# Manufacturing Brighter Futures

May 29, 2024



Enable widespread patient access to life-saving cell and gene therapies

The status quo in cell and gene therapy for most patients today represents death or serious disability



**Tim Hunt** Chief Executive Officer, Alliance for Regenerative Medicine (ARM) Globally, less than 4% of patients who could benefit from approved CAR-T therapies have been able to access them

# 900K\*

Addressable Patient Population

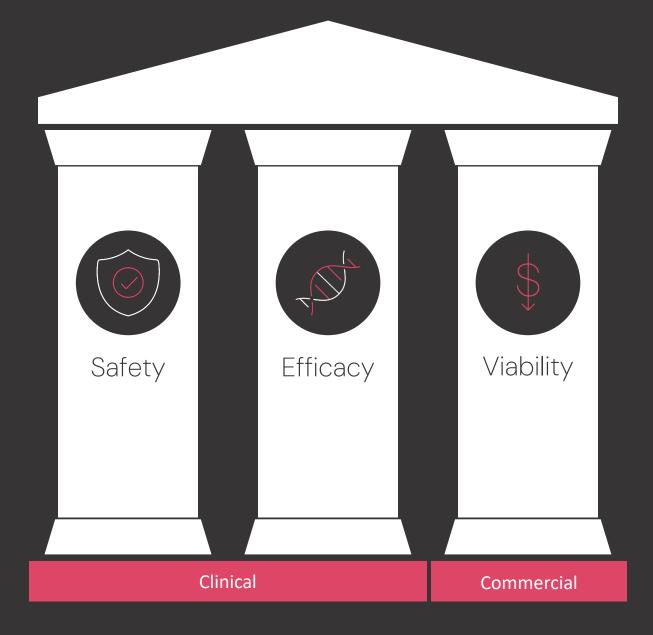


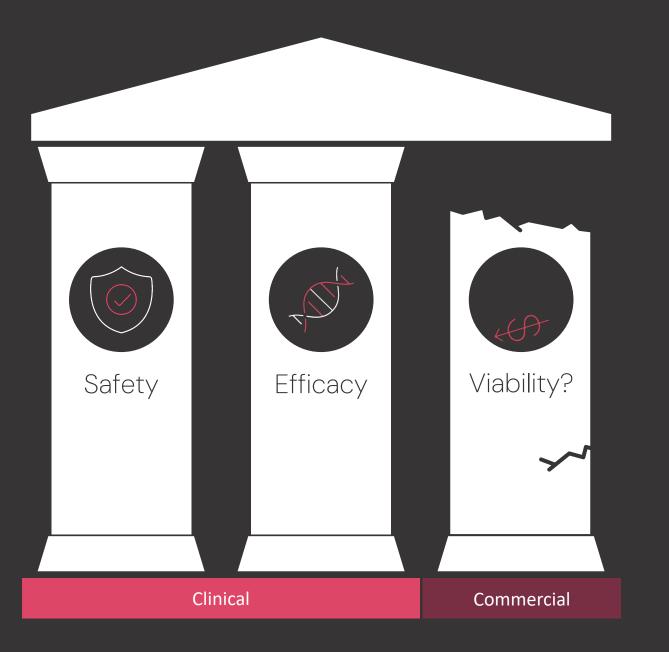
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\*Projected : Includes ALL, DLBCL, Multiple Myeloma, CLL and FL. Source: McKinsey, Statnews. Source: Ori Biotech Internal Research at www.oribiotech.com

# Small molecules and biologics

If we develop safe and effective products, we assume they will be commercially viable





# Cell and gene therapy

Developing safe and effective products is not enough to ensure that cell and gene therapies will be viable

Cell and gene (developers) are essentially manufacturing companies... You have to prioritize that first in addition to clinical and research



#### Andrew Obenshain

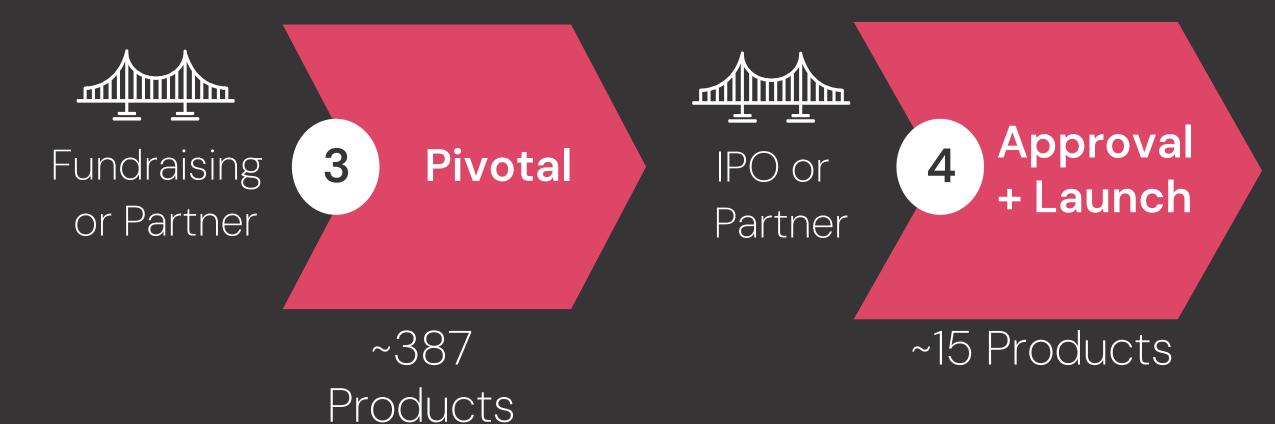
Chief Executive Officer, Bluebird Bio ARM State of the Union 2024



