



Make Your Own Technology

An exciting, innovative delivery platform
for Antibodies and Therapeutic Proteins

February 26, 2024

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Executive Summary

Potential paradigm shift in delivery of antibodies and therapeutic proteins for treatment of chronic diseases and prevention of infectious diseases.

MYO
TECHNOLOGY



Intramuscular electroporation of plasmid DNA enables durable, in vivo production of antibodies and therapeutic proteins

Company and financing overview

\$4M Series Seed

\$24M Series A

\$13M Nondilutive grants

Located in New York City

13 Employees

- ✓ Patent-protected platform technology
- ✓ Substantial commercial potential identified across a variety of indications
- ✓ Established clear proof of concept
- ✓ Able to support a wide variety of payloads
- ✓ Solves supply and distribution challenges associated with biologics
- ✓ Early clinical development funded by government agencies in Zika prevention



Rachel A. Liberatore, PhD
President & Chief Scientific Officer

BA in Molecular Biology from Princeton University |
PhD in Cellular & Molecular Biology from Columbia University | 10+ years in scientific leadership and team management



Yaoxing Huang, PhD
Co-founder and Scientific Advisor
Associate Professor, Columbia University

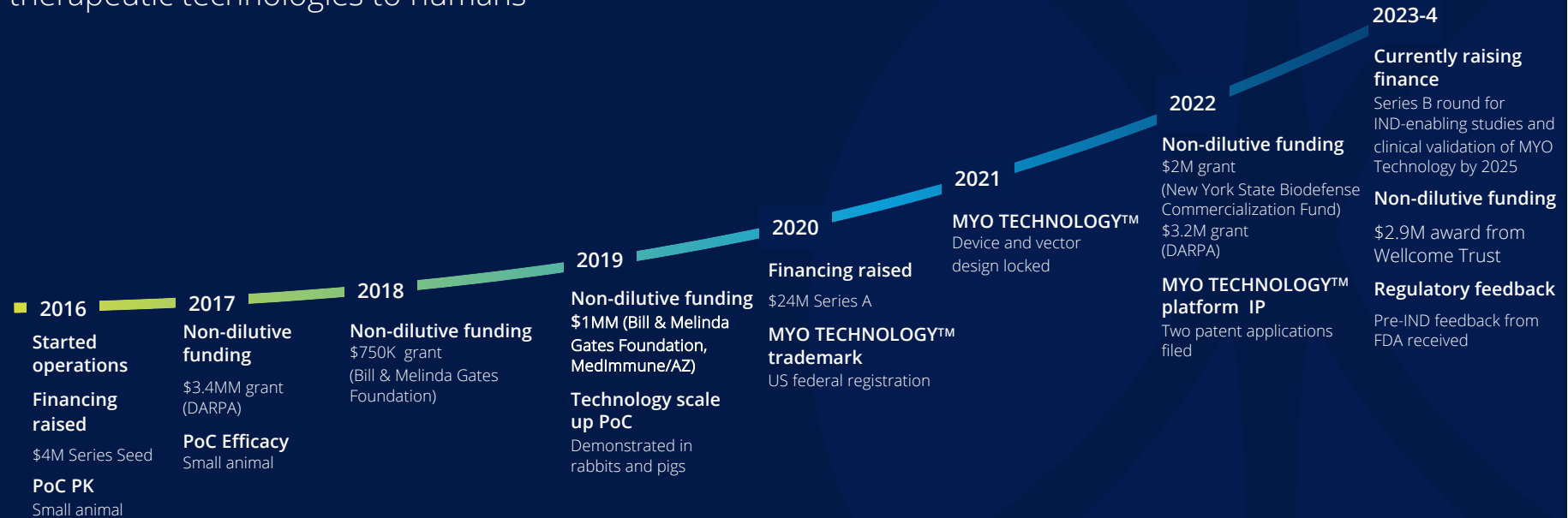


David D. Ho, MD
Co-founder and Scientific Advisor

Professor, Columbia University & Director,
Aaron Diamond AIDS Research Center |
Scientific Founder, TaiMed Biologics (US FDA approved Trogarzo®) | Time Man of the Year

