LIANBIO

January 2023

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Investment Highlights

We are a global biopharmaceutical company dedicated to developing and commercializing paradigm-shifting medicines for patients with unmet medical needs in Greater China and other Asian markets

Bringing a pipeline of innovative therapies into the rapidly growing Greater China market

Established pharmaceutical in-licensing and development platform well positioned to capitalize on positive market trends and momentum

Multiple near-term catalysts across a diverse late, mid and early-stage pipeline
Five clinically validated therapeutic candidates, nine in-licensed assets

Experienced cross-border team with BD, alliance management, clinical development, regulatory and commercial expertise and track record

Key validating and differentiating partnerships with Pfizer and BridgeBio

Strong financial position with cash runway through the end of 2024; cash balance[1] of $331.8 million as of September 30, 2022

1. Which includes cash, cash equivalents and marketable securities

China is the Second Largest Pharmaceutical Market Today, with Innovation Agenda Propelling Strong Growth

Substantial unmet medical needs persist in China

- **Aging population > 1.4Bn, with a high disease burden** compared to developed countries[1]
- **"Healthy China 2030" sets clear healthcare industry KPIs** from the government[2]
  - Improve key therapeutic area mortality rates, including CV and oncology
- Despite increased R&D activity, still **few China-originated first-in-class and best-in-class drugs approved**

Fostering innovation: continued momentum in policy and industry evolution

Comprehensive policies enacted to foster innovation

- China's five-year plan includes innovation priorities in TAs such as oncology and CV[3]
- Accelerated review and approval timelines of patented pharmaceuticals[4]

Expanding coverage and broadening access for innovative drugs

- Growth in basic medical insurance and commercial health insurance[5]
- NRDL now updated annually

Biotech ecosystem growth

- Improving capital markets and fund flows into Chinese biotech
- Increase in number of CROs, bioparks, biotechs, clinical trial centers

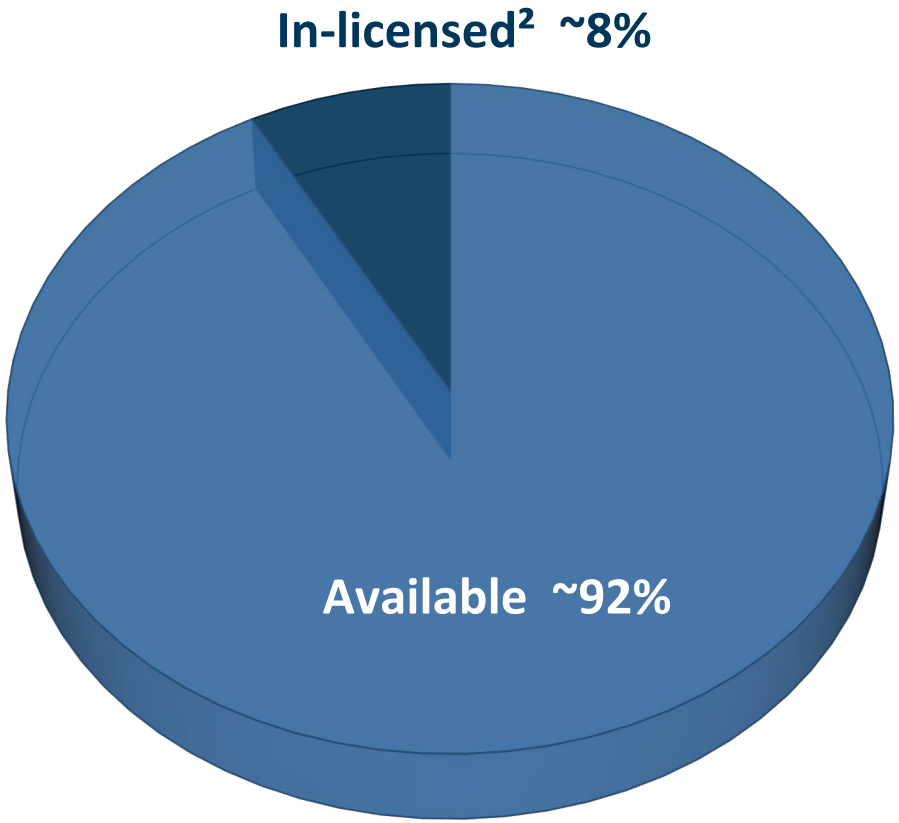
Healthcare infrastructure upgrades

- Upgrades to private and public hospitals and community health centers[6]
- Increasing number of healthcare professionals

1. GBD Global Healthdata Exchange 2019; 2. "Healthy China 2030" released by China State Council in July 2019; 3. "14th Five-Year Plan (2021-2025) and the Long-Range Objectives Through the Year 2035; 4. GBI; review time calculated as time interval between NDA submission date and approval date 5. MoHRSS; NRCMS; CIRC website; China Insurance Yearbook; Xrate 6.53 RMB/USD; 6. "Comprehensive Reform of Public Hospitals Notification of Subsidy Fund Budget" released by Ministry of Finance in Nov 2020

China Biotech Sector at Potential Inflection Point for Significant Growth…

Potential U.S./EU Biotech In-licensing Opportunities for China[1]



Early Innings:

- **< 10% of western innovative biotech medicines tapped for China**, and majority of in-licensed programs are concentrated in oncology
- Western biotechs seeking **strategic access to China as part of global enrollment acceleration** and commercial opportunity

1. CapitalIQ. assumes only one opportunity per company based on 10,794 total US/EU biotech companies as of July 2021. 2.Based on 857 cross-border deals from 2015-2020 per ChinaBio.

