BRINGING THE
PROMISE OF CELL
THERAPY TO LIFE



Chimeric Therapeutics AGM 14 November 2023

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# Bringing the Promise of Cell Therapy to Life

# 2 CLINICAL TRIALS WITH POSITIVE PHASE 1 RESULTS

- Positive CHM 1101 Phase 1A data in recurrent brain cancer
- Positive CHM 0201 Phase 1A data in Acute Myeloid Leukemia and Advanced Colorectal Cancer
- Areas of high unmet medical need

# ADVANCING CLINICAL DEVELOPMENT

- Innovative cell therapy portfolio with 3 novel technology platforms
- 4 ongoing clinical trials in solid tumours and blood cancers
- Multiple clinical catalysts in the next 12-18 months

#### **EXPERIENCE AND EXPERTISE**

- Team of experienced cell therapy experts
- Proven development and commercialization expertise
- Success in business development opportunities

# CHIMERIC: A FOUNDATION FOR LONG TERM SUCCESS 3 NOVEL PLATFORM TECHNOLOGIES

CHM 1101 (CLTX CAR T)

2020

Best in class NK cell platform for blood cancers and solid tumours



CHM 2101 (CDH17 CAR T)

2021

First in class CDH17 CAR T for gastrointestinal cancers



CHM 0201 (CORE NK)

2022

Best in class NK cell platform for blood cancers and solid tumours





# CHM 1101: POSITIVE PHASE 1A CLINICAL DATA IN RECURRENT, LATE-STAGE BRAIN CANCER

DISEASE CONTROL RATE

55%

Disease Control Rate (DCR) in heavily pretreated patients

Exceeding historical disease control rates of 20-37%<sup>1</sup>

**SURVIVAL** 

~10 months

Median survival in patients that achieved disease control

14+ months

Survival in two patients that achieved disease control

~7 month survival expectation after first recurrence<sup>2</sup>

SAFETY

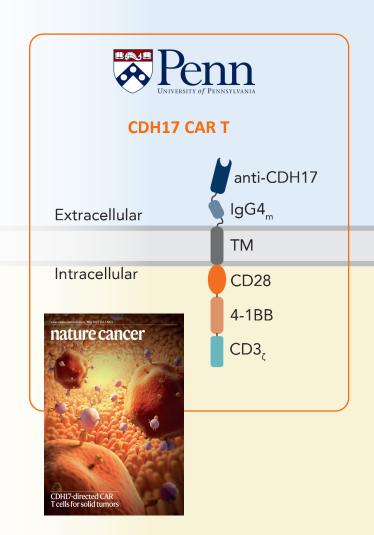
Generally, well tolerated

- No Dose Limiting Toxicities
- No Cytokine Release Syndrome
- No Tumour Lysis Syndrome

### CHM 2101: FDA IND CLEARANCE

#### A FIRST-IN-CLASS 3RD GENERATION CAR T FOR GASTROINTESTINAL TUMOURS

- FDA clearance for Chimeric IND for a Phase 1/2 clinical program in Colorectal Cancer, Gastric Cancer and Neuroendocrine Tumours
- First-in-class CAR T cell therapy licenses from world renowned cell therapy center, the University of Pennsylvania after a decade of optimization
- Preclinical evidence published in Nature Cancer demonstrated complete eradication of tumor cells with no relapse in seven different types of cancer
- Clinical trial to be initiated in 2024 at leading US institutions





### CHM 0201 for BLOOD CANCERS AND SOLID TUMOURS

### **ADVANCING NOVEL PHASE 1B CLINICAL COMBINATIONS**

CHM 0201 Phase 1A

COMPLETE TO ENROLLMENT

Established Safety

Complete Response in Acute
Myeloid Leukemia

Demonstrated Persistence & Expansions

CHM 0201 + VACTO
Phase 1B

ONGOING ENROLLMENT

Initiated January 2023

Acute Myeloid Leukemia and Colorectal Cancer

First Study to Investigate NK Cells + Vactosertib

CHM 0201 + AZA/VEN
Phase 1B

**ENROLLMENT LATE 2023** 

FDA IND Clearance

Acute Myeloid Leukemia

First NK Combination Study in Front Line AML

NEW CLINICAL TRIAL







Making Cancer History®



# **CHIMERICTHERAPEUTICS**

# VALUE PROPOSITION FOR INVESTORS

1.