Our story

Considered, steady progress built on a solid foundation to advance the delivery of first-in-class and best-in-class DNA therapeutic technologies to humans



There remain high levels of unmet need for the optimal development, supply, and use of antibodies and therapeutic proteins today

A faster, smoother, and cost-effective delivery platform would address many of these needs

Challenges that remain

Therapies with **short half-lives** require **frequent dosing** and ongoing monitoring, leading to inefficient clinic workflows and a risk of suboptimal efficacy and tolerability



Specialized production, purification and cold-chain requirements for transport and storage, drive long production lead times, complicate supply and distribution, and result in an unnecessarily **high COGs**



Meeting the need

Improving clinical utility for healthcare professionals by significantly reducing clinical workload with **easier, less frequent dosing**, negating the need for regular infusion clinics, reducing the burden of patient monitoring, and guaranteeing **real world efficacy** that meets that seen in clinical trials as patient adherence is no longer part of the equation

Reducing the COGs and ensuring flexible, fast product supply, distribution and storage by **simplifying manufacturing processes**, negating the need for specialized production facilities and removing any onerous **distribution and storage** criteria

Our MYO Technology™ delivery platform enables an individual to make their own antibodies and therapeutic proteins, improving on frequent dosing regimens required by conventional delivery of recombinant proteins



Bio Blueprints

Proprietary DNA plasmid encoding antibody or protein therapeutic



Delivery Device

Proprietary electroporation device



Antibody Factory

An individual's muscle cells, produce antibodies or therapeutic proteins following in vivo electroporation



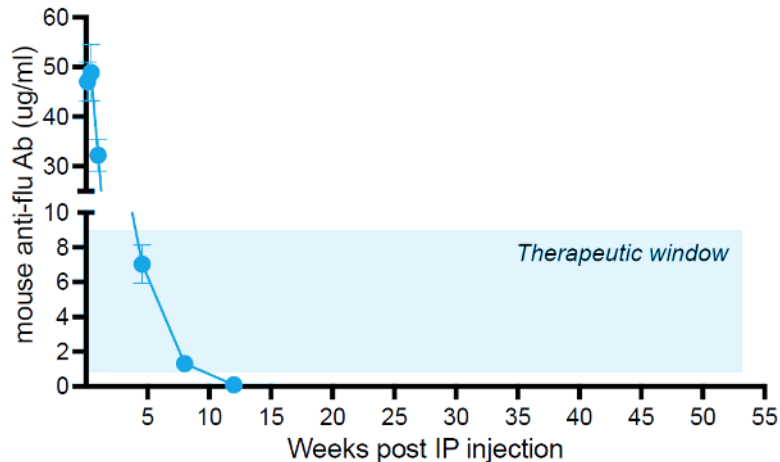
Antibodies/Therapeutic Proteins

Circulate systemically following secretion by muscle cells

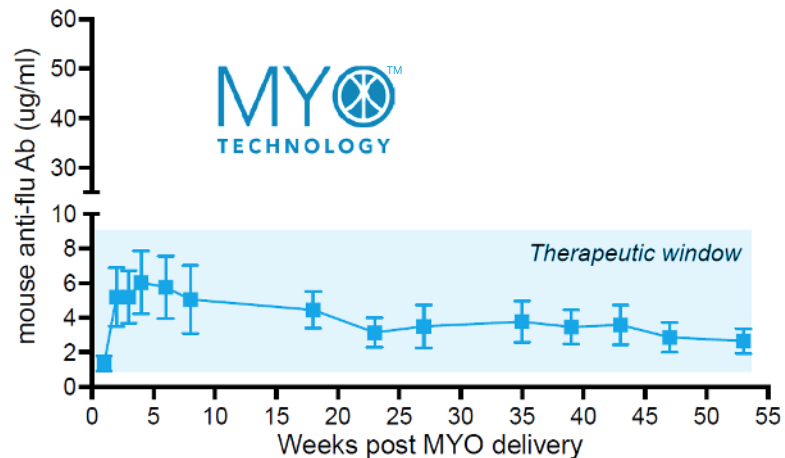
We continue to build the evidence to demonstrate the high value of our technology platform