…LianBio Positioned to Potentially Lead Next Generation of Chinese Biotechs: Speed, Scale and Sustainable Growth

Differentiated Access to Innovation

- **Relationship with our founder** provides expanded BD opportunities, with unparalleled sourcing, access and clinical/scientific due diligence capabilities

PERCEPTIVE ADVISORS

- BD approach informed by
  - Deep scientific expertise
  - Region-specific development insights
  - Regulatory and commercial insights

Cross-Border Execution Platform

- Management team with deep experience and **proven track records across global and Chinese biopharma** companies
- Robust asset and alliance management with bilingual **U.S.-based team dedicated to alliance management**
- **Maximizing asset value** locally and globally through **bespoke development strategies**
  - Ability to facilitate potentially faster market entry through **bridging studies and accelerated pathways**
  - **Unique in-market indications** and combination strategies for **global-first expansion** studies

Commercial Model Provides Optionality

- **Integrated commercial infrastructure** built around **core therapeutic areas**, products and market segments
- **Optionality** to leverage commercial partnerships for **broad access** to select assets

Pfizer

- Commercialization strategies **beyond hospital channels** provide broadened opportunities

Pipeline of Innovative Medicines – 5 Clinically Validated Therapeutic Candidates

Global Development Status[1] — Clinically Validated

Therapeutic Area	Program	Indication	Phase 1	Phase 2	Phase 3/ Pivotal	Approved	Next step in China	Partner
Cardiovascular	Mavacamten[2]	Obstructive Hypertrophic Cardiomyopathy (oHCM)					China Phase 3 trial ongoing, enrollment completed August 2022	Bristol Myers Squibb
		Non-obstructive Hypertrophic Cardiomyopathy (nHCM)					Conduct registration enabling trial	
		Heart Failure with Preserved Ejection Fraction (HFpEF)					Conduct registration enabling trial	
Ophthalmology	TP-03	Demodex Blepharitis					China Phase 3 trial ongoing	tarsus
		Meibomian Gland Disease					Join future global Phase 3 trial	
Oncology	NBTXR3[3]	Head and Neck Squamous Cell Carcinoma (HNSCC)[2]					Global NANORAY-312 Phase 3 trial ongoing in China	NANOBIOTIX
		Solid Tumor IO Combinations					Join future global Phase 3 trial	
	Infigratinib[4]	Second-line Cholangiocarcinoma w/ FGFR2 Fusions					Approved in Bo'ao region through early access program	bridgebio
		Gastric Cancer w/ FGFR2 Fusions and other FGFR-Driven Tumors[5]					China Phase 2a proof of concept trial ongoing	QED therapeutics
	BBP-398	Advanced Solid Tumors					China Phase 1 monotherapy trial ongoing	navire, bridgebio
		Non-Small Cell Lung Cancer (NSCLC)					Conduct China Phase 1 Osimertinib combo trial	Bristol Myers Squibb
Inflammatory Disease	Omilancor	Ulcerative Colitis					Join potential future global Phase 3 trial	LANDOS BIOPHARMA
	NX-13	Ulcerative Colitis					Join potential future global Phase 3 trial	
	LYR-210	Chronic Rhinosinusitis (CRS)					Conduct China standalone Phase 3 trial	LYRA THERAPEUTICS

1. The commercialization of each of our product candidates will require regulatory approval in the respective jurisdiction in which we intend to market such product candidate; however, obtaining and maintaining regulatory approval in one jurisdiction does not guarantee we will be successful in obtaining or maintaining regulatory approval of the product candidate in other jurisdictions that are material to the success of LianBio. 2. Mavacamten has received FDA approval in the US, which is not a part of our licensed territory, for the treatment of NYHA class II-III obstructive HCM 3. NBTXR3 has received European market approval (CE mark) in the EU, which is not a part of our licensed territory, for the treatment of locally advanced soft tissue sarcoma. At present, we are not pursuing NBTXR3 in relation to this STS indication. 4. Infigratinib has received FDA approval in the US, which is not a part of our licensed territory, for the treatment of previously treated, unresectable locally advanced or metastatic cholangiocarcinoma with FGFR2 fusion or other rearrangement. 5. Ongoing Phase 2a gastric cancer and other FGFR-driven tumor standalone clinical trial in China. Separate investigator sponsored Phase 2 clinical trial of infigratinib in FGFR-driven tumors is ongoing in the United States.

