



Ultra long-lasting, DNA-Based Delivery of Incretin Receptor Agonists

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Forward-looking statements

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MYO Technology

Paradigm shift in delivery of antibodies and therapeutic proteins
for treatment of chronic diseases and prevention of infectious diseases

MYO[®]

TECHNOLOGY



Bio Blueprints
Proprietary DNA plasmid



Delivery Device
Proprietary electroporation device



Antibody Factory
Muscle cells produce proteins following in vivo transfection



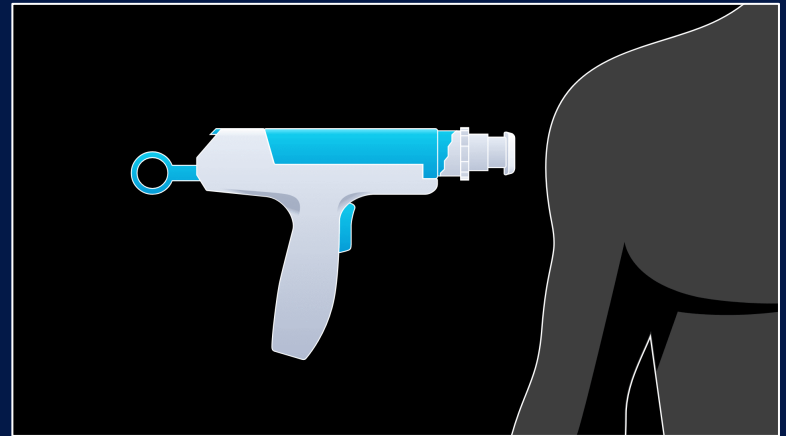
Antibodies/Therapeutic Proteins
Circulate systemically following secretion

MYO Technology Delivery Device

Generator



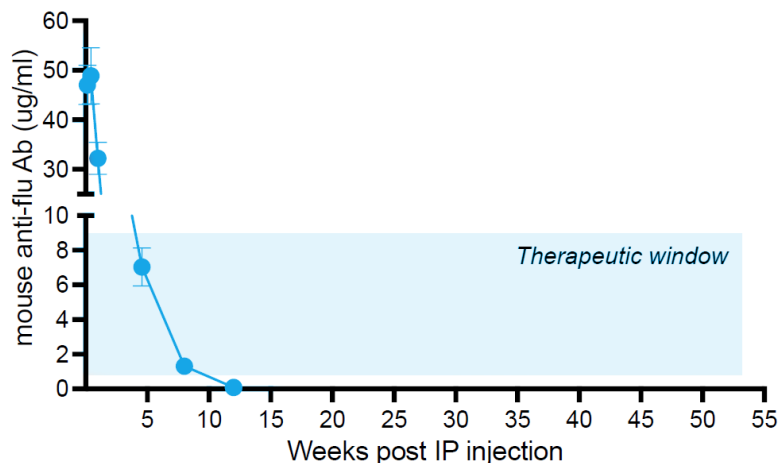
Handpiece



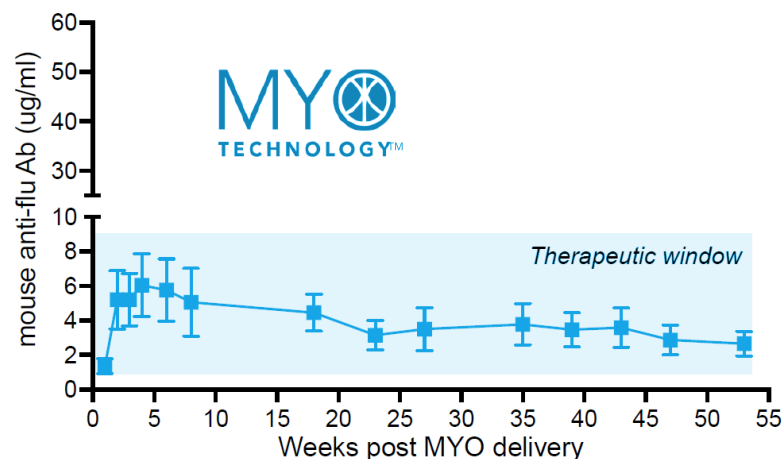
MYO Technology enables ultra durable protein expression