In vivo animal data demonstrates promise for durable delivery of a payload within the therapeutic window

Traditional antibody protein delivery



MYO delivery of antibody genes



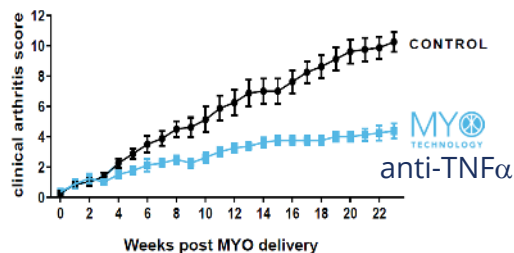
Our MYO Technology™ has compelling advantages compared to other novel delivery platforms when applied to the management of systemic, chronic disease and prevention of infectious disease

<u>Feature</u>	MYO™ Technology	mRNA	AAV Gene Therapy
Durability	Months to years	Weeks to months	Years/Permanent
Redosable	****	***	*
Large genetic payload	****	***	**
Large scale manufacturing	****	**	*
Freedom from cold chain	****	*	*
Clinical safety of technology	***	***	*
Opportunity	Systemic activity Chronic disease Tx, Infectious disease Px	Localized activity Short half-life Vaccines	Specific tissue target Genetic disease Tx

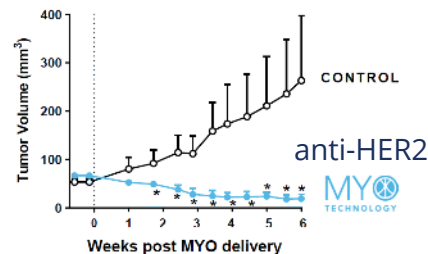
MYO Technology™ has the potential for broad applicability

Compelling in vivo animal efficacy studies, using well accepted models, show applicability for diverse range of diseases from oncology to autoimmune