# ADVANCED, UNDERVALUED PORTFOLIO

Chimeric's pipeline and portfolio are highly undervalued in relation to other cell therapy companies

2.

# ADVANCING DEVELOPMENT TO VALUE REALIZATION

Chimeric's assets are entering Phase 1B, where big pharma M&A is now focused

3.

# NAVIGATING THE CHALLENGES

Chimeric is responding to todays biotech environment with program prioritization, cash preservation and business development

# 1. ADVANCED, UNDERVALUED PORTFOLIO

Chimeric has an industry leading cell therapy portfolio with novel technology platforms in clinical stage development





Novel technology platforms



Assets with positive clinical data



Ongoing clinical trials

## 2. VALUE REALIZATION OPPORTUNITY FOCUSES ON PHASE 1B/2

#### CHIMERIC IS ADVANCING DEVELOPMENT TO BIG PHARMA'S FOCUS

In today's market, big pharma M&A has shifted its focus from preclinical assets to assets that are clinically de-risked with Phase 1 A/B data

Chimeric currently has 3 ongoing Phase 1B clinical trials.

As development advances to enhance M&A opportunity, funding requirements become more significant.

> PRE **CLINICAL**

1-2 years

In vitro and in vivo experiments with lead asset

**PHASE** 2/3 **REGISTRATION TRIAL PHASE** 

> **BIG PHARMA** PHASE OF **INTEREST**

3-4 years

1A/ B CLINICAL

**TRIALS** 

Establishing safety, efficacy and the recommended dose

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- CHM 1101
- CHM 0201 + VACTO
- **CHM 0201 + AZA/ VEN**

3-5 years

to support

commercial

approval

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Registration trial

**REGULATORY APPROVAL** 

1 year

Approval by global regulatory authorities to commercialize

3-5 years

**ASSET** 

**DISCOVERY** 

Discovery and lead optimisation of assets

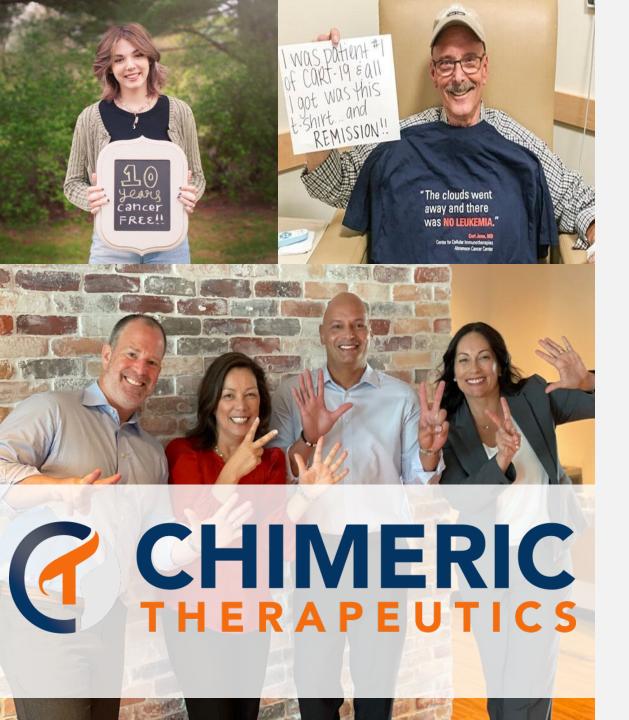
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### 3. NAVIGATING THE CHALLENGES

To enable clinical advancement, Chimeric has responded to the challenges of todays market with program prioritization and cash preservation while creating value through business development







- POSITIVE CLINICAL DATA in GBM and AML
  - Two assets with positive Phase 1A clinical data
- ADVANCING CLINICAL DEVELOPMENT
- Four ongoing clinical trials in solid tumours and blood cancers
  - **NEAR TERM MILESTONES**
- Multiple clinical catalysts in next 12-18 months
- IMPACTFUL BUSINESS DEVELOPMENT
- Two collaborative clinical trials with limited CHM funding

- INDUSTRY LEADING TEAM
- Experienced team with significant cell therapy development and commercialization expertise

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