Traditional antibody protein delivery



MYO delivery of antibody genes

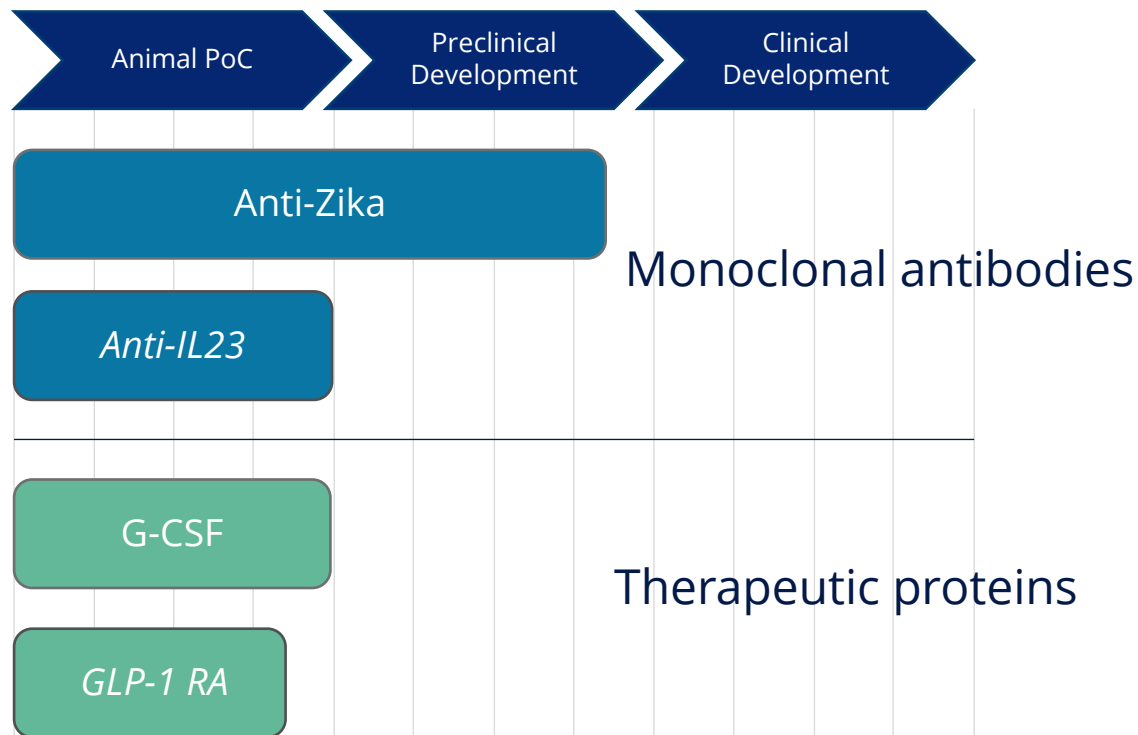


Unpublished, internal data

MYO Technology is a perfect fit for chronic disease treatment and infectious disease prevention

Feature	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

Pipeline: mAbs and therapeutic proteins

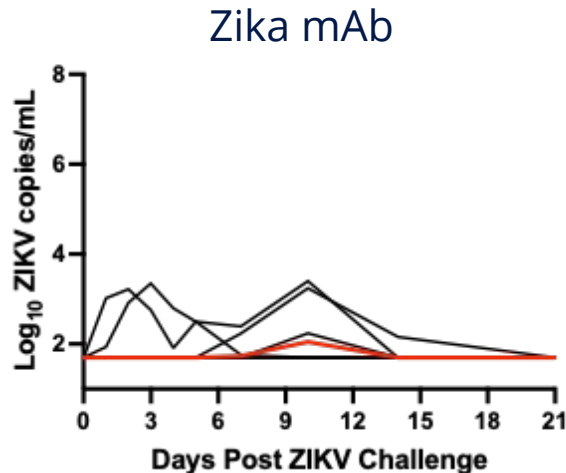
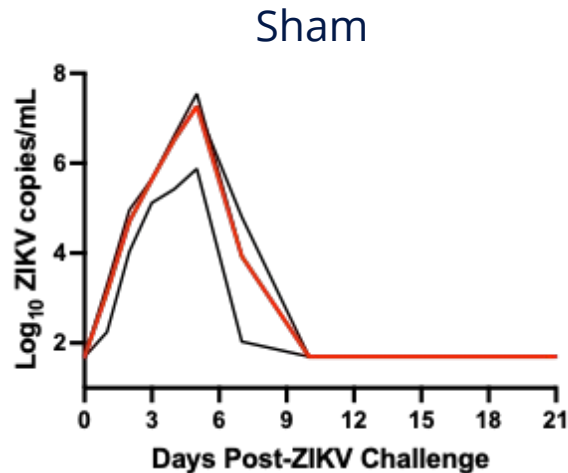


Delivery of an anti-Zika virus mAb with MYO Technology protects rhesus macaques in a Zika challenge model