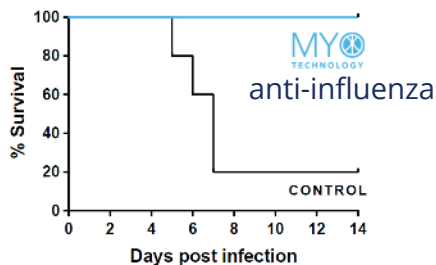
Arthritis



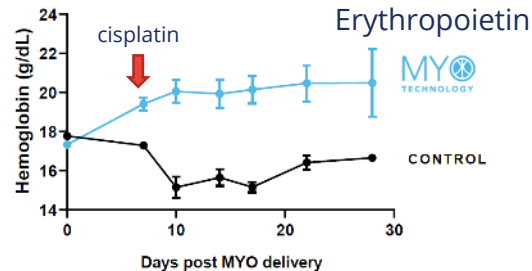
Breast Cancer



Influenza



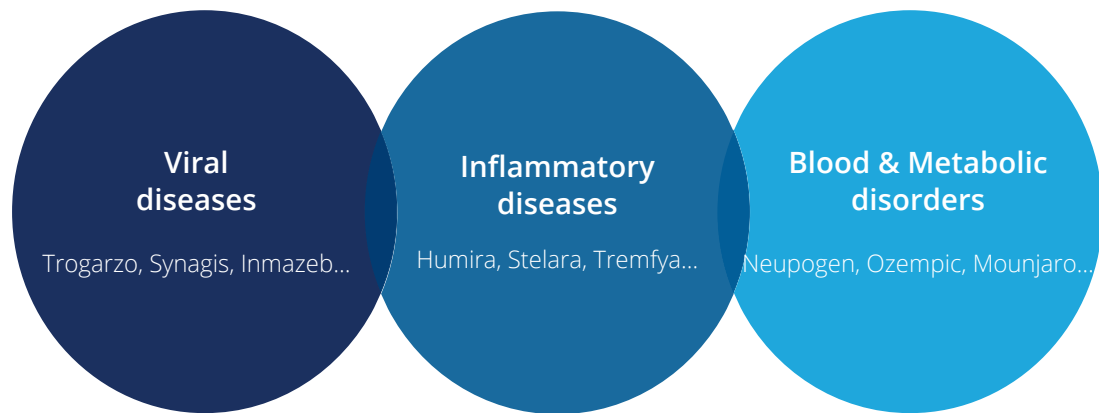
Anemia



Targeting high value, high growth markets

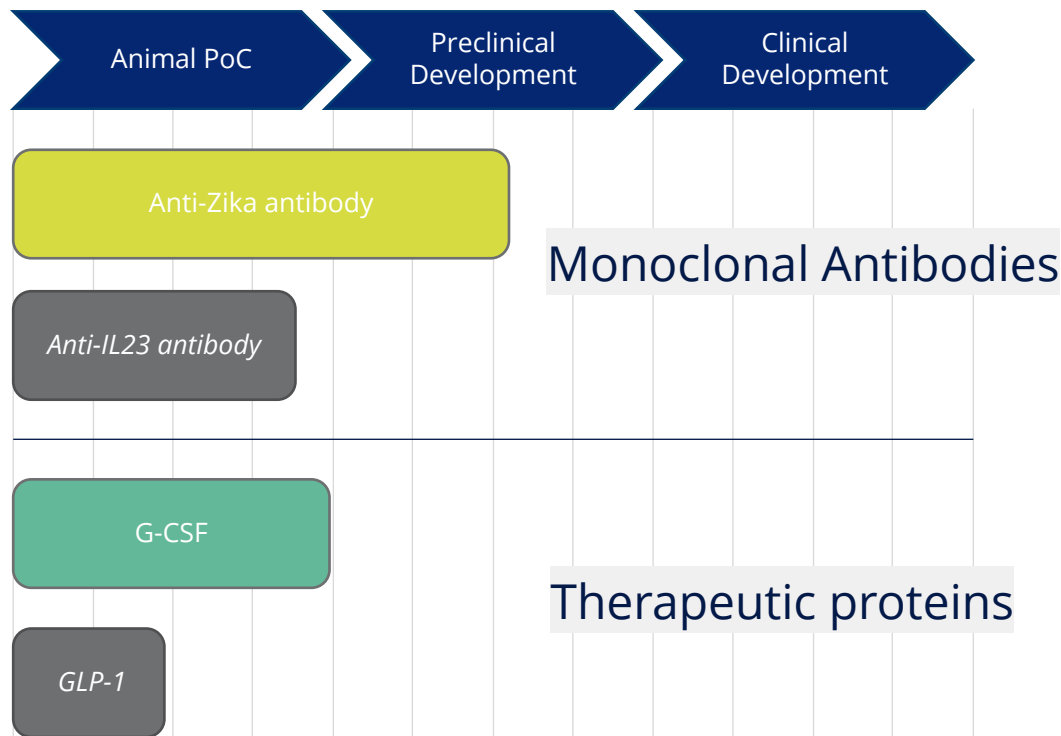
The market for antibodies and therapeutic proteins is very large with wide utility.

Growing rapidly, the global market for antibodies was valued at \$210B in 2022, with an expected CAGR of 11.04% from 2023 to 2030.



