以**阿斯利康**长期合作为基石，实现年均可持续**固定收入**贡献，并持续获得**项目选择权**及**里程碑**等可持续收入
- 拓展至更多**全球前二十大跨国药企**...
- **Hub and Spoke**模式孵化多家创新药企
- **Windward Bio**、某**NASDAQ**上市公司少数股权
- 与**中国科学院院士**共同孵化前沿技术创新药企

Beyond Harbour • Into Future

初心为舵，创新为帆
驶出港湾，奔赴蓝海



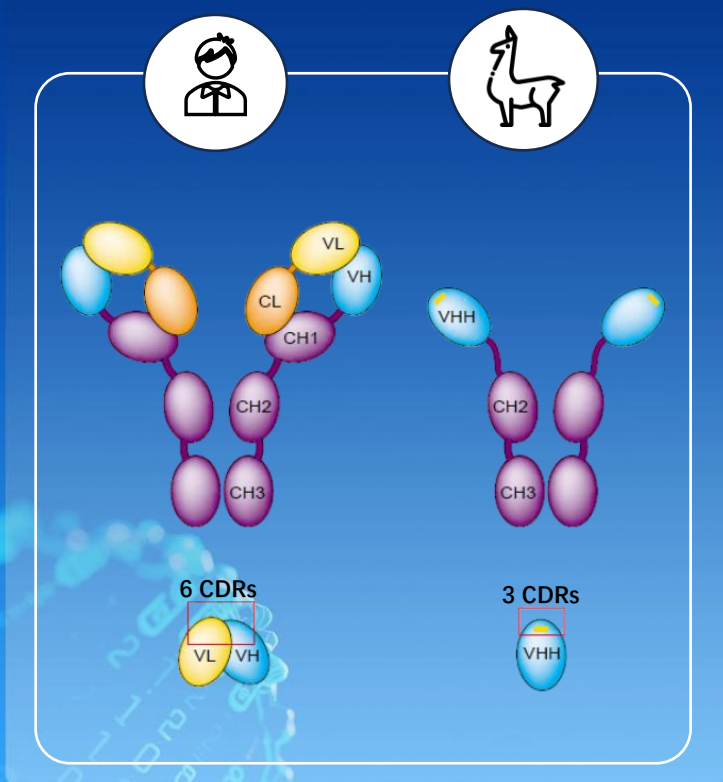
加速抗体研发的未来：AI HCAb模型发布

Accelerating Next-Gen Antibody Discovery:
The AI HCAb Model Launch

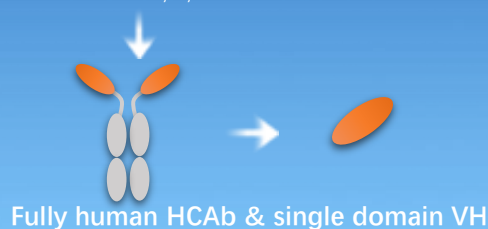


Human HCAb from Harbour Mice® is a New Class of Biologics as Foundation for Complicated Modalities

Naturally occurring classical antibody from Human and Heavy Chain Only Antibody (HCAb) from Camelidae

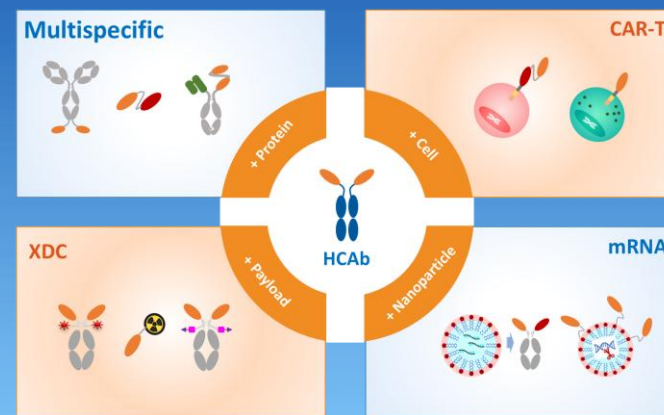


Harbour Mice® HCAb is the first transgenic mice platform to develop human HCAb

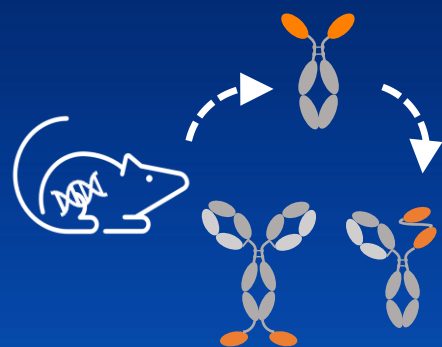


Human HCAb is foundation to generate various complicated modalities for next-gen innovative biologics

- ✓ Clinically validated
- ✓ Global IP protection
- ✓ Excellent affinity and properties
- ✓ Plug and play



Cutting-edge Antibody Platforms Enable Sustained Invention of Novel Molecules



IDEA



Antigen
Design



Immunization



Discovery



in vitro



in vivo



Engineering



CMC

IND

How AI is Reshaping the Antibody Discovery?

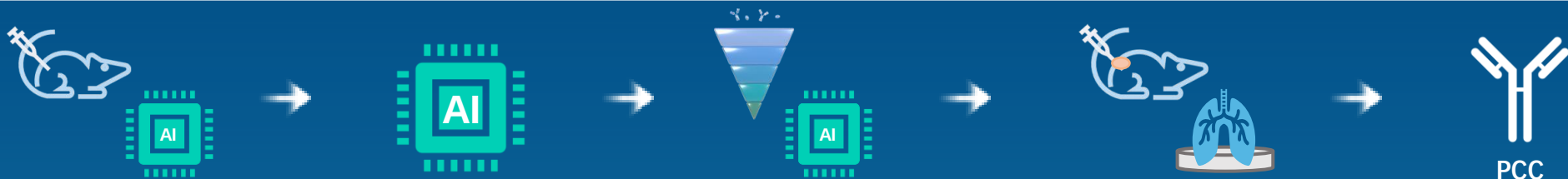
Level 0



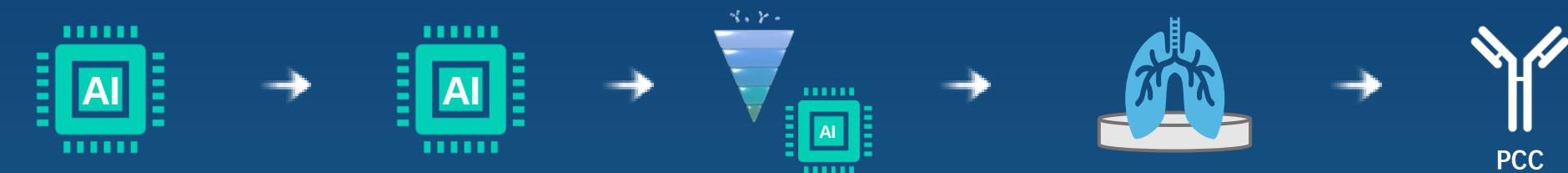
Level 1



Level 2



Level X



End-to-End Workflow from Sequence Generation to Wet Lab Validation



1. AI HCAb Sequence Generation

Generate candidate antibody sequences using a fine-tuned LLM



2. AI Guided Design

AI HCAb Classification Model
HCAb classification model to filter sequences



AI HCAb Developability Prediction Model
select the sequences with good properties



Structural Analysis
structure prediction by multiple models and rank high-potential sequences



3. Wet Lab Validation

Produce selected candidates and analyze their biological activity in the lab.

Generation of Human HCAb: The First Puzzle

Facts and Mysteries about Harbour Mice® HCAb

❖ Facts

- Human VH can't stand alone without VL.
- “Hallmark residues”: signature of Camel VHH different from Human VH.
- Harbour Mice® HCAb incorporates natural human germline V,D,J genes, and generate soluble and functional human HCAb without camel hallmark residues.

❖ Mysteries

- How does human HCAb fold into soluble structure ?
- What is the difference between the VH from human HCAb and the VH from human H2L2 ?

First puzzle: how to distinguish VH of human HCAb and VH of H2L2

Example:

anti-CTLA4 human HCAb VH (PDB: 7DV4) and anti-SARS-CoV-2 human antibody C098 VH (PDB: 7N3G) have similar sequences (90% identity) and structures (0.7 Å RMSD)

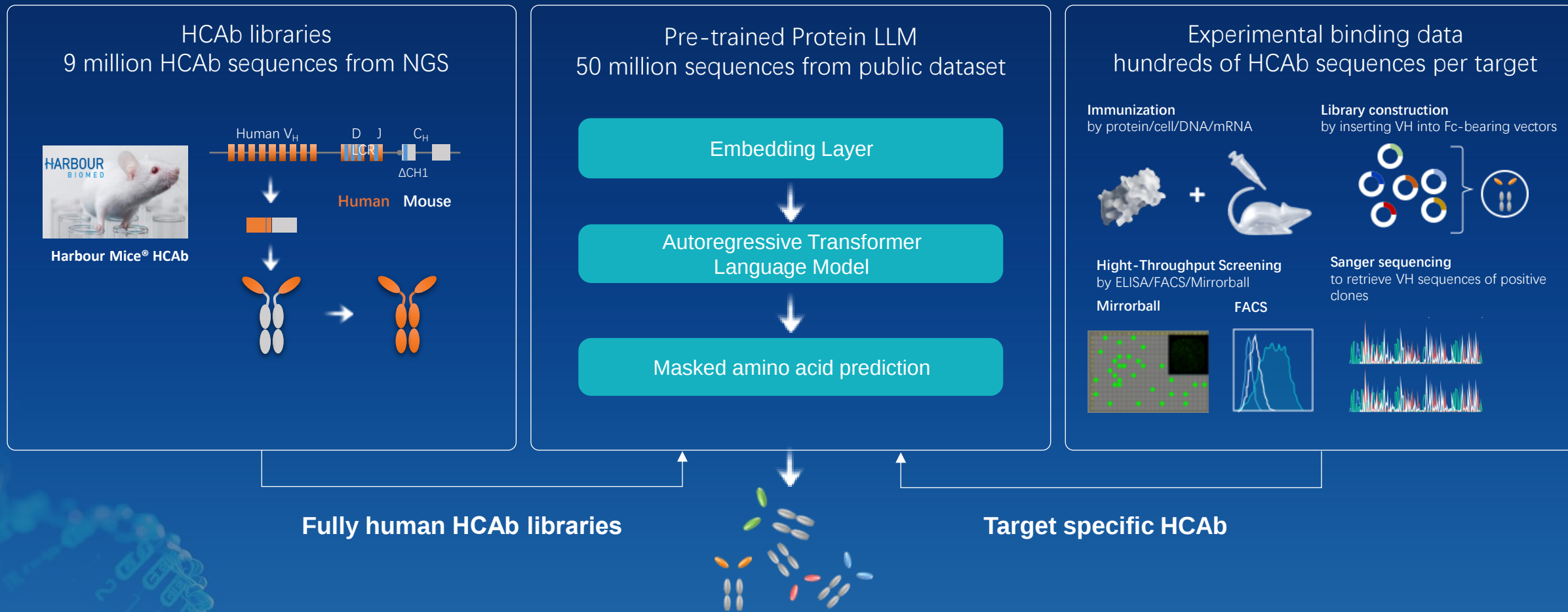
VH structures
superposed (PDB:
7N3G, 7DV4)



VH sequence alignment

	20	40	60	
7DV4 - HCAb VH	EVQLVESGGGLIQPGGSLRLSCA	VSGFTVSKNYMSWVRQAPGKGLEWVSV	YSGGSKTYA	60
7N3G - FAB VH	EVQLVESGGGLIQPGGSLRLSCA	ASGFTVSSNYMSWVRQAPGKGLEWVSV	YSGGSTYYA	60
	80	100		
7DV4 - HCAb VH	DSVKGRFTISRDNSKNTLYLQMNSLRAEDTAVYYCAR	AVPHSPSSFD	IWGQGTMTVTVSS	119
7N3G - FAB VH	DSVKGRFTISRDNSKNTLYLQMNSLRAEDTAVYYCAR	DL-YSSGGTD	IWGQGTMTVTVSS	118