Three Key Pillars for Patient Reach and Sustainable Growth

#1 Establish Commercial Footprint with Lead Assets

*** All clinically validated***

- CAMZYOS (mavacamten) capsules — Mavacamten
- TP-03
- NBTXR3
- TRUSELTIQ (infigratinib) capsules — Infigratinib

#2 Leverage Infrastructure

- BBP-398
- LYR-210
- Omilancor
- NX-13

#3 Expand Pipeline via Additional BD

- Deepen existing TA franchises
- Strategic multi-asset partnerships
- Combination opportunities

Oncology | Cardiovascular | Ophthalmology | Inflammatory Diseases

LianBio has Rapidly Established a Platform to Serve as a Partner of Choice and Gateway to China

9 assets in-licensed, 6 clinical trials ongoing

2019

- Initial investment from Perceptive Advisors
- Global strategic partnership with BridgeBio
- In-licensing agreement with QED Therapeutics

2020

- Announced mavacamten and BBP-398 partnerships
- Closed $310m Series A financing
- Entered into strategic collaboration with Pfizer

2021

- In-licensing agreement with ReViral Therapeutics
- In-licensing agreement with Tarsus Pharmaceuticals
- In-licensing agreement with Nanobiotix
- In-licensing agreement with Landos Biopharma
- Initiated infigratinib P2a gastric cancer trial in China
- In-licensing agreement with Lyra Therapeutics
- Completed $325m Nasdaq IPO
- Initiated and completed dosing in PK study of mavacamten in Chinese healthy volunteers
- Infigratinib approved for 2L CCA in Bo'ao region

2022

- Initiated P3 EXPLORER-CN trial of mavacamten in China
- Mavacamten granted Breakthrough Therapy Designation for oHCM in China
- Formed Scientific Advisory Board
- Submitted mavacamten NDA in Singapore
- Submitted infigratinib NDA in Hong Kong
- Completed enrollment in P3 EXPLORER-CN trial of mavacamten in China
- Initiated global P3 NANORAY-312 trial of NBTXR3 in LianBio territories
- Submitted mavacamten NDA in Hong Kong
- Initiated P3 LIBRA trial of TP-03 in China
- Initiated P1 monotherapy trial of BBP-398 in China
- Pfizer opted into LianBio rights to sisunatovir

Collaboration Agreement | Financial/Corporate Milestone | Clinical Development Milestone | Regulatory Milestone

Experienced Cross-Border Management Team Supported by a Highly Regarded Board of Directors

Management Team

Yizhe Wang, Ph.D.
Chief Executive Officer; Board Member

Yi Larson
Chief Financial Officer

Pascal Qian
China General Manager

Michael Humphries
Chief Scientific Advisor

Brianne Jahn
Chief Business Officer

Nathan Chen
VP, Regulatory Affairs, Pharmacovigilance and Project Management

Levvy Lv, D. Eng
VP, Clinical Operations & Translational Development

Board of Directors

Konstantin Poukalov
Managing Director – Strategy, Perceptive Advisors; Executive Chairman, LianBio

Yizhe Wang, Ph.D.
Chief Executive Officer, LianBio

Adam Stone
Chief Investment Officer, Perceptive Advisors

Tassos Gianakakos
Former Chief Executive Officer, MyoKardia

Neil Kumar, Ph.D.
Chief Executive Officer, BridgeBio

Susan Silbermann
Former Global President, Emerging Markets, Pfizer

Wei Wei Chen
Former Vice President, Chief Financial Officer, Starbucks China

Significant Commercial Leadership Experience Across Diverse and Relevant Therapeutic Areas, Including Global and China Launch Execution

Select commercialization experience

Cardiometabolic

- Entresto™ (sacubitril/valsartan) tablets 24/26mg • 49/51mg • 97/103mg
- Plavix Clopidogrel 75mg/300mg
- Lipitor atorvastatin calcium tablets
- Norvasc Amlodipin
- Fraxiparine®
- Avandia rosiglitazone maleate
- Betaloc
- Volibris ambrisentan
- Pradaxa

Oncology

- 达伯舒 信迪利单抗注射液 Sintilimab Injection
- Verzenio abemaciclib
- AFINITOR (everolimus) tablets
- Votrient™ 200 mg
- SUTENT 50 mg SUNITINIB Zytostatikum
- GEMZAR (gemcitabine)
- Arzerra® (ofatumumab) Injection, for intravenous infusion 20 mg/mL
- ALIMTA pemetrexed
- 爱优特® 呋喹替尼胶囊

Inflammatory Disease

- olumiant (baricitinib) tablets 2 mg
- taltz (ixekizumab) injection 80 mg/mL
- Benlysta (belimumab) Intravenous Use 120 mg/vial Subcutaneous Use 200 mg/mL