Clinical proof-of-concept with a monoclonal antibody (Zika) and a therapeutic protein (G-CSF) will support expansion into multiple, high value markets

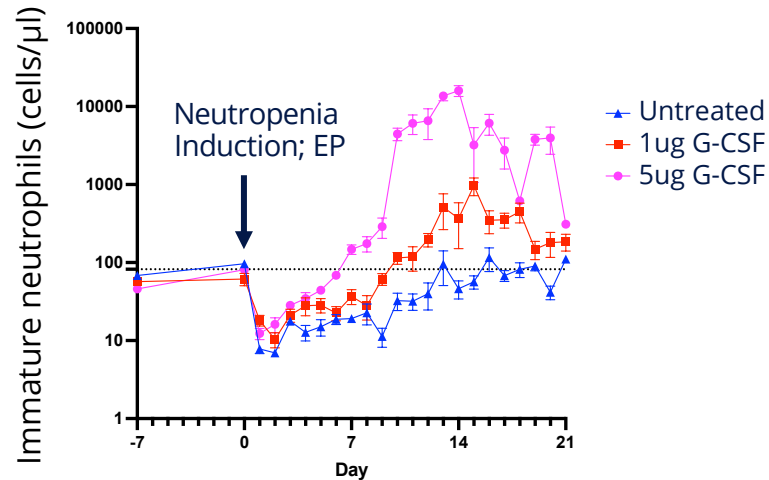
Phase 1 data with prototype molecules (anti-Zika monoclonal antibody and G-CSF) will demonstrate the potential for the MYO Technology platform



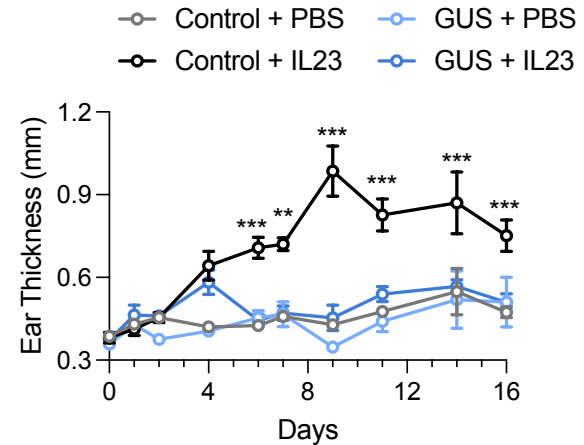
Proof of concept data for G-CSF and anti-IL-23 delivery with MYO Technology

Animal models of neutropenia and plaque psoriasis demonstrate the potential for MYO Technology in these indications

MYO G-CSF: DNA dose-dependent treatment of anti-Ly6G-induced neutropenia

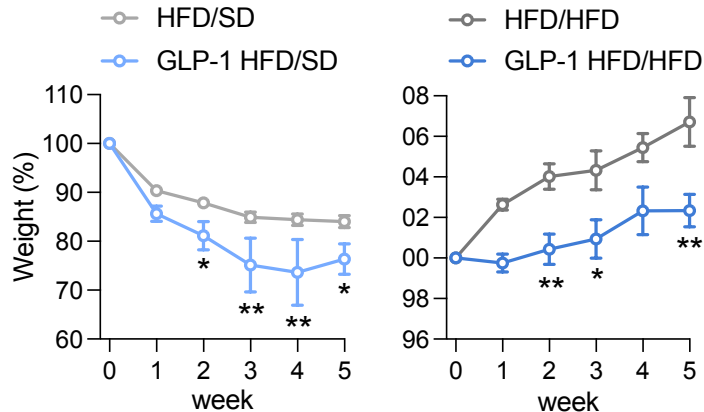
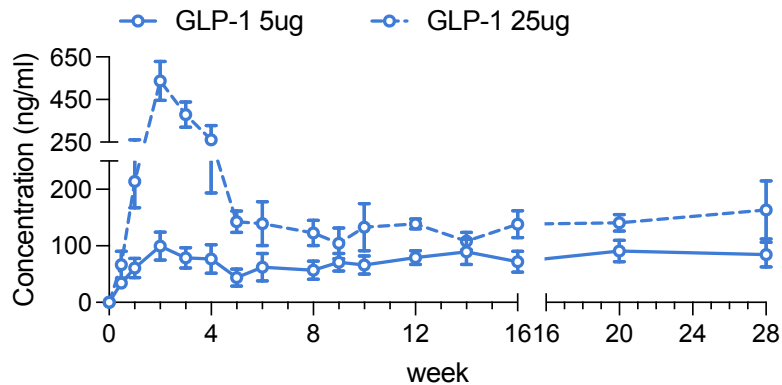
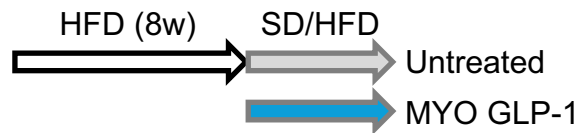