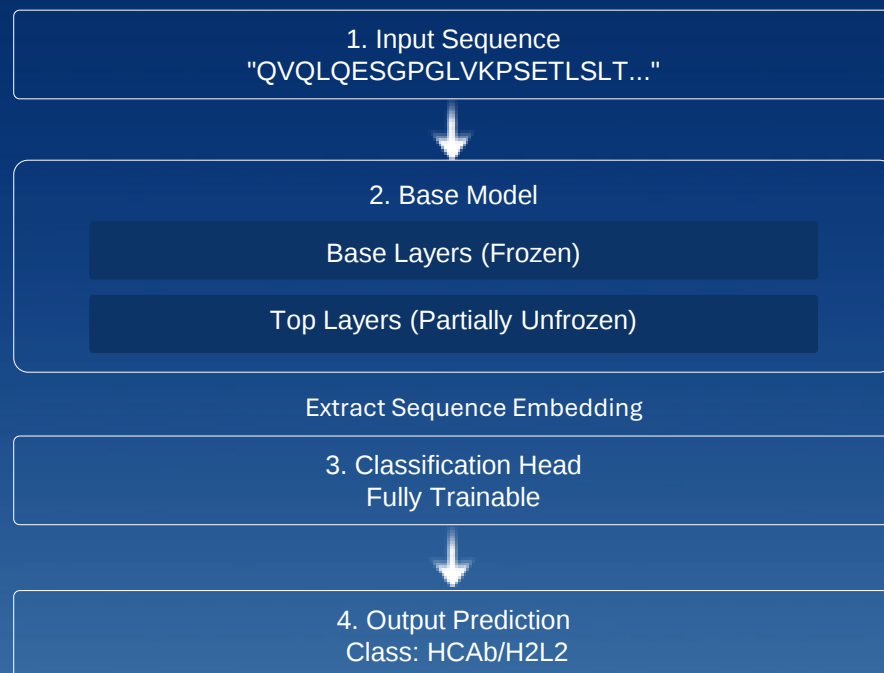
AI HCAb Sequence Generation Model



Compared to conventional immunization, the AI-assisted approach can generate an order of magnitude more candidates with a significantly higher success rate

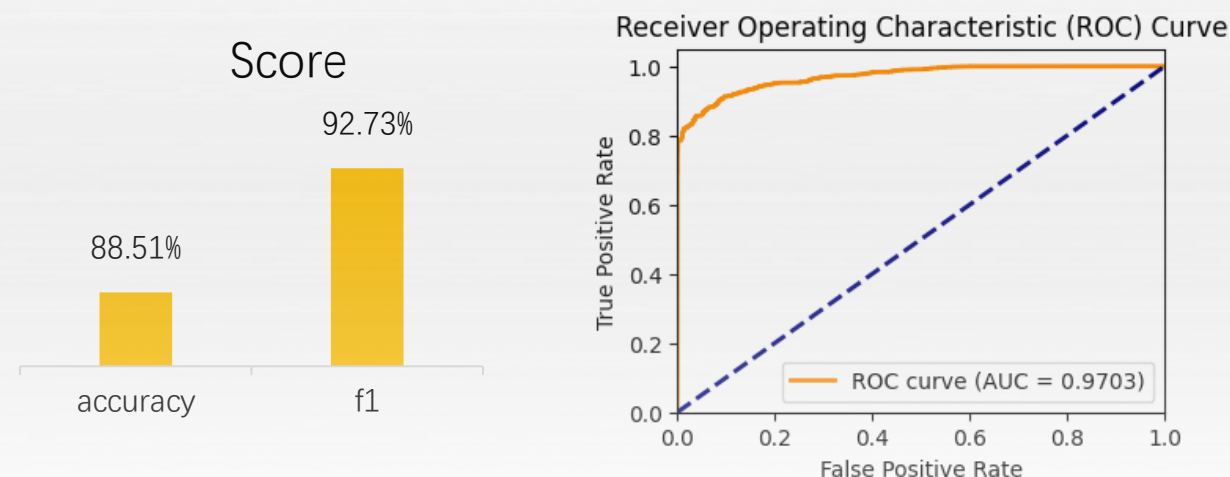
AI HCAb Classification Model

Partial Fine-tuning Architecture



HCAb classification model in test-set

Testing dataset: 14k

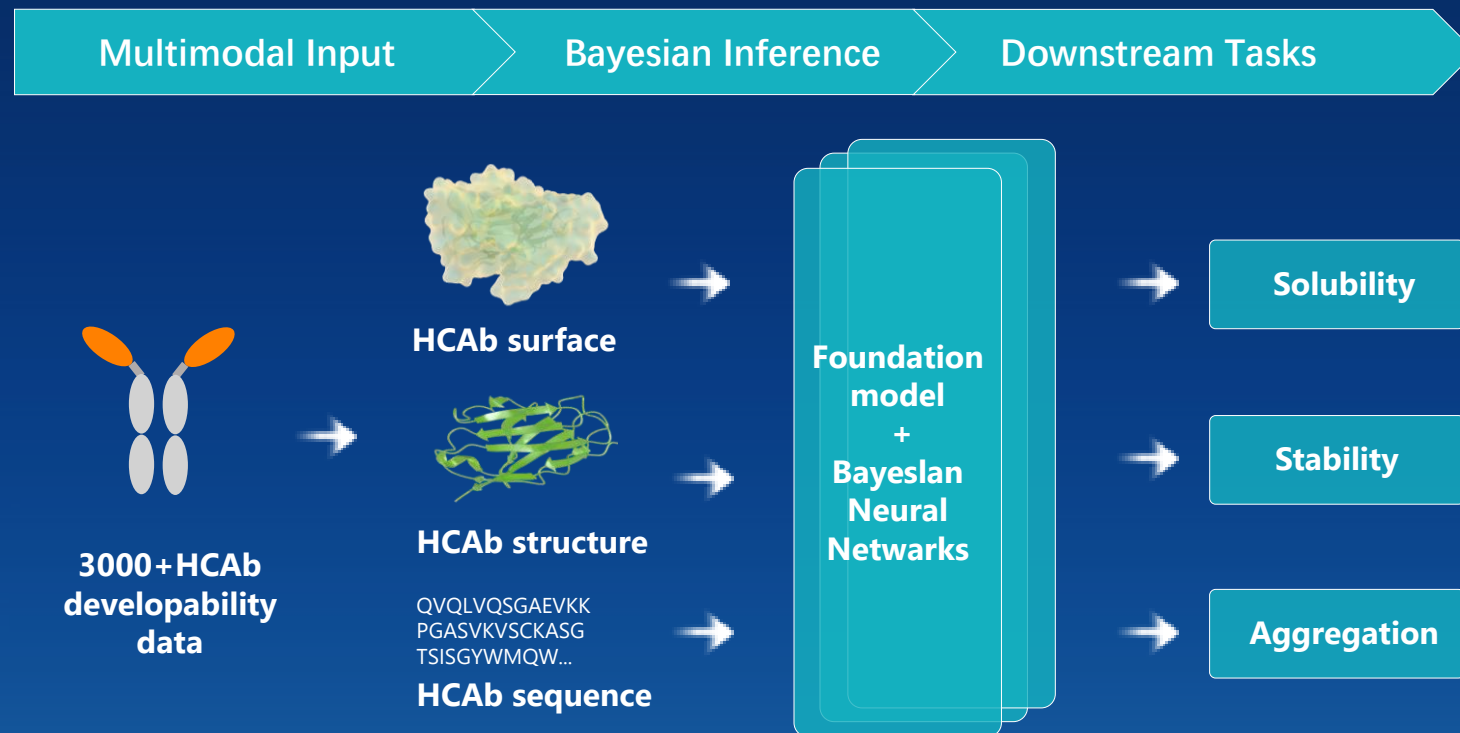


True Label Predicted Label

Sequence

H2L2	H2L2	EVQLLESGGGLVQPGGSLRLSCAAS	GFTFSSYA	MSWVRQAPGKLEWVSA	ISGSGGRT	YYADSVKGRFTISRDNKNTLYLQMNSLRAEDTAVYYC	AKHEGLGVFDY	WGQGT	LTVVSS
HCAb	HCAb	EVQLLESGGGLVQPGGSLRLSCAAS	GFTFSSYA	MSWVRQAPGKLEWVSA	ISGSGG	STYYADSVKGRFTISRDNKNTLYLQMNSLRAEDTAVYYC	EKVDIVATD	WGQGT	LTVVSS
		FR1	CDR1	FR2	CDR2	FR3	CDR3		FR4

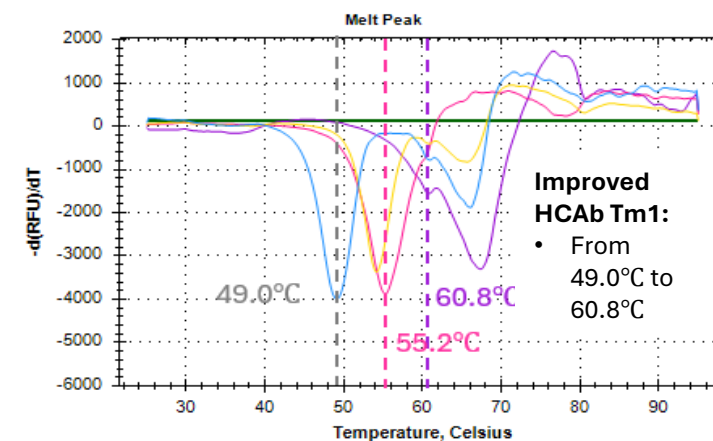
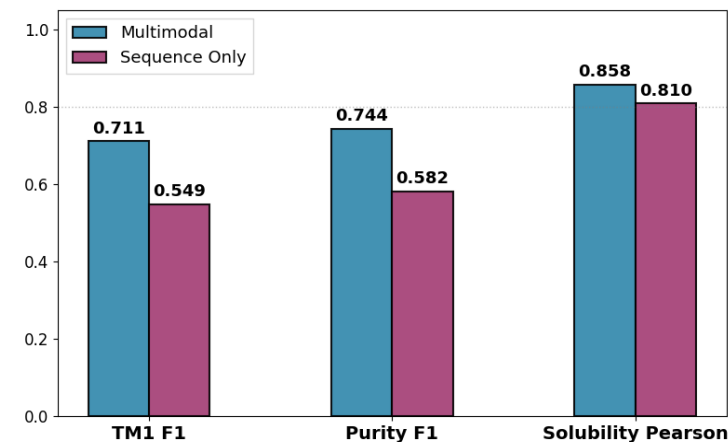
AI HCAb Developability Prediction Model



Our model achieves SOTA performance in the prediction of

- HCAb Stability (Tm1 by DSF)
- HCAb Solubility (HIC-HPLC)
- HCAb Aggregation (SEC-HPLC)