Ophthalmology & Other

- Xalatan® latanoprost ophthalmic solution
- Latanoprost
- ADVAIR DISKUS®
- Pulmicort®
- ZYPREXA IntraMuscular Olanzapine for Injection
- Cialis® (tadalafil) tablets 2.5 mg, 5 mg, 10 mg, 20 mg
- SINGULAIR 10mg Tablets montelukast sodium
- Cymbalta® duloxetine HCl
- TIENAM®
- RELVAR ELLIPTA fluticasone furoate/vilanterol
- Viread® 300mg tablets tenofovir disoproxil fumarate
- ANORO® ELLIPTA® umeclidinium/vilanterol
- strattera™ atomoxetine HCl
- EPIVIR (lamivudine) ORAL SOLUTION 10 mg/mL
- VFEND®
- Ventolin® (albuterol sulfate)

Pipeline

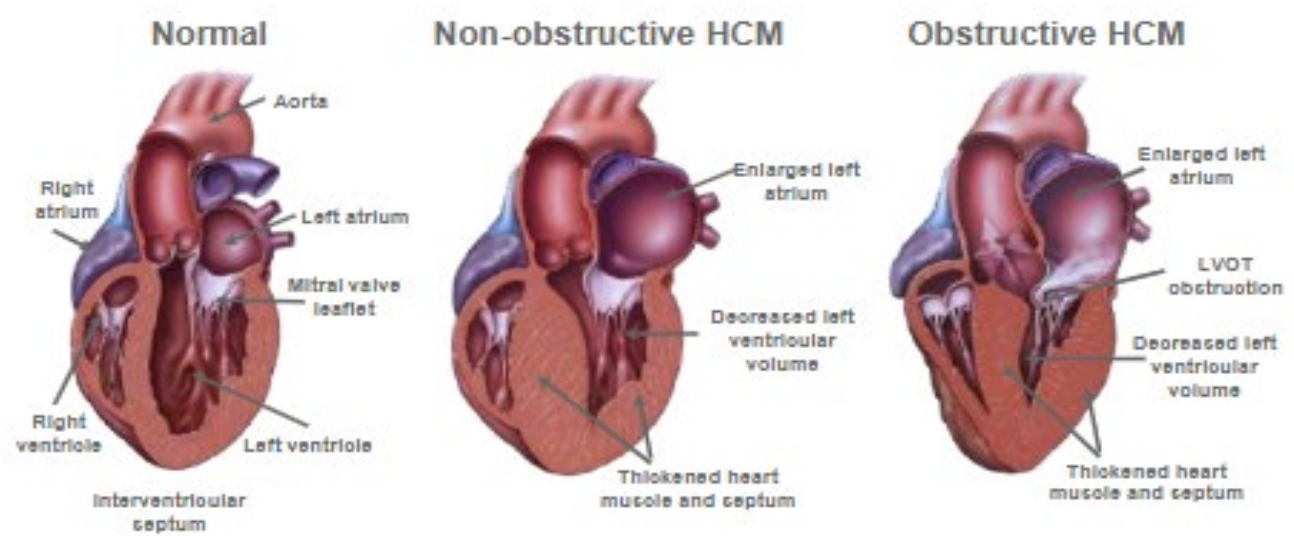
Mavacamten for the Treatment of HCM and HFpEF

- Mavacamten is a **myosin inhibitor** that targets excessive contractility and impaired relaxation, myocardial energetics and compliance
- In development for the treatment of obstructive hypertrophic cardiomyopathy (**oHCM**), non-obstructive hypertrophic cardiomyopathy (**nHCM**) and heart failure with preserved ejection fraction (**HFpEF**)

China Opportunity

- **1.1M - 2.8M HCM** patients in China (67% oHCM / 33% nHCM)
- **3.7M HFpEF** patients, 10-20% of whom may potentially be addressed by mavacamten

Hypertrophic Cardiomyopathy



- **Obstructive HCM (oHCM)**: Characterized by dynamic LV outflow tract obstruction, in which the enlarged and diseased muscle blocks the flow of blood from the left ventricle to the rest of the body.
- **Non-Obstructive HCM (nHCM):** No significant LV outflow tract obstruction (<30 mm Hg) at rest or with provocation. Driven by diastolic impairment due to the enlarged and stiffened heart muscle.

Clinical Activity Demonstrated in oHCM and nHCM

Clinical Data Summary

oHCM:

- Phase 3 EXPLORER-HCM trial demonstrated patients on treatment experienced statistically significant and clinically meaningful improvements
  - Primary endpoint: Improvement of symptoms and functional capacity (improvement in NYHA class and peak VO2)
  - Well-tolerated; safety results were comparable to placebo; only 2% drop out rate

nHCM:

- Phase 2 MAVERICK-HCM trial demonstrated physiologic benefit with dose dependent reduction in serum levels of NT proBNP, with potentially greater benefit in more severe disease