MYO anti-IL-23: DNA-based delivery of guselkumab potentially inhibits IL-23-induced psoriasis



Proof of concept data for GLP-1 delivery with MYO Technology

DNA-based delivery of a GLP-1 receptor agonist results in sustained expression and promotes weight loss and suppresses weight gain in an animal model of diet-induced obesity



Three pillars of opportunity identified for our MYO Technology™ platform

Multiple opportunities exist for biopharma partners to realize the full potential of drugs in large and rapidly growing markets

New molecule requiring long-term delivery

Antiviral antibodies and other molecules for which durability is critical

Life cycle Management

Established brands wanting to extend dominance

Differentiation in a competitive market

New/generic brands wanting to make an impact in crowded markets

A track record of accomplishment, planning for future success

Significant milestones achieved following each funding round



Seed (\$4M) to Series A



Series A (\$24M) to present



Planned for Series B (\$27M)

- PK and Efficacy demonstrated in small animal models
- Technology scale up demonstrated in large animal models
- Additional non-dilutive funding secured (\$3.4M DARPA, \$1.75M Bill & Melinda Gates Foundation)

- MYO Technology device and plasmid design locked
- 2 patent applications filed (device, plasmid)
- Manufacturing of GLP and clinical devices initiated
- First GMP plasmid manufacturing campaign initiated
- FDA feedback on Pre-IND questions and briefing package received
- NHP studies initiated
- Further non-dilutive funding secured (\$2M New York State Biodefense Commercialization Fund, \$3.2M DARPA, \$2.9M Wellcome Trust)

- Completion of IND-enabling studies and Phase 1 clinical study for lead clinical development program (*fully funded with non-dilutive funds*)
- IND-enabling activities and Phase 1 clinical study for first internally funded clinical development program
- Proof-of-concept animal studies for second and third planned internal clinical development programs
- Biopharma partnerships
- Board of Directors enhancement for next phase of growth

Clinical and Business Development Team Partners



- Martin Markowitz, MD – Clinical
 - Clinical Director and Principal Investigator on ~80 clinical trials
- Jim Ackland – Regulatory, preclinical
 - >45 years experience in development and regulatory affairs for biopharmaceuticals
- Meredith Brown-Tuttle, FRAPS – Regulatory, biologics
 - >31 years experience in regulatory affairs for biologics
- Sheila Ramerman, RAC-US, RAC-Devices – Regulatory, medical devices
 - >30 years experience in electromedical devices
- Joanne Kelley, LLB – Business Development
 - Former Vice President of Business Development, Head of Transactions at AstraZeneca
- Jill Ogden, PhD – Business Development
 - >30 years commercial/transactional expertise in biopharmaceutical industry

Thank you

Help us to transform the lives of people globally
via the delivery of therapeutic antibodies and proteins
enabled by our MYO Technology™, easing and
expanding the use of these therapeutics.

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