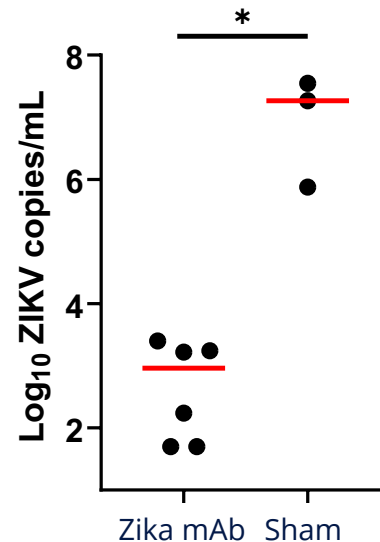
Animals treated with MYO Technology had ~10,000-fold reduction in peak viral loads



Plasma viral loads



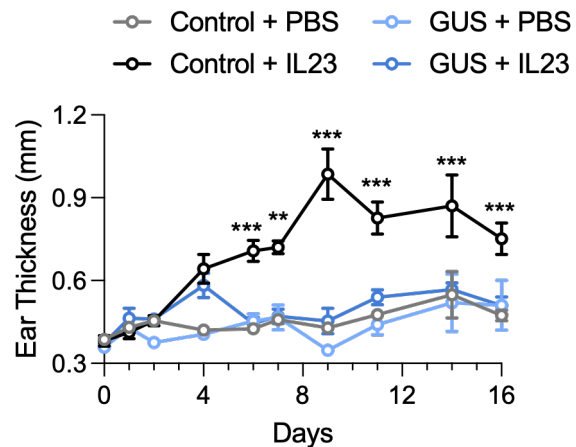
Peak plasma viral loads



Preclinical PoC for G-CSF and anti-IL-23 delivery with MYO Technology

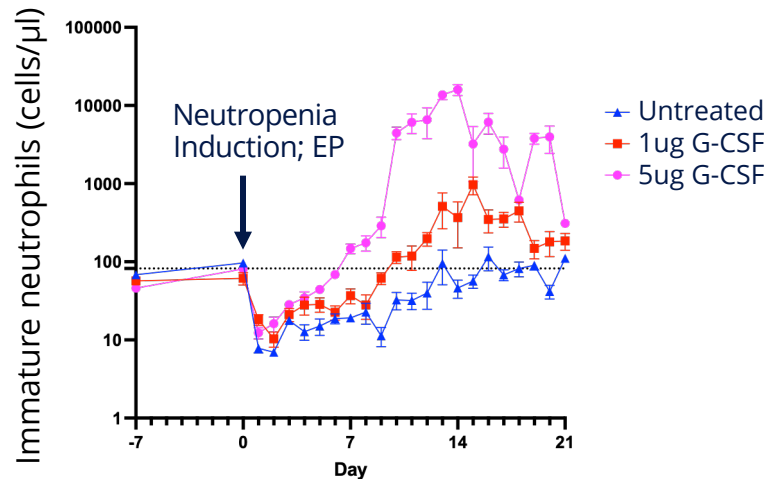


MYO anti-IL-23: DNA-based delivery of guselkumab potentially inhibits IL-23-induced psoriasis