Model evaluation in wet-lab experiment



Target-specific AI-Powered HCAb Discovery

AI-Powered Antibody Discovery Paradigm 1.0

HCAb Harbour Mice®
immunized with Target



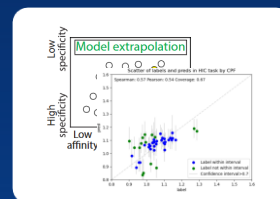
Data collection



NGS for pre- and post-
enrichment antibody
repertoire



Predict binder
sequences from
NGS data



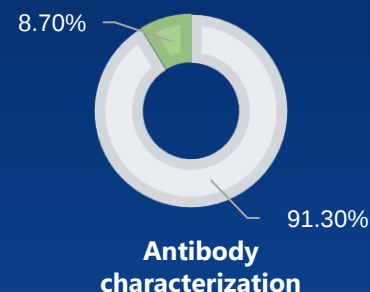
Score other
properties by AI
models



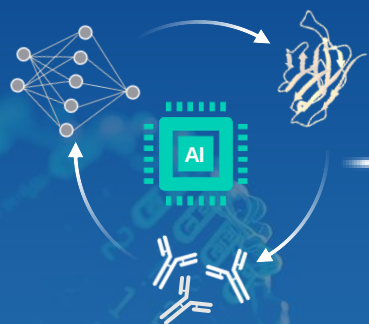
Recombinant
antibody
production

Wet-lab evaluation

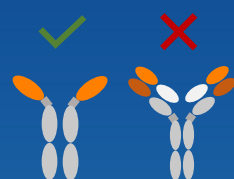
■ NonBinder ■ Binder



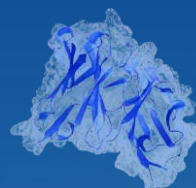
AI-Powered Antibody Discovery Paradigm 2.0



HCAb generative model



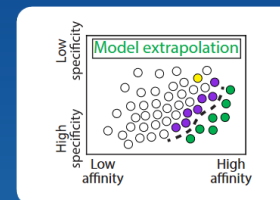
HCAb filter



Structure
modeling



Developability



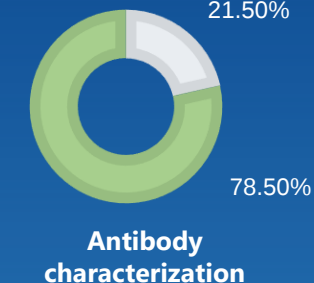
Affinity ranking



Recombinant
antibody
production

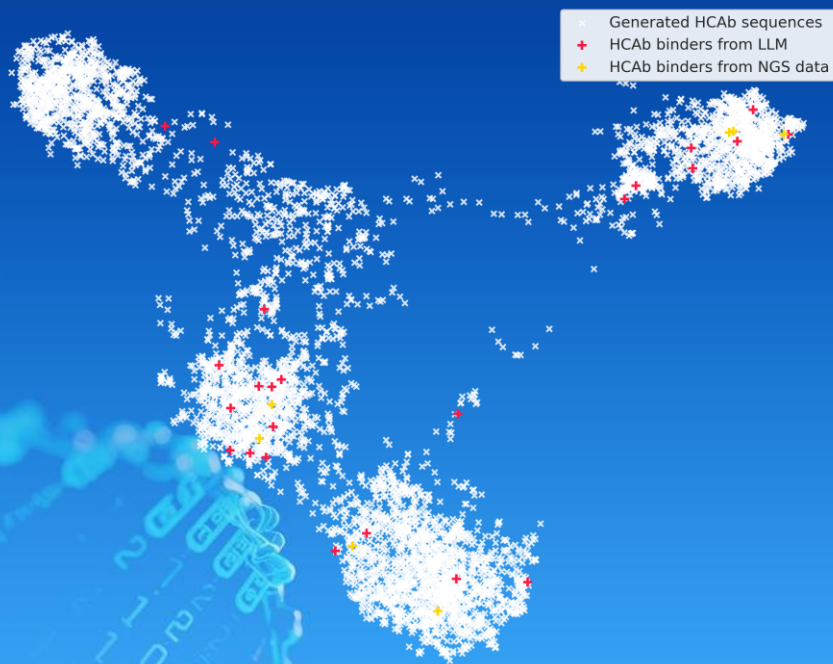
Wet-lab evaluation

■ NonBinder ■ Binder

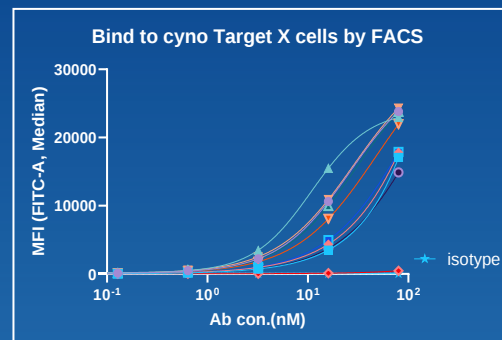
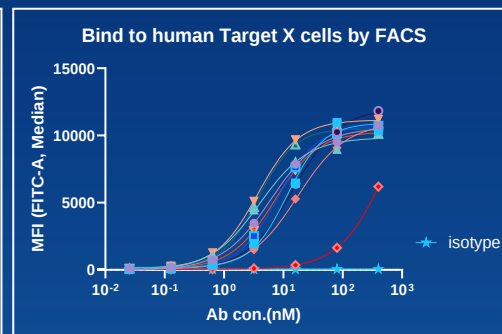
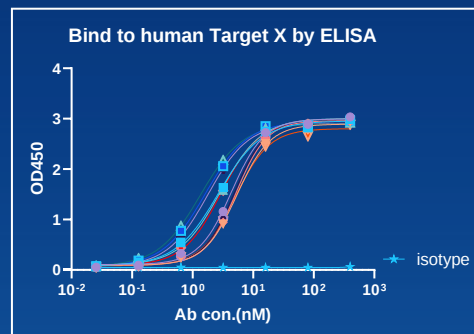


Superior Wet-Lab Performance of the AI HCAb Model

AI HCAb model generates vast sequence diversity beyond known binders, enabling the discovery of novel binders with superior properties.



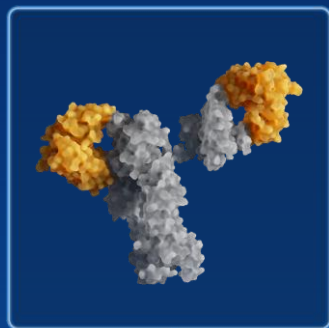
Selected HCABs with specific binding activities to both human and cyno monkey



Summary of produced HCABs:

- Average yield > 700 mg/L
- 20 HCABs ≤ 10 nmol (EC50)

AI Powered Dry to Wet for Next Generation of Antibody R&D Platform



Leverage Harbour BioMed proprietary HCAb technology and inhouse data, build first-in-class generative model to make fully human heavy chain only antibodies.



Embedding Layer

Autoregressive Transformer Language Model

Masked amino acid prediction



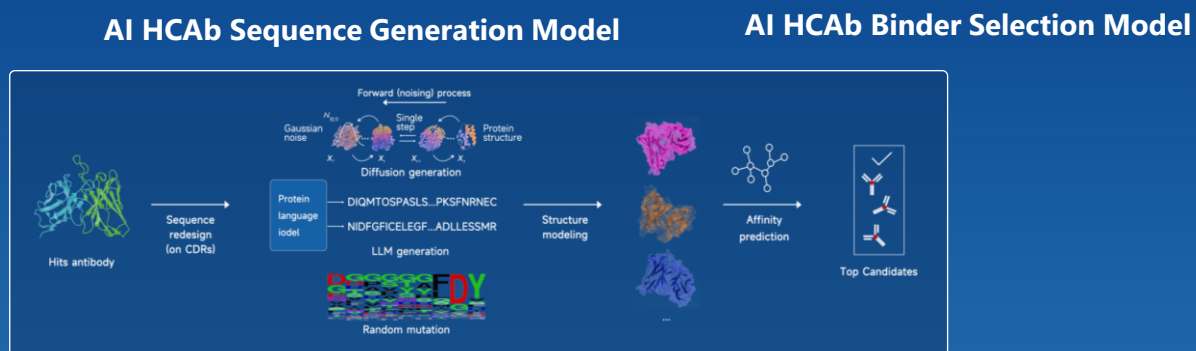
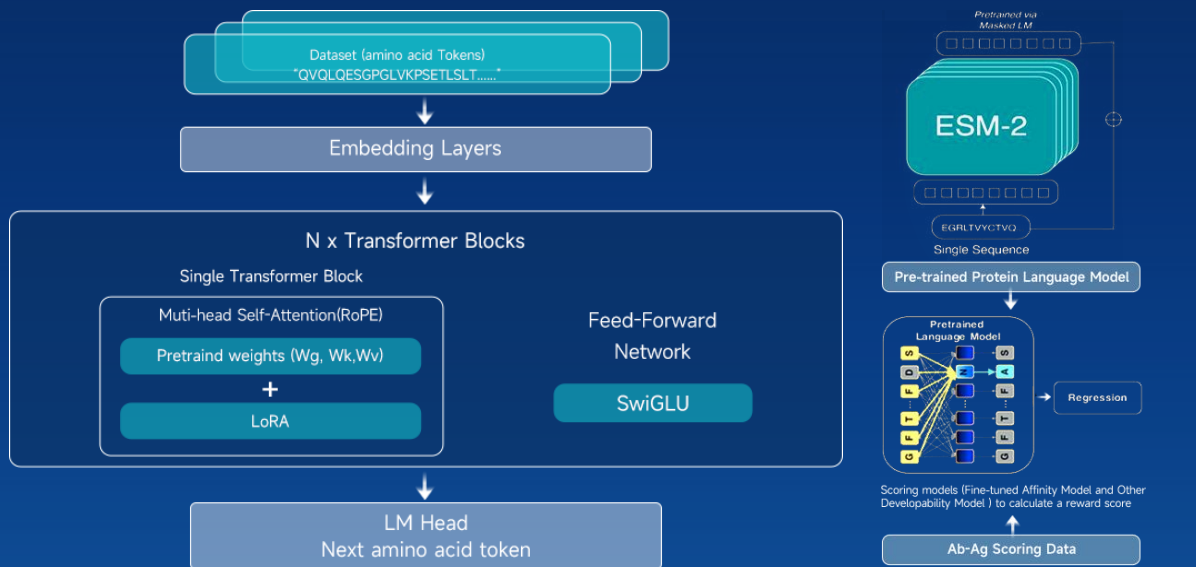
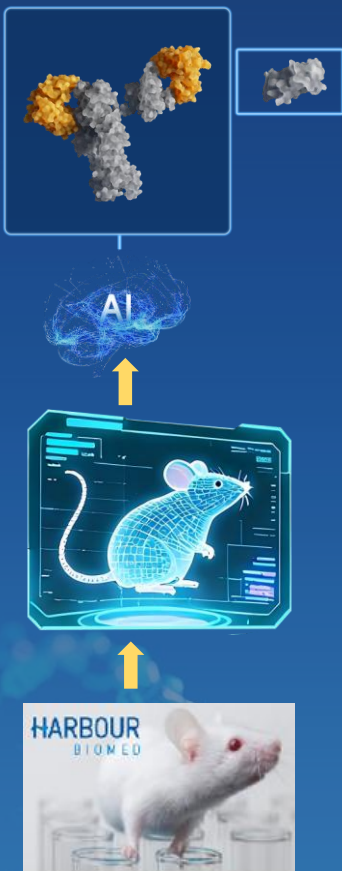
Automated Protein Expression and Characterization System

Utilizing high-throughput automation instruments significantly expands the capacity of molecule generation and characterization, enhancing the probability of identifying candidates, generating large-scale, high-throughput wet-lab data which can further enhance AI models.

We're not only discovering drugs, we're discovering the unmet medical needs with data-driven, predictive process. We welcome all kinds of partners to co-create the ecosystem with us.