EXPLORER-HCM

Change from Baseline to Week 30			
	Mavacamten (n=123)	Placebo (n=128)	P-value
Primary Endpoint			
Composite functional, n (%) **EITHER** ≥1.5 ml/kg/min increase in pVO2 with ≥1 NYHA class improvement **OR** ≥3.0 ml/kg/min increase in pVO2 with no worsening of NYHA class	45 (37%)	22 (17%)	**0.0005**
Secondary Endpoints			
Post-exercise LVOT peak gradient, mmHg, mean (SD)	-47 (40)	-10 (30)	**<0.0001**
Peak VO2, mL/kg/min, mean (SD)	1.4 (3.1)	-0.1 (3.0)	**0.0006**
NYHA improved ≥ 1 class, n (%)	80 (65%)	40 (31%)	**<0.0001**
KCCQ-CSS, mean (SD)	13.6 (14.4)	4.2 (13.7)	**<0.0001**
HCMSQ-SoB score, mean (SD)	-2.8 (2.7)	-0.9 (2.4)	**<0.0001**

Source: Olivotto et al, Lancet 2020; Ho et al, J Am Coll Cardiol. 2020

Mavacamten Registration Pathway

Global Stage of Development

- **oHCM:** BMS received approval in the U.S. for the treatment of symptomatic NYHA Class II-III oHCM to improve functional capacity and symptoms
  - BMS presented additional supportive data April 2022:
    - Phase 3 VALOR-HCM study demonstrated mavacamten significantly reduced the need for septal reduction therapy (SRT) in patients with severely symptomatic oHCM who had been appropriate for SRT
    - Phase 3 EXPLORER-LTE study demonstrated sustained improvements in clinically meaningful CV outcomes at weeks 48 and 84
- **nHCM:** MyoKardia completed Phase 2 double-blind, placebo-controlled MAVERICK trial in symptomatic nHCM patients; BMS to initiate Phase 3 nHCM trial in 2022
- **HFpEF:** BMS initiated a Phase 2 trial of mavacamten in HFpEF in Feb 2021

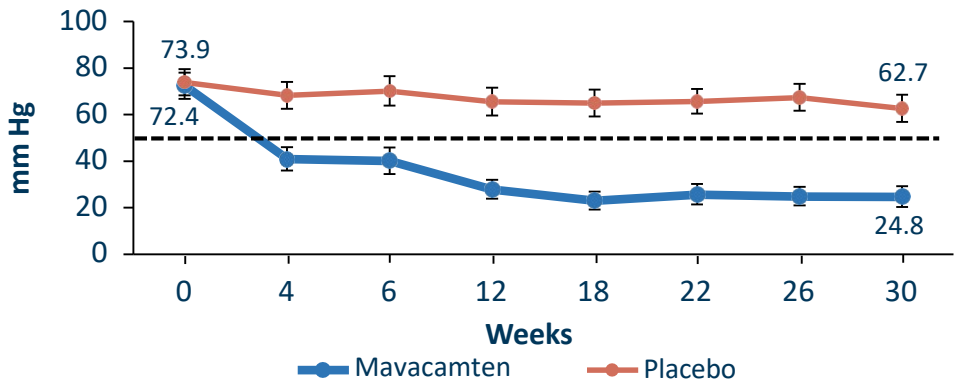
China Development Plan

oHCM: P3 EXPLORER-CN standalone trial enrollment completed August 2022, topline data anticipated mid-2023; PK trial complete

- EXPLORER-CN design mimics EXPLORER-HCM, with some changes to account for **China-specific considerations**
  - Primary endpoint: Valsalva LVOTg
  - Secondary endpoints: resting LVOTg, NYHA and KCCQ
  - EXPLORER-CN initiated January 2022
- PK study complete, favorable tolerability & PK profile demonstrated
- Breakthrough Therapy Designation granted in China February 2022

EXPLORER-HCM

Mean (95% CI) Valsalva LVOT gradient



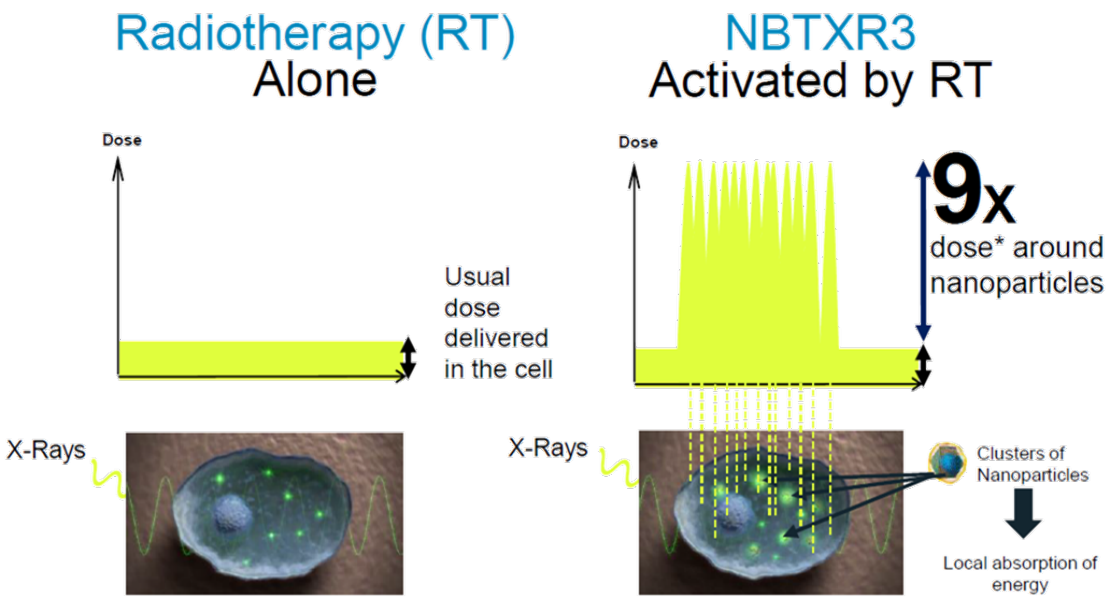
Number of patients at visit

Mavacamten	123	117	118	118	116	118	120	117
Placebo	128	119	119	125	122	125	124	124