Unpublished, internal data

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

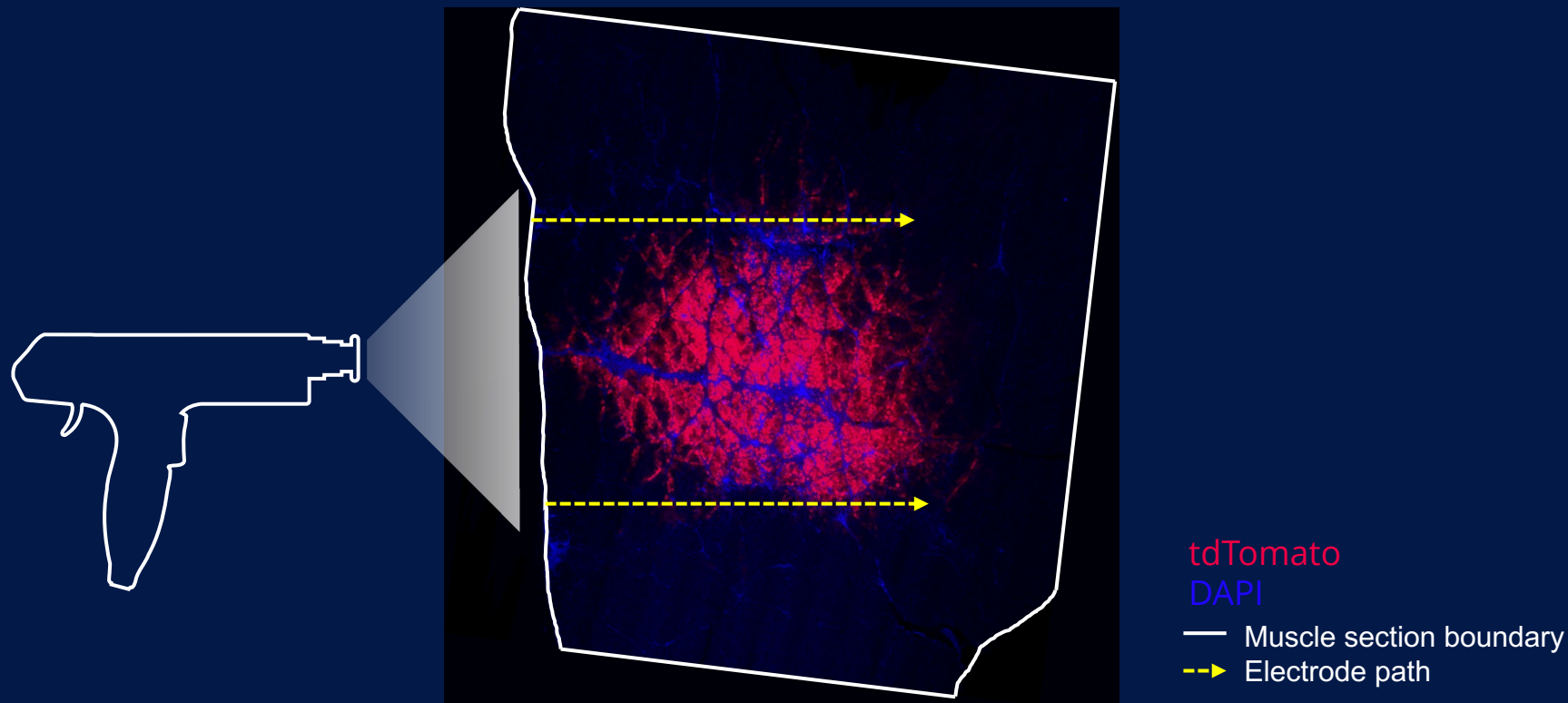


Drozd MM, et al. Molecular Therapy (2025), <https://doi.org/10.1016/j.ymthe.2025.03.038>