AI Accelerates a New Era of Antibody Therapeutics Across Multiple Fields

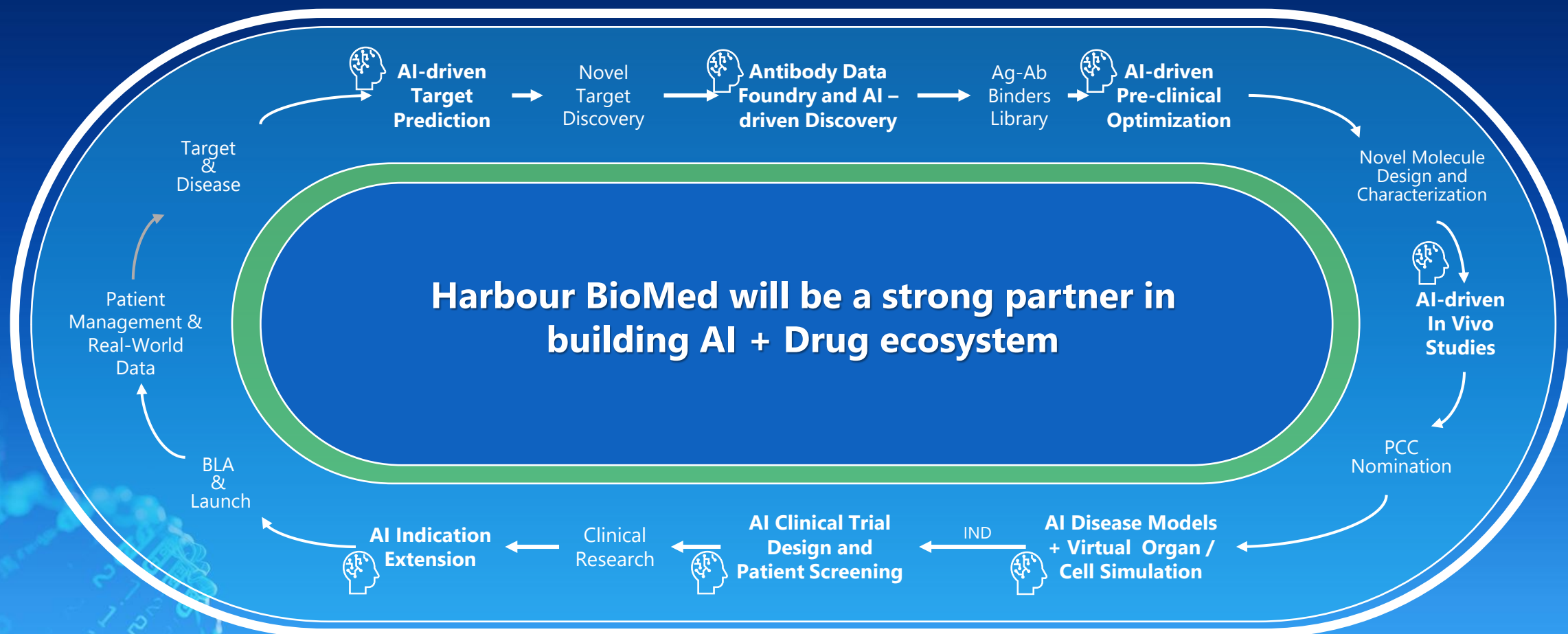
HCAb Harbour Mice immunized with Target



Fully Human VH Antibody Innovative Applications

- Multi specific antibody drugs
- Targeted delivery of radiopharmaceuticals
- In vivo CAR-T technology
- Inhaled therapeutics for pulmonary diseases
- Oral therapeutics for gastrointestinal diseases

Shaping the Future of Drug Discovery



诺纳生物创新战略及远景

Nona Biosciences Innovation Strategy and Vision

洪涤 博士
诺纳生物首席执行官



诺纳生物是以技术创新为先的平台公司

HARBOUR
BIOMED

和铂医药

专注于推进公司全球产品管线及变革性疗法的研发

- 在免疫与炎症领域布局广泛且具高度差异化的产品管线

HBM9378 HBM7020

- 通过整合外部资源加速产品开发进程

HBM7022 HBM9033

AstraZeneca

Pfizer Seagen®



诺纳生物

运用尖端技术平台，及开放创新模式，赋能全球医药创新

- 诺纳生物成立于2022年
- 不断创建生物医药尖端技术平台

Bispecific Abs/mRNA/ADCs/CAR-T/NK...

- 为全球医药创新客户提供 Idea to IND (I to I™) 全链创新
- 持续构建创新孵化的生态体系

moderna

LCB
LegoChemBio

DualityBio
映恩生物

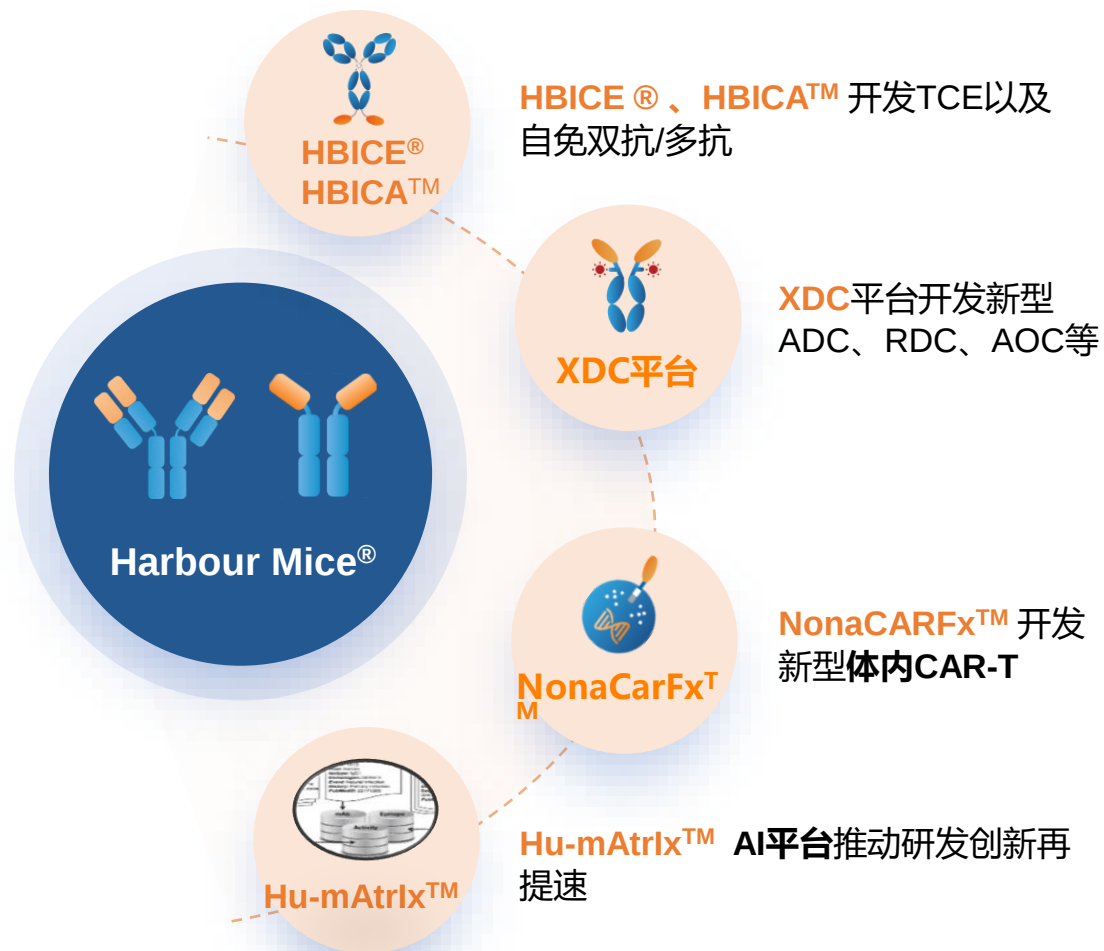
Dragonfly
THERAPEUTICS

MYTHIC
THERAPEUTICS

Dana-Farber
Cancer Institute

诺纳生物四大技术平台关键业务进展

HARBOUR
BIOMED



关键业务进展*



诺纳生物成功提名
2025年度医学界诺贝尔奖
盖伦奖 “Best Startup”

110+
合作伙伴

19+
分子进入临床
开发阶段

320+
合作项目

2家
已孵化生物科技
子公司

120+
早研团队

+173%
25H1订单金额**
较24H1增加

*累计至2025年6月30日数字
**研究及技术许可服务相关订单金额

诺纳生物2028：加强临床前研发能力，拓展临床概念验证新平台

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加强自身“IND Enabling”平台建设，积极拓展临床概念验证业务

GLP
毒理

CM
C

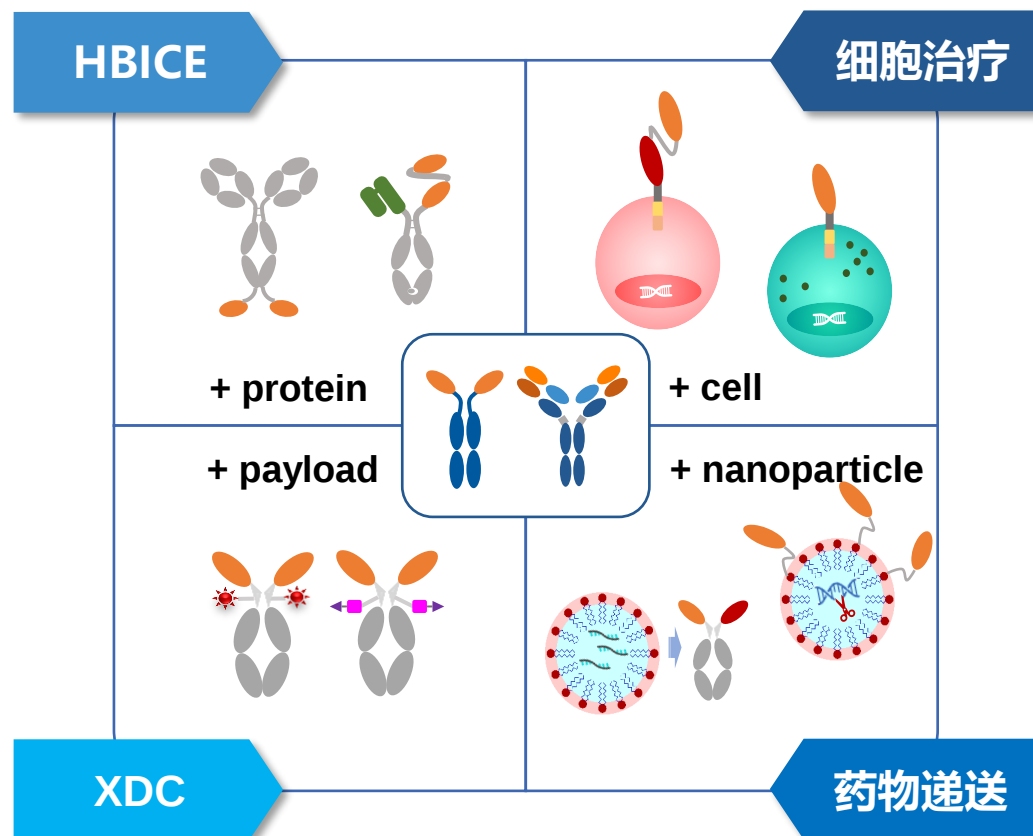
临床概
念验证



诺纳生物2028：构建下一代抗体工程创新技术集群

HARBOUR
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诺纳生物四大技术平台



诺纳生物将构建更多样的技术集群

HIBCE®

类器官

XDC

药物递送

体内 CART

siRNA

细胞治疗

下一代TCE

中枢神经递送

mRNA

抗体工程AI

AOC

...