NBTXR3 for the Treatment of Solid Tumors

NBTXR3 is a radioenhancer designed to enhance the efficacy of radiotherapy without resulting in additional side effects on surrounding healthy tissue

Illustrative example of NBTXR3 activity



*Dose enhancement determined by Monte Carlo simulation (CEA Saclay, France). In vitro data. Nanobiotix data.

China Opportunity

- **1.3M patients** receive radiation therapy annually as part of their cancer treatment[1]
- **Up to 925K** patients across potential target indications
  - Locally advanced head and neck cancer: ~25K
  - Non-IO potential solid tumor indications: ~150K
  - IO combination potential solid tumor indications: ~750K

1. Based on 2018 data

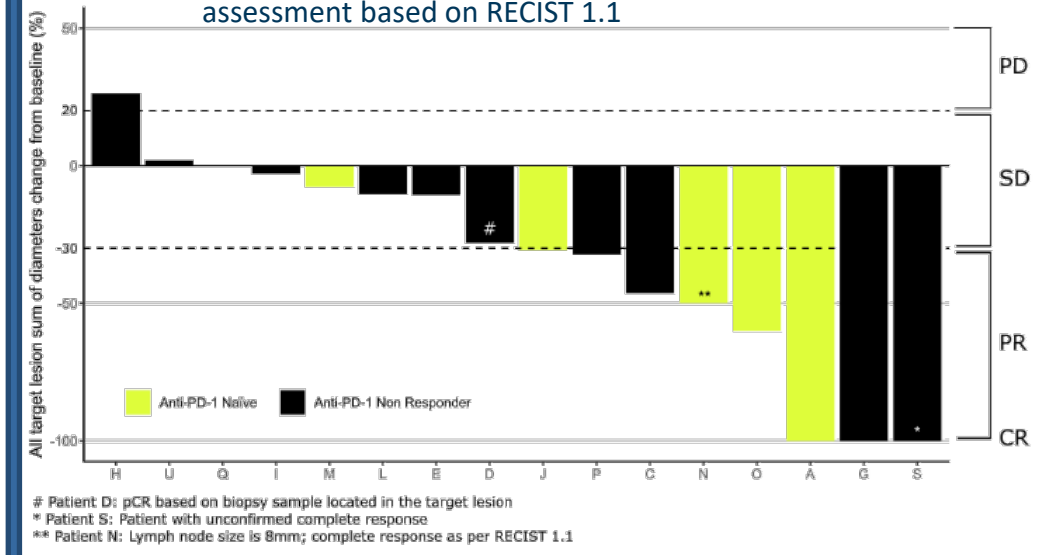
Broadly Applicable MOA: Clinical Proof of Concept Demonstrated Across Multiple Tumors and IO Combination

Nanobiotix Key Clinical Data

- **NBTXR3 + RT in soft tissue sarcoma**
  - CE mark approval in EU based on Phase 3 study showing 16.1% CRR w/ NBTXR3 +RT vs. 7.9% CRR w/ RT alone
- **P1 Expansion Study 102: NBTXR3 + RT in locally advanced head and neck cancer (n=41 evaluable patients)**
  - 85.4% ORR
  - 63.4% CRR
  - mOS 18.1 months
  - mPFS 10.6 months
- **P1 Study 1100: NBTXR3 + anti-PD-1+ RT in patients with HNSCC, lung metastases and liver metastases (n=16 evaluable patients)**
  - PD-1 naïve ORR: 80% target lesion
  - PD-1 prior non-responder ORR: 45% target lesion
  - Target lesion disease control rate: 94%

Study 1100: Best Observed Target Lesion Response (Evaluable Population n=16)

Best observed target lesion response as per investigator assessment based on RECIST 1.1



China Development Strategy

- Enrollment ongoing in LianBio territories in global Phase 3 NANORAY-312 trial of NBTXR3 in **in locally advanced HNSCC**
- Additional future trials to include IO combination approaches

TP-03 for the Treatment of Demodex Blepharitis (DB) and Meibomian Gland Disease (MGD)