~1998 Products

~1812 Products





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Data shown above is from Beacon (Hanson Wade Database), ARM, and FDA



Success

# Who is

# Oribiotech

## Partnered

with leading Large Pharma, Biotech, Academic Medical Research, and CDMOs

# Labs in

Philadelphia, London and Cambridge (UK) – More coming soon!

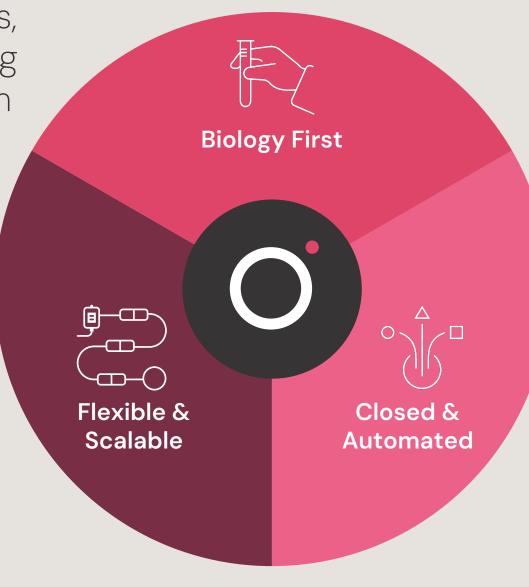
# Founded in 2015

Mission to enable widespread patient access to CGT

Team with 100+ Years operational experience

## 83 Person Team

CGT CDMO, Therapy Developer, Large Pharma, Technology, and Engineering Experience Over the past 5 years, we have been working to develop a platform that is...



to support you and the other heroes of cell therapy.

# Oribiotech



# Ori Platform Technology: Bioreactor, LVC, SVC, and OriConnect™





## Automate Better Biology<sup>™</sup>



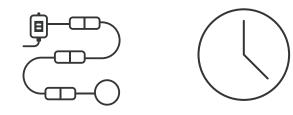
- Supports different cell suspension culture types and workflows
- Begin with ~50M cells (at 50ml) and yield ~12B cells (at 1L)
- Achieves higher cell yields and transduction efficiencies than the "gold standard"
- (Delivers) 170x fold expansion with no signs of exhaustion



- Allows for activation, transduction, expansion, and harvest in one platform
- Reduces manufacturing process times by 25% (average of 1-2 days)
- Delivers cells with >95% viability and potent cancer killing activity
- Delivers target cell yield with a 50% reduction in MOI



## **Accelerate Product Development**

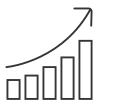


- Enables a seamless transition from R&D into the clinic and from the clinic to commercial scale manufacturing
- Brings products to market faster by reducing process development and tech transfer time
- Empowers your scientists, offering them the freedom to optimize and the flexibility to innovate
- Reduces batch failure rates to hit target dose more consistently

- Eliminates the need for comparability during scale up, shaving years off drug development
- Helps you reach your next value inflection point more quickly
- Puts control back in the hands of your expert scientists via customizable software
- Enables complete product traceability via our digital, cloud native platform



# Scale Your Impact<sup>™</sup>



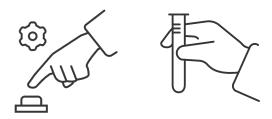
- Increases throughput to ~1,000 doses per year in 1,000 sq ft
- Reduces labor costs by more than
  50%
- Enables manufacturing in less expensive, lower grade clean rooms
- Designed to support full robotic automation and advanced analytics (AI, ML)



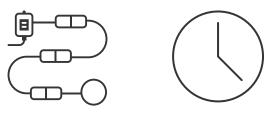
- Unlocks multi-site manufacturing capability
- Reduces manufacturing costs by 30-50%
- Delivers a > 50% reduction in manufacturing facility footprint
- Integrates with industry leading software (e.g. EBMR, ELN, SCADA etc.) via APIs



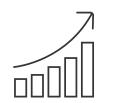
# **Automate Better Biology**



# **Accelerate Product Development**



# **Scale Your Impact**





# I bet you probably wouldn't believe me...

# **Completed Platform Validation**



Characterization runs completed (in house and at partner sites in NA and UK) Testing platform's ability to address donor and patient variability, showcasing the robustness of system outputs across different starting material

> 40

Donors

/ Patients

Different processes including CAR-T, TCR-T, TILs, CD34+, with CAR-M and others on the horizon

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Unique

Processes



5 therapy developers, 4 CDMOs, and 1 AMC

# I expected automation, I expected more process insights, but I never expected better biological performance right out of the gate



#### Jason Bock

Co-Founder and Chief Executive Officer, CTMC – A Joint Venture Between Resilience + MD Anderson Cancer Center





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Let's Manufacture Brighter Futures Together.