诺纳生物2028：AI平台推动医药创新全面进入人工智能时代

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临床前

HBM药物研发历史数据



数字化拼图



Hu-mAtrix™

数字化和铂小鼠

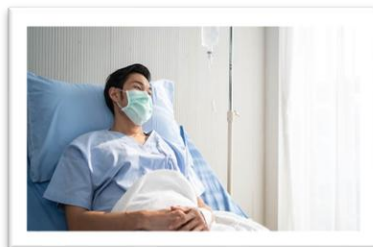


数字化抗体药物生成

- ✓ 提高药物研发成功率
- ✓ 减少研发时间
- ✓ 降低研发阶段费用

临床

患者人群



- ✓ 与政府、医院进行基础设施建设
- ✓ 获取临床大数据

数字人



数字化临床试验

- ✓ 提高临床试验的效率
- ✓ 极大降低临床试验的费用

诺纳生物2028：建立“自动化-AI”新药发现螺旋进化循环

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诺纳生物2028年愿景：构建A³医药创新“新基建”平台

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2028年诺纳生物将进化成为通过装载人工智能内核的抗体工程技术集群，构建从立项到临床 Idea to IND (I to ITM) 生物医药创新“新基建”平台，为全球医药创新客户赋能，并构建生物医药创新孵化生态。

凭借全球领先技术平台，持续赋能全球生物医药合作伙伴，加速下一代创新疗法的开发与应用

赋能全球医药创新客户

A³技术矩阵

Antibody Engineering
抗体工程

AI Plus
人工智能平台

Automation
自动化湿实验室

建立诺纳创新孵化生态

以全球领先技术平台为基石，与全球顶尖学术及科研机构开展战略合作，共同孵化前沿疗法

2028年实现目标10亿人民币常规化营收规模



开创更前沿的
技术平台



拓展更广阔的
全球合作



构建更开放的
产业生态

持续为病患和投资者带来更有价值的产品和服务

和铂医药研发战略和远景

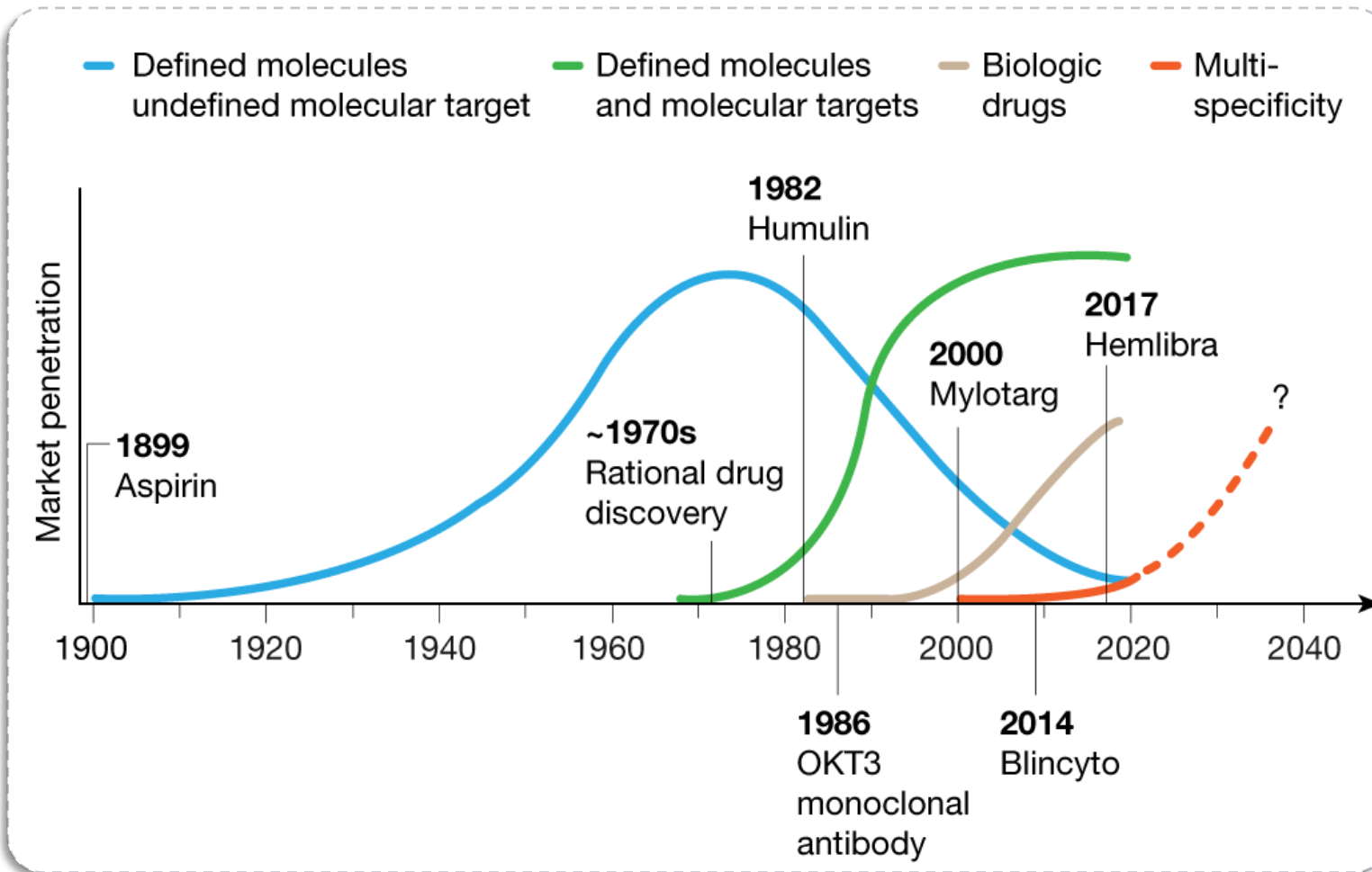
Develop the Next Generation Multispecific Medicines
- *The Right Technology for the Right Therapeutics*

戎一平 博士

和铂医药执行董事、首席科学官



“1T1D”Era Passed, A New Wave of Transformative Innovation - Multispecific Medicine is Now Upon Us



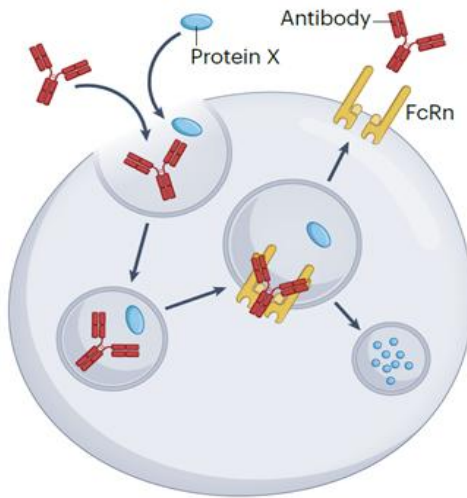
Raymond J. Deshaies, “Multispecific drugs herald a new era of biopharmaceutical innovation”, Nature (2020)

Multispecific Molecular Medicines Require Specific Targeting Moiety

Representative Drug Classes

Fc/Albumin fusion protein
(Etanercept=TNFR-Fc,
Abatacept=CTLA4-Fc)

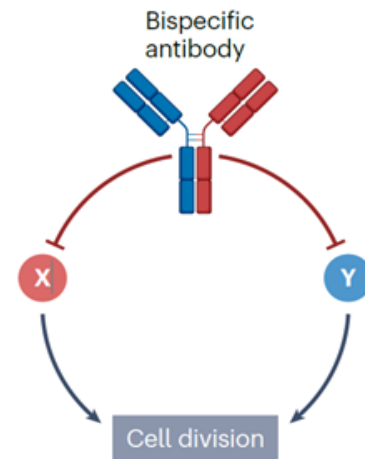
Stabilizers
Enhance persistence



Rescue endocytosed protein

Bispecific antibody
(Amivantamab-aEGFRxcMet
Faricimab-aAng2xVEGFA
Tirzepatide-GLP1xGIP)

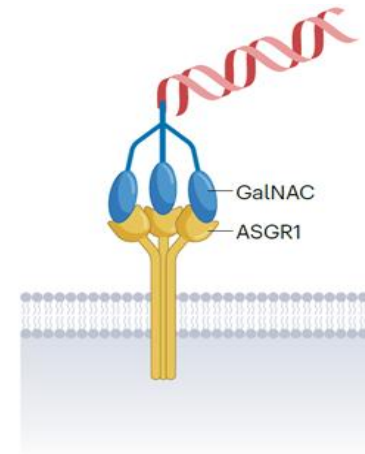
Multi-actives
Overcome redundancy



Block cancer growth

XDC
(ADC, RDC, AOC, DAC, ISAC)

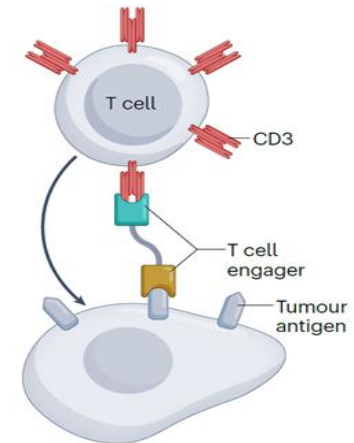
Tetherbodies
Enhance therapeutic index



Silence liver protein

TCE PROTAC
(Blinatumomab-CD19xCD3,
Risdiplam-SMA2RNA.U1snRNP
Emicizumab-aFIXa/FX)

Matchmakers
Overcome
“undruggability”



Kill tumor cell

Proximity Dependent Mode of Actions

Raymond J. Deshaies, “How multispecific molecules are transforming pharmacotherapy”, Nature (2025)

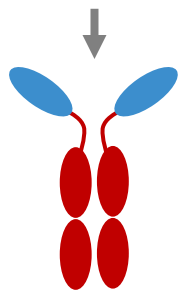
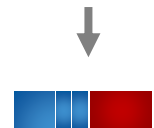
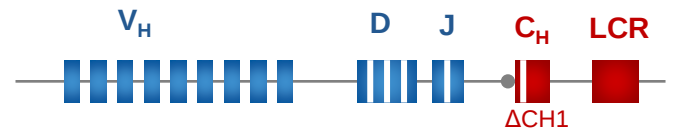
Human VH from HCAb Technology is Ideal Targeting “Missile” for Multispecific Medicine

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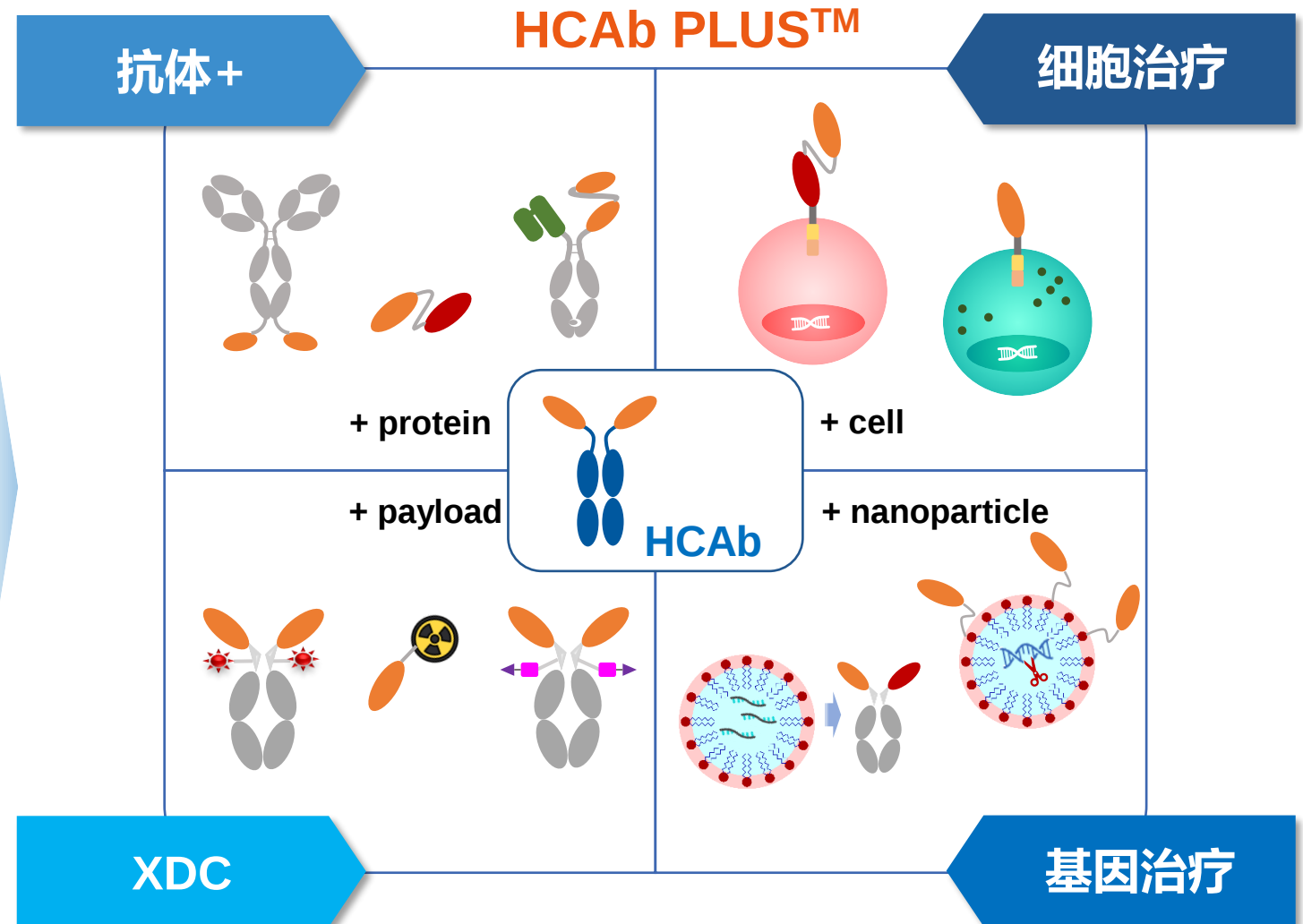
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Harbour Mice® HCAb