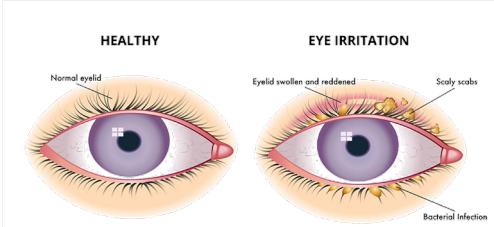
TP-03 (lotilaner ophthalmic solution) is a GABA-Cl channel blocker in development for the treatment of Demodex blepharitis (DB), meibomian gland disease (MGD)

China Opportunity

- **43M DB** patients
- **73M Demodex-driven MGD** patients
  - ~50% of Demodex-driven MGD patients also have DB

Target Indications

Demodex Blepharitis (DB)

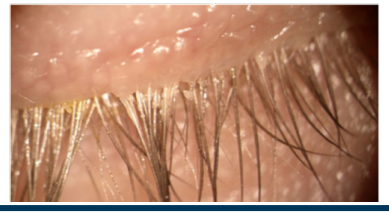


- Blepharitis is characterized by eye inflammation, burning, and tearing, and may be accompanied by a specific type of debris called “collarettes”
- A significant proportion of blepharitis cases are caused by eyelash follicle infestation by the Demodex parasite

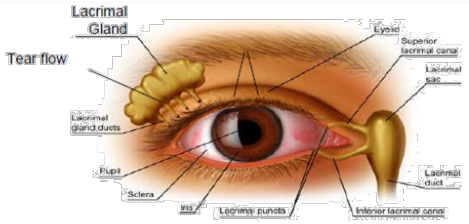
Collarettes Are Pathognomonic Sign of Demodex Infestation

Collarettes Are Composed of Mite Waste Products and Eggs[1]

- Regurgitated undigested material combined with epithelial cells, keratin, and mite eggs
- Contain digestive enzymes, which cause irritation



Meibomian Gland Disease (MGD)



- Common eye condition where the glands do not secrete enough oil or when the oil they secrete is of poor quality
- If left untreated, MGD can cause or exacerbate dry eye symptoms and eyelid inflammation
- Symptoms include dryness, burning, itching, stickiness/ crustiness, watering, light sensitivity, red eyes, foreign body sensation

All Pre-Specified Primary and Secondary Endpoints were Met in Tarsus's Saturn-1 and Saturn-2 Pivotal Trials

Tarsus completed two successful pivotal trials with consistency across endpoints

	Saturn-1 (Pivotal Phase 2b/3) N=421	Saturn-2 (Pivotal Phase 3) N=412	Combined Pivotal Data N=833
Primary Endpoint: Complete Collarette Cure	**44%** vs. 7% (p<0.0001)	**56%** vs. 13% (p<0.0001)	**50%** vs. 10%
Clinically Meaningful Collarette Cure (Grade 0 or 1)	**81%** vs. 23% (p<0.0001)	**89%** vs. 33% (p<0.0001)	**85%** vs 28%
Mite Eradication	**68%** vs. 18% (p<0.0001)	**52%** vs 14% (p<0.0001)	**60%** vs 16%
Lid Erythema Cure	**19%** vs. 7% (p<0.0001)	**31%** vs. 9% (p<0.0001)	**25%** vs 8%

Approximately 90% of patients experienced a clinically meaningful benefit with respect to collarettes, collarette grade improvement and mites per lash

Source: Tarsus Pharmaceuticals

Development and Regulatory Status

U.S.

- **Tarsus submitted NDA for TP-03 in DB to U.S. FDA in September 2022**
- Phase 2a MGD trial initiated August 2022

China

- LianBio conducting pivotal study to support regulatory approval in China, data expected Q4 2023
  - PK cohort (n=12)
  - P3 China standalone trial (N=150, 1:1 randomization)
    - Co-primary endpoints: collarette cure (0-2 collarettes per eyelid) at day 43, mite eradication at day 43
    - Secondary endpoints: composite cure of collarette and erythema (0-2 collarettes per eyelid and grade 0 erythema) at day 43

Additional Pipeline Programs

Infigratinib (FGFR-selective TKI) for the treatment of FGFR-driven cancers

- Phase 2a clinical trial in FGFR2-amplified gastric and other FGFR-driven cancers ongoing in China, where gastric cancer has a disproportionately higher prevalence
- Approved in 2L CCA in the US in May 2021

BBP-398 (SHP2 inhibitor) for the treatment of MAPK pathway-driven solid tumors

- Differentiated profile with a shorter-half life, attractive PK/PD and clean tox
- Phase 1 monotherapy clinical trial ongoing in China; Phase 1 trial in combination with EGFR-inhibitor in NSCLC planned in China

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LYR-210 (implantable drug matrix) for the treatment of chronic rhinosinusitis (CRS) with 3.4M medically refractory patients in China

- Implantable drug matrix designed to consistently and locally elute mometasone furoate (steroid) to inflamed mucosal sinus tissue for up to six months with a single administration for surgically naïve patients
- Clinically validated with Ph2 statistically significant symptom improvement vs. control at 16, 20 and 24 weeks

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Omilancor (LANCL2 agonist) for the treatment of IBD