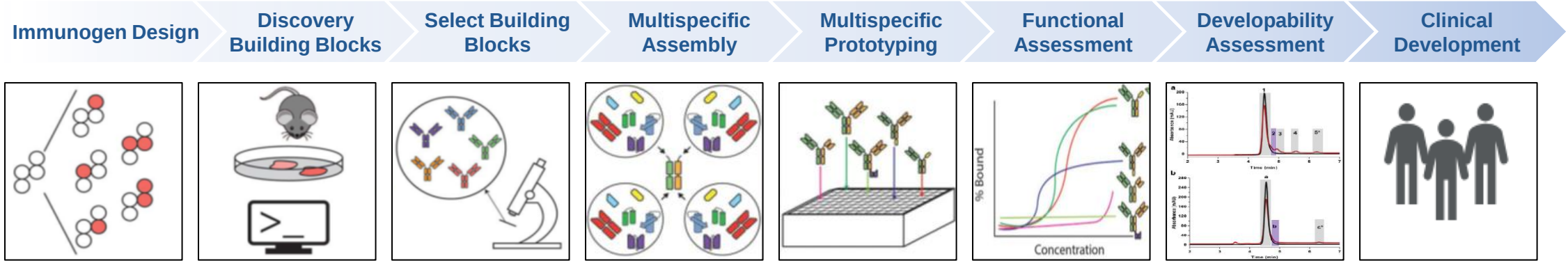
Fully human HCAb

Human VH or
single domain Ab



HCAb Plus Technology is a Systemic Modality Solution from Idea to IND

Multispecific Medicine Discovery Roadmap



- ✓ mRNA/DNA/inducible immunizations
- ✓ Antigen design

- ✓ Harbour Mice
- ✓ AI Hu-mAtrix™

- ✓ Single B cell tech
- ✓ NonaCarFx screening

- ✓ Multispecific protein engineering
- ✓ Tissue specific VH guided delivery

- ✓ 3D organoid models
- ✓ Humanized animal models
- ✓ In-vitro functional assays

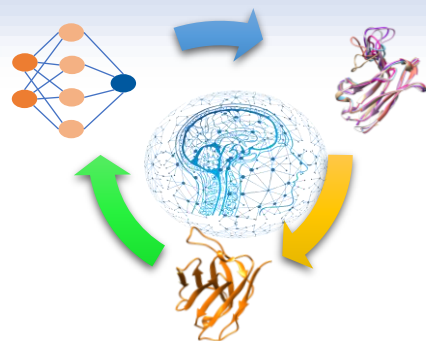
- ✓ Cutoff biophysical parameters
- ✓ Mammalian/phage/yeast display for lead optimization
- ✓ AI Hu-mAtrix aided LO

- ✓ IIT, ph1, 2, 3, BLA

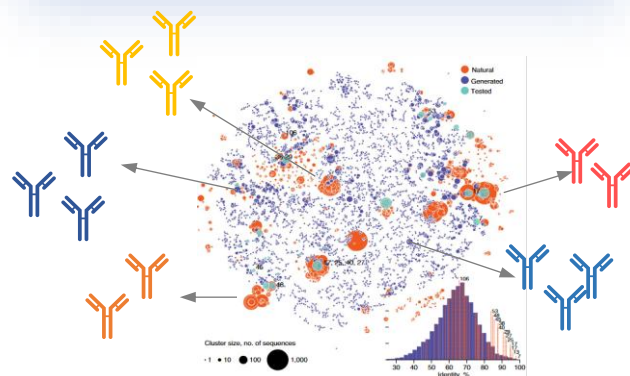
Harbour HCAb Plus Technology – The Solution for Multispecific Medicines or Complex Biologics

AI-aided Drug Discovery for Induced-Proximity to Solve Problems that Single-target Drugs Can Not

Generative Models



Sequence Space



Geometry-dependent proximity is critical

Mid/Low TAA density

12 – 20 nm

TAA Fab CD3 VH

High TAA density

TAA VH

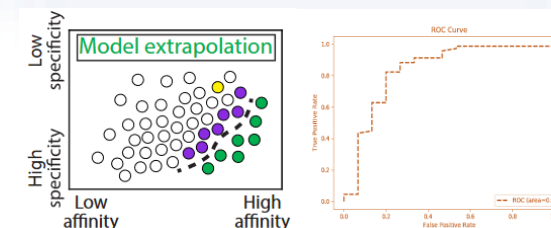
CD3 Fab

Tumor 15 nm T cell

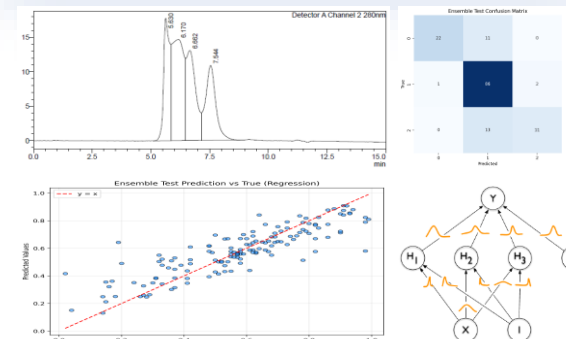
CD3 VH

TAA VH

Predictive Models

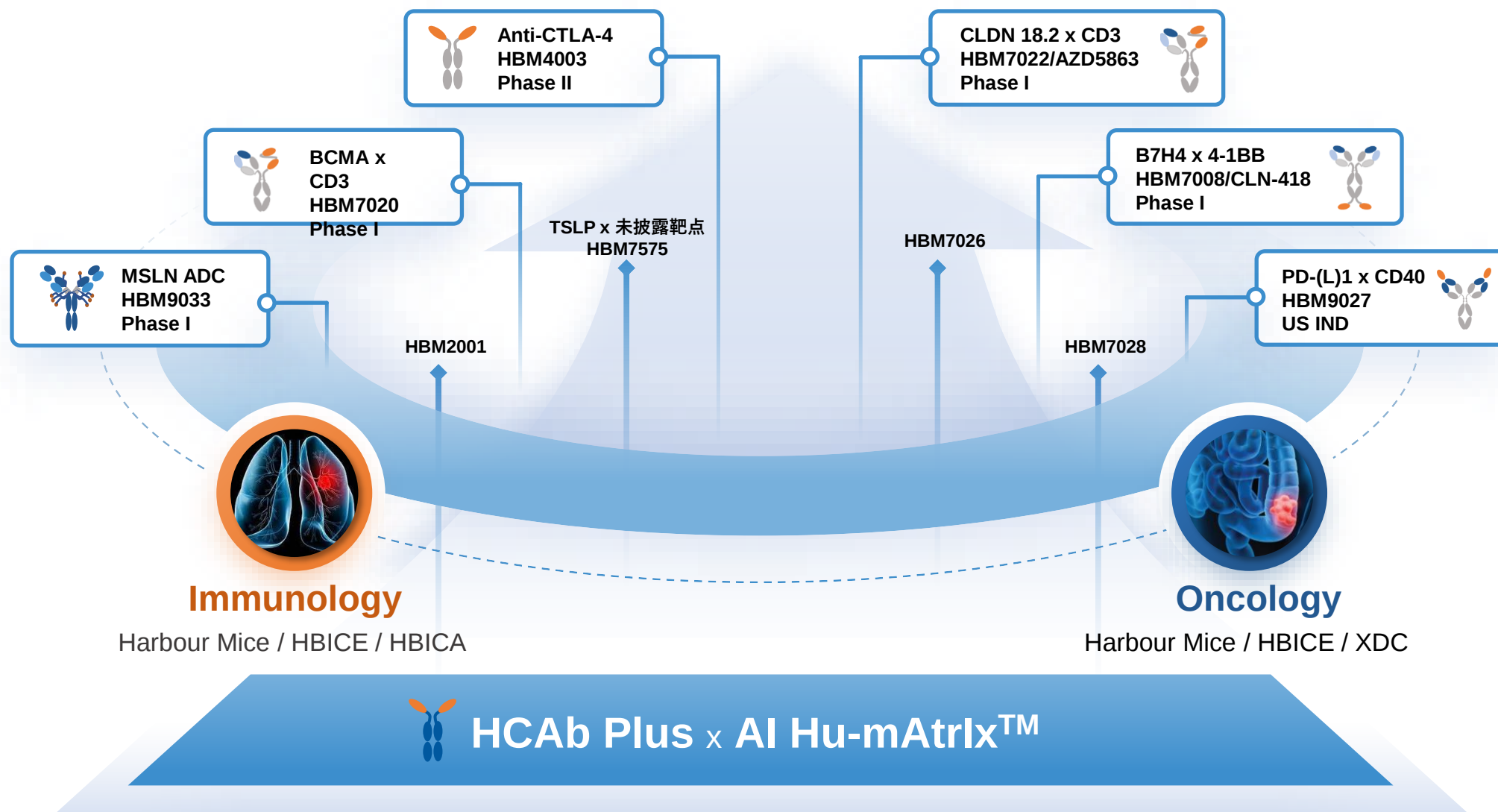


Developability, Thermal Stability, Aggregation, etc.



Platform-Led Pipeline Strategy: HCAb Plus for Next Generation Multispecific Therapeutics

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HBM Therapeutic & Strategy Focus

HBM focuses on 2 key therapeutic areas to address significant unmet medical needs and drive differentiated innovation.

Immunology & Inflammation



Respiratory

- COPD
- Asthma



Dermatology

- Atopic Dermatitis
- Hidradenitis Suppurativa
- Vitiligo
- Alopecia Areata



GI

- Ulcerative Colitis
- Crohn's Disease
- Celiac Disease



Systemic I & I

- SLE
- Multiple Sclerosis
- IgA Nephropathy
- Fibrosis

Science Focus

- **Pioneering Novel Multispecific Antibodies:** Restoring immune balance by targeting complementary pathways across the Th1/Th2/Th17 axis and key regulatory circuits (e.g., Treg, IL-2, TGF- β).
- **Resetting the Immune System with HBICE™:** Enabling superior safety and efficacy in autoimmune diseases through deep, specific, and durable depletion of pathogenic cells.

Abbr.

GI – Gastrointestinal; SLE – Systemic Lupus Erythematosus; DKD – Diabetic Kidney Disease; COPD – Chronic Obstructive Pulmonary Disease; MAFLD – Metabolic Dysfunction-Associated Fatty Liver Disease; MASH – Metabolic Dysfunction-Associated Steatohepatitis

Oncology



Solid Tumor

- Colorectal Cancer
- Pancreatic cancer
- Gynecology cancer
- Lung cancer

Science Focus

- **Conquering Immunologically Cold Tumors:** Targeting tumors with low immune cell infiltration and inherent resistance to existing checkpoint inhibitor therapies.
- **Deploying a Diversified Modality Toolkit:** Advancing novel platform, including XDC, multispecific antibodies, cytokine fusion, and combination immunotherapies.
- **Improving Overall Survival in Refractory Cancers:** Integrating direct tumor killing with sophisticated tumor microenvironment (TME) modulation and renormalization strategies to extend patient survival.

Immunology: Address Complementary Mechanisms of Action in I&I Diseases by Multispecific Antibodies

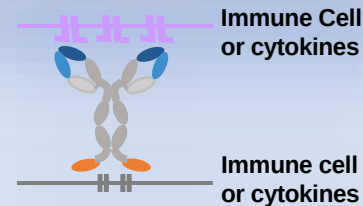
Immunology diseases are multifactorial, strongly heterogenous disorders driven by distinct inflammatory pathways

Blocking multiple pathways in Type 2 (T2)-low endotypes of asthma/AD



HBICA

HCAb-base Bispecific Immune Cell Antagonist

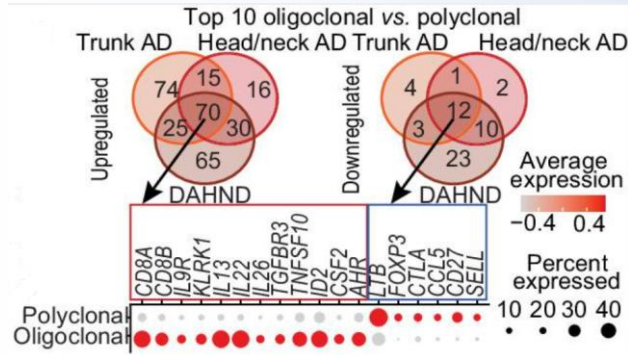
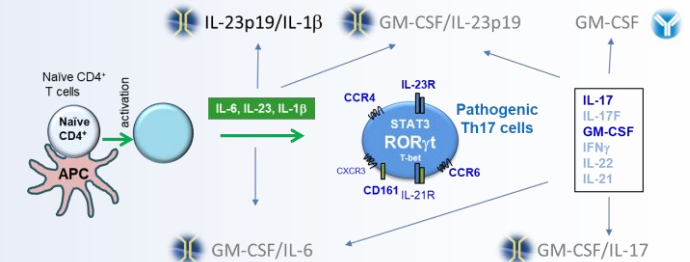
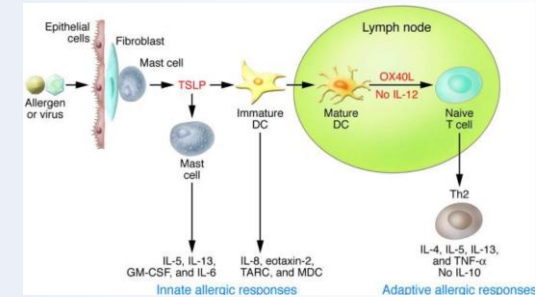


2+2 symmetric HBICA



Dual blocking immune modulator

- Fully human aa seq
- Intact Fc with HLE
- Bivalent blocking
- favorable developability



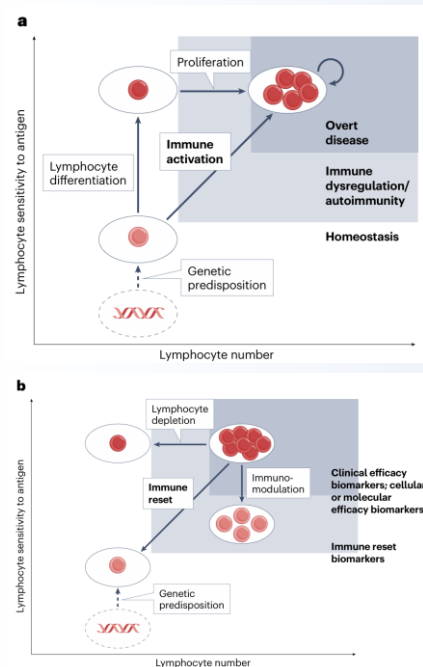
Nat Commun. 2024; 15, 2839

Immunology: Achieve Immune Reset by Pathogenic Immune Cell Depletion Mediated by HBICE/HBIME

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Working model of immune reset

Achieve deeper and longer depletion in tissue
B or LLPC



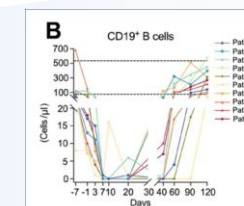
Nat Rev Immunol 25, 528–541

HBICE
HCAb-based Bispecific Immune Cell Engager
HCAb-based Bispecific Myeloid Cell Engager

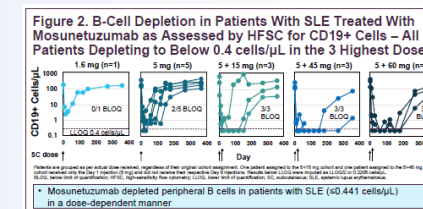
Immune Cell Pathogenic Cell

N+1 asymmetric HBICE/HBIME

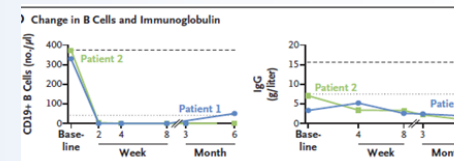
B/DC/T cell depletion therapy



Wang W, et al. Ann Rheum Dis. 2024



EULAR-2025



Zhao J, et al. NEJM. 2025

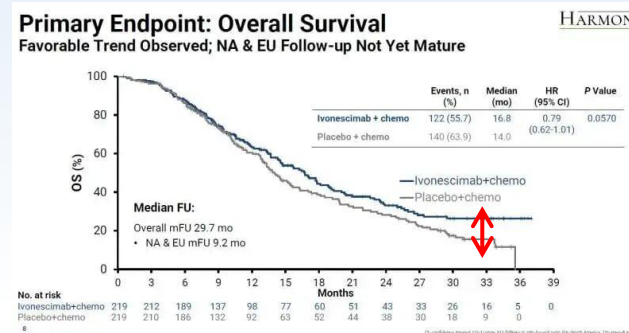
Oncology: Discover Next-Gen Multispecific PD-(L)1 Plus for Solid Tumor Patients

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Improving Therapeutic Window by Modulating IO, Angiogenesis and Stroma of Cancers with Cornerstone PD-(L)1 Plus Therapies

- ✓ mAb → bsAb → **Multispecific Medicines**
- ✓ **Improve OS** as ultimate endpoint
- ✓ Better **safety** for combination to beat SOC



Dr Jonathan Goldman, IASLC

**HBM4003 CTLA-4
Combo with PD-1**

MSS CRC Ph2 Positive Readouts

**HBM9027 PD-L1 x CD40
DC Cell Engager**

IND US/CN Approved

**PD-1/VEGF/undisclosed 1
Trispecific Antibody**

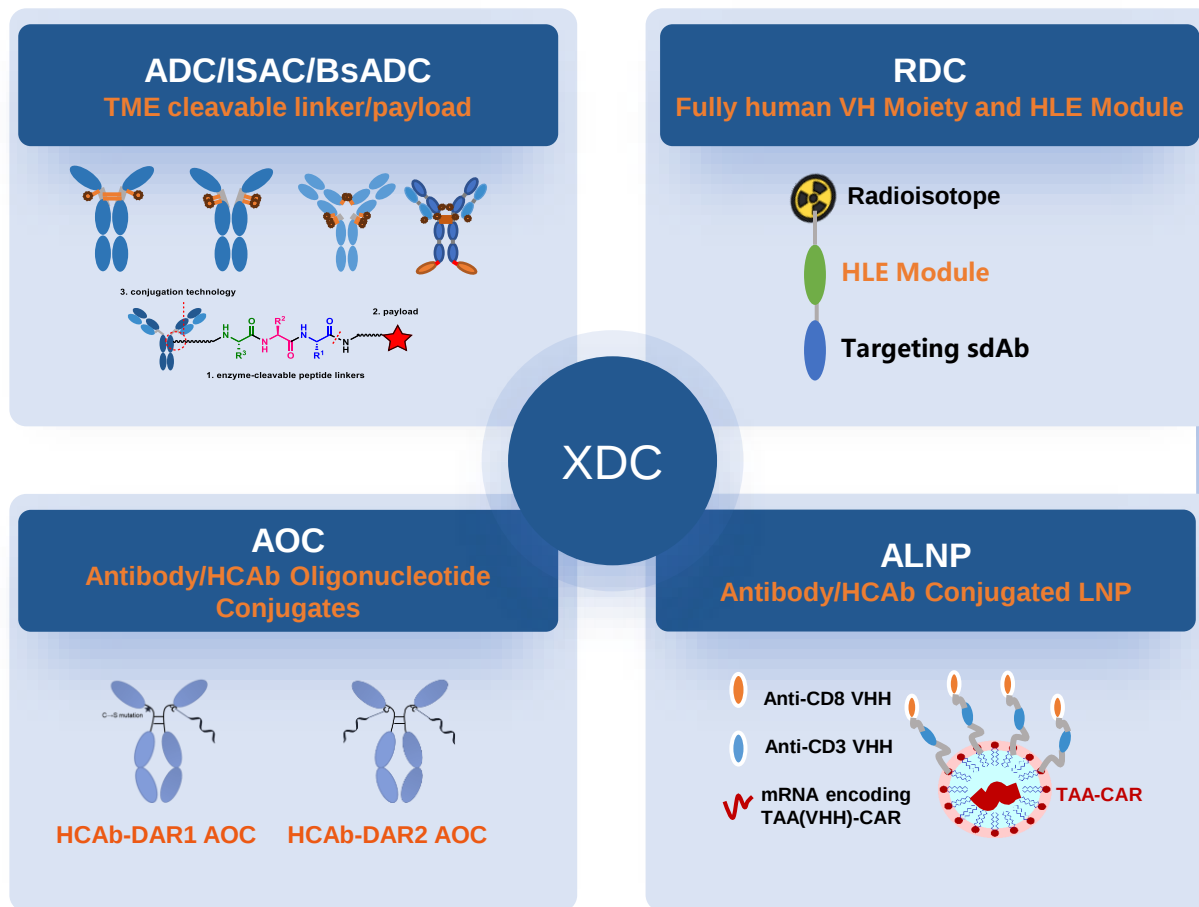
IND 2026

**PD-(L)1/VEGF/
undisclosed 2
Trispecific Antibody**

IND 2026-2027

Oncology: Improve XDC for Cold Tumors with Stroma Barrier

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Ongoing Approaches

1. **ADC:** Tumor microenvironment specific cleavable linker payload for soluble and stroma targets in PDAC and HCC etc.
2. Explore novel targets for CRC/NSCLC ADC or Bispecific ADC
3. **AOC:** Target intracellular oncogenes with specific cell delivery
4. **Ab-LNP:** Tissue specific delivery of drug to improve therapeutic window