- Oral, gut-restrictive mechanism (lack of systemic exposure) designed for a safe and convenient route of administration for treatment of moderate to severe IBD
- Rapidly growing IBD incident population in China

NX-13 (NLRX1 agonist) for the treatment of IBD

- In Ph1a safety study, NX-13 was shown to be well tolerated

Major Validating Milestones Highlight Strength of LianBio Business Development Engine

Partner	LianBio Partnership Date	Asset Milestone Post-Partnership
MYOKARDIA Mavacamten Bristol Myers Squibb	Aug 2020	✓ **Oct 2020:** MyoKardia **acquired by BMS for $13.1B** ✓ **Apr 2022:** BMS received **U.S. FDA approval** of mavacamten for patients with symptomatic oHCM
REVIRAL Sisunatovir Pfizer	Mar 2021	✓ **Apr 2022:** Reviral enters agreement to be **acquired by Pfizer for up to $525M** ✓ **Dec 2022:** **Pfizer opts in** to LianBio rights to develop and commercialize sisunatovir in mainland China, Hong Kong, Macau, and Singapore
bridgebio QED therapeutics Infigratinib	Oct 2019	✓ **May 2021:** QED received **FDA approval** of infigratinib for patients with previously treated cholangiocarcinoma
tarsus TP-03	Mar 2021	✓ **Jun 2021:** **Positive pivotal results** in Tarsus's SATURN-1 trial (P2b/3 DB) – all primary and secondary endpoints met ✓ **May 2022:** **Positive pivotal results** in Tarsus's SATURN-2 trial (P3 DB) – all primary and secondary endpoints met
bridgebio navire BBP-398 Bristol Myers Squibb	Oct 2019	✓ **May 2022:** BridgeBio and BMS enter into **BBP-398 strategic collaboration**

Strategic Partnerships Provide Optionality with Differentiated Access to Commercial Infrastructure and Pipeline Opportunities

A differentiated strategic collaboration that provides sourcing, development and commercial optionality

- Provides LianBio and partners optionality to access **Pfizer's established commercial infrastructure** with a highly compliant, secure commercial engine
- At LianBio's election and Pfizer's ROFN, we can jointly develop and commercialize certain LianBio products
- Companies are also working together to **source, select and develop/register leading products for China**
  - Pfizer will contribute up to $70M of non-dilutive capital for in-licensing and co-development activities
  - First opted into LianBio rights to develop and commercialize sisunatovir in mainland China, Hong Kong, Macau, and Singapore in December 2022

Preferential access to an innovative pipeline of more than 20 product development candidates

- BridgeBio is developing transformative medicines to treat patients who suffer from genetic diseases and cancers with clear genetic drivers. BridgeBio is **advancing a broad, innovative pipeline** across rare disease, oncology, dermatology, and other indications
- LianBio already holds China rights to two of BridgeBio's oncology assets, **infigratinib** and **BBP-398**

Multiple Clinical and Regulatory Milestones Achieved in 2022

Therapeutic Area	Program	Milestone / Catalyst	
Cardiovascular	Mavacamten	▪ Initiated Phase 3 EXPLORER-CN clinical trial in patients with oHCM	✓ Jan 2022
		▪ Mavacamten granted BTD for oHCM in China	✓ Feb 2022
		▪ *U.S. FDA approval for the treatment of symptomatic oHCM (BMS)*	✓ April 2022
		▪ Completed PK trial in China, demonstrating favorable safety, tolerability and PK profile	✓ May 2022
		▪ Completed enrollment in Phase 3 EXPLORER-CN clinical trial in patients with oHCM	✓ Aug 2022
Ophthalmology	TP-03	▪ *Saturn-2 pivotal trial readout (Tarsus)*	✓ May 2022
		▪ Initiated Phase 3 clinical trial in patients with Demodex blepharitis in China	✓ H2 2022
Oncology	NBTXR3	▪ *Global trial initiation of Phase 3 NANORAY-312 clinical trial in head and neck cancer (Nanobiotix)*	✓ Jan 2022
		▪ Initiated China portion of Phase 3 NANORAY-312 clinical trial in patients with head and neck cancer	✓ H2 2022
	BBP-398	▪ Initiated Phase 1 monotherapy clinical trial in Chinese patients with advanced solid tumors	✓ H2 2022

Partner milestones

Investment Highlights

We are a global biopharmaceutical company dedicated to developing and commercializing paradigm-shifting medicines for patients with unmet medical needs in Greater China and other Asian markets

Bringing a pipeline of innovative therapies into the rapidly growing Greater China market

Established pharmaceutical in-licensing and development platform well positioned to capitalize on positive market trends and momentum

Multiple near-term catalysts across a diverse late, mid and early-stage pipeline
Five clinically validated therapeutic candidates, nine in-licensed assets

Experienced cross-border team with BD, alliance management, clinical development, regulatory and commercial expertise and track record

Key validating and differentiating partnerships with Pfizer and BridgeBio

Strong financial position with cash runway through the end of 2024; cash balance[1] of $331.8 million as of September 30, 2022

1. Which includes cash, cash equivalents and marketable securities