Lead Candidates



HBM9033
MSLN ADC
Ph1

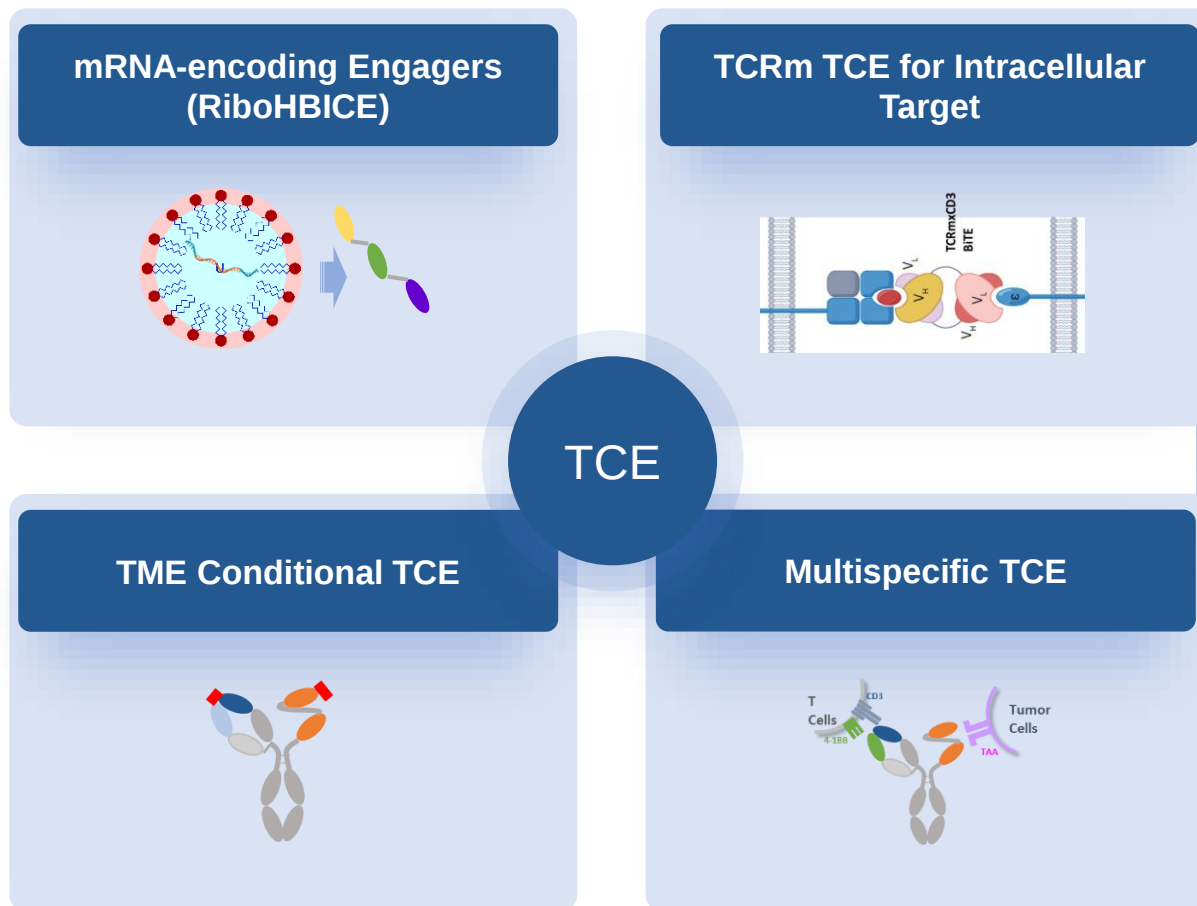


FIC ADC



BsADC

Oncology: Nextgen TCE for Solid Tumor Mono- and Combo-Therapies

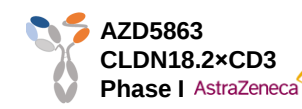
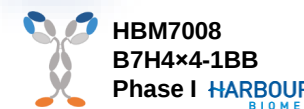


Ongoing Approaches

1. Overcome TAA escape resulted resistance and/or T cell exhaustion due to lack of signal 2/3
2. Achieve multi-in-one function by mRNA encoding RiboHBICE
3. Access intracellular targets (e.g. neoantigens) by TCRm TCE
4. Improve safety window by conditional activated TCE
5. Explore novel targets and combination therapy for TCE/MCE
6. Novel engineering to control specific activation, on-off modulation and geometry dependent cytotoxic killing

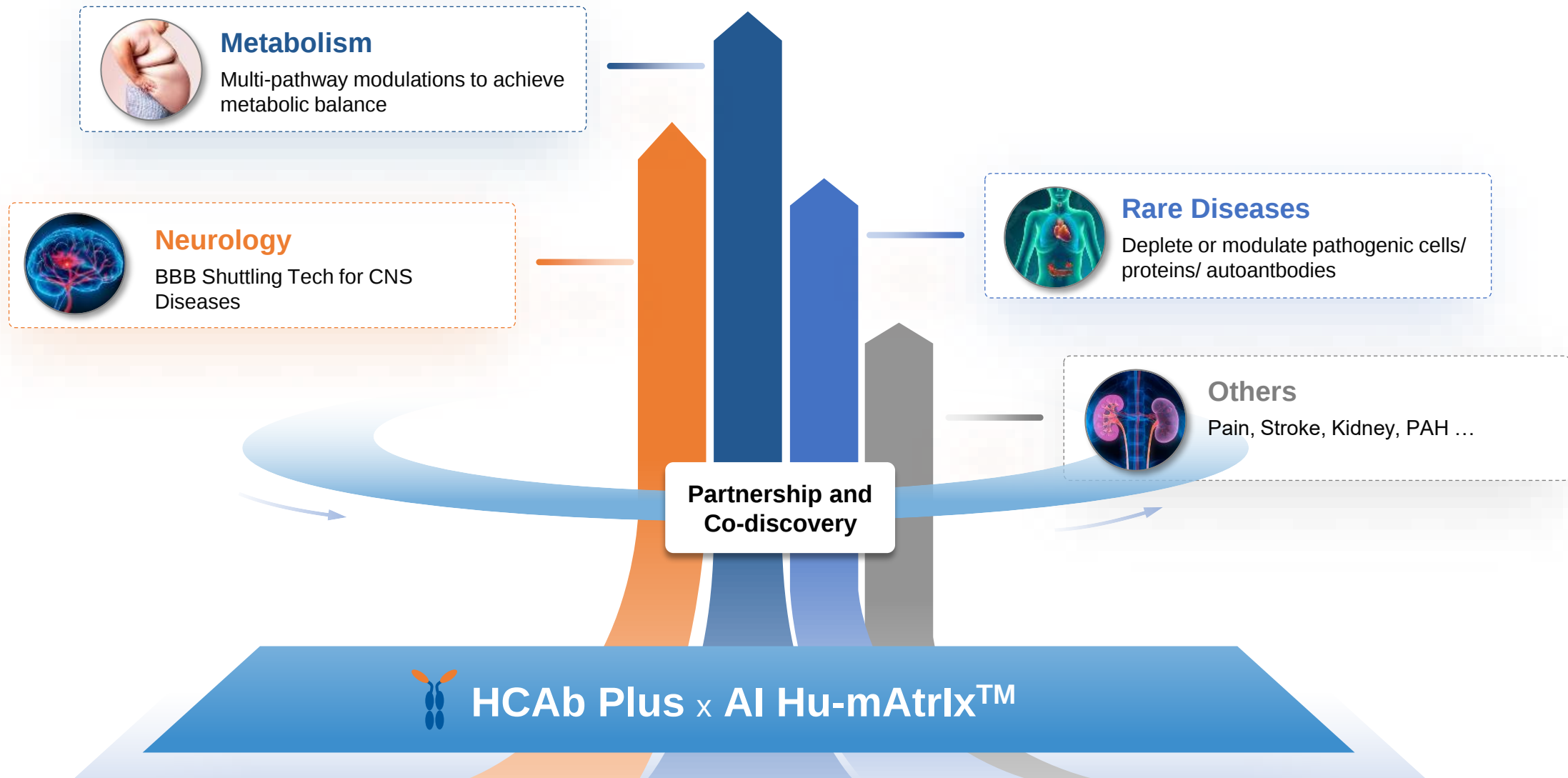


Lead Candidates



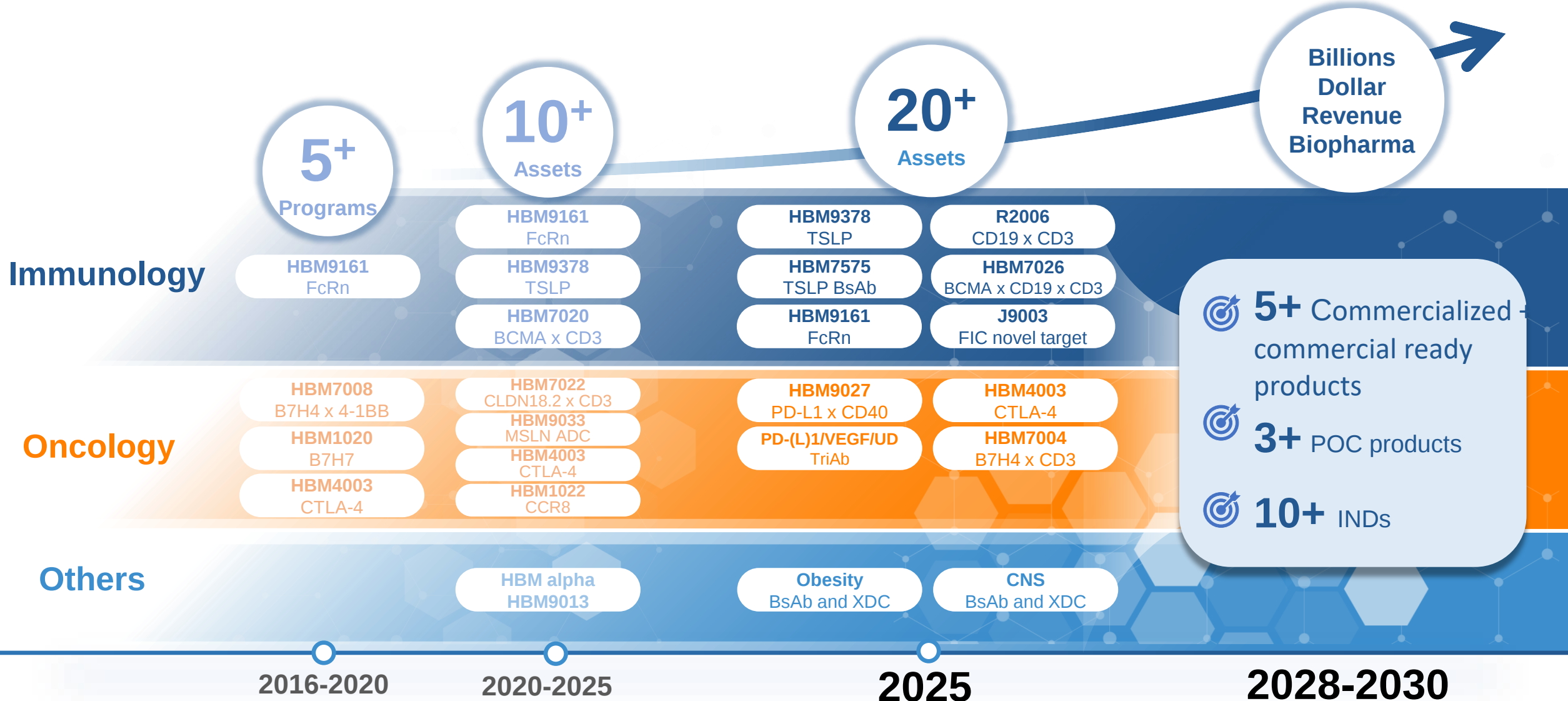
Platform Driven Approach for Other Therapeutic Areas by Open Innovation

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From Platform Biotech to Product Development and Commercial Company

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Thank You