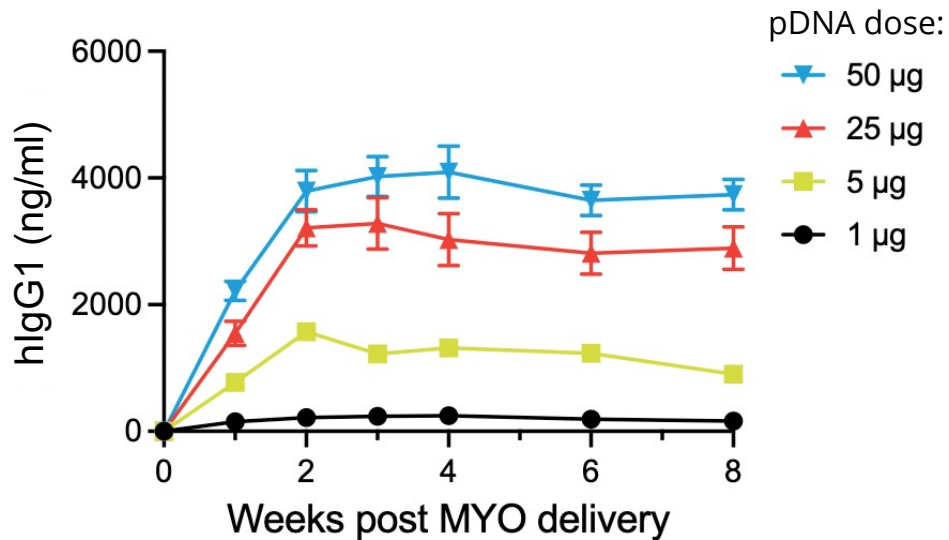
MYO Technology

features

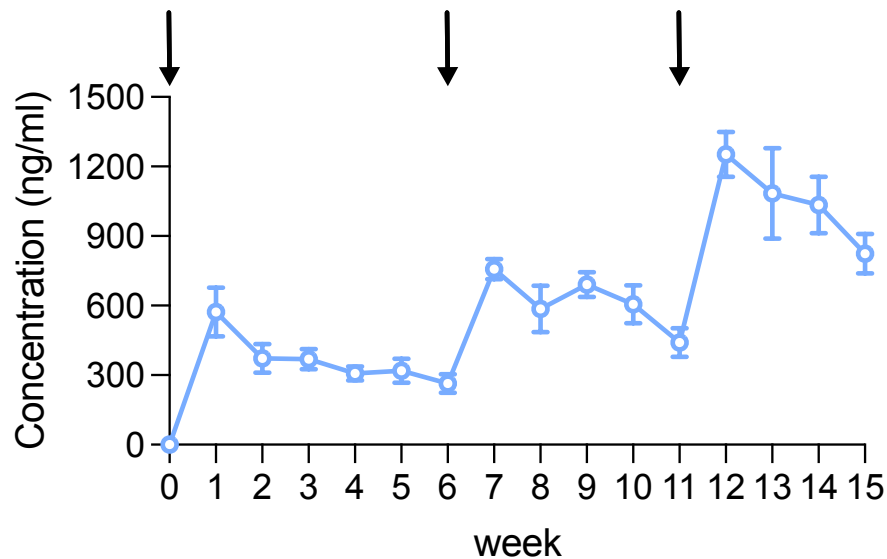
In vivo electroporation efficiently transfects muscle cells within the footprint of the device



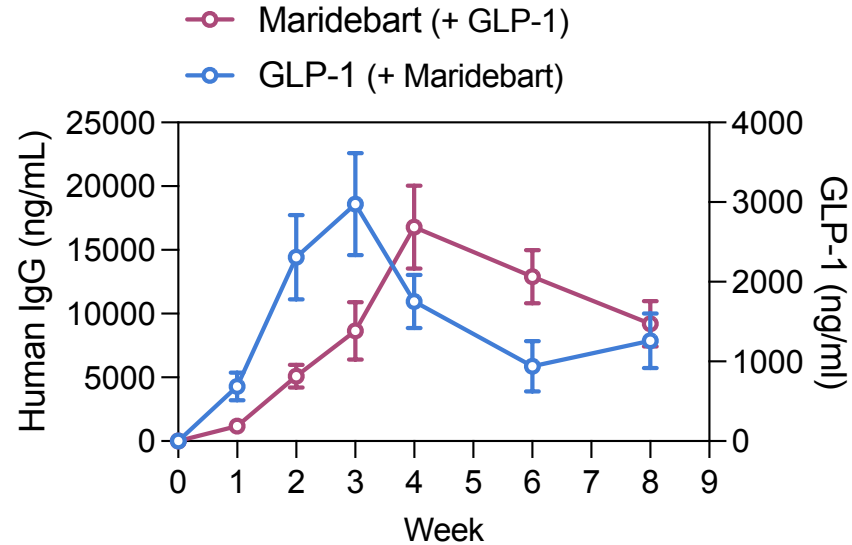
Protein expression correlates with the amount of pDNA



MYO Technology permits robust repeated dosing



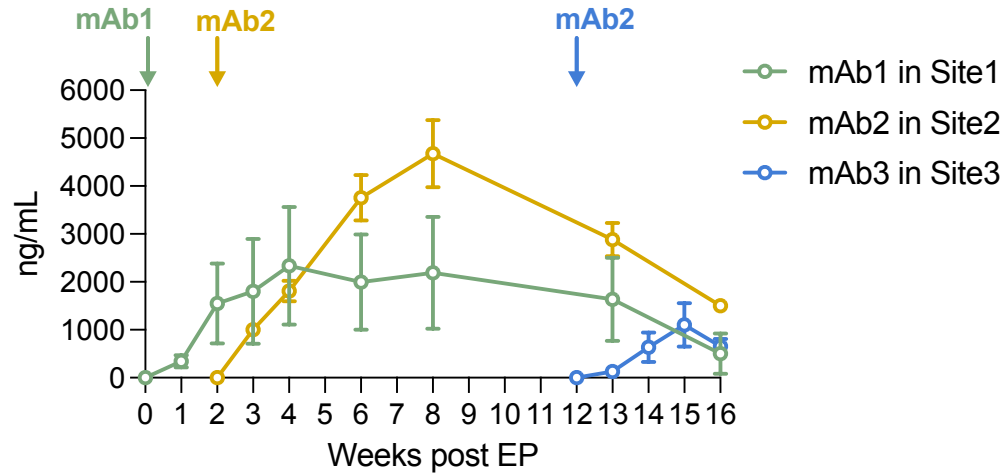
MYO Technology permits simultaneous delivery of multiple molecules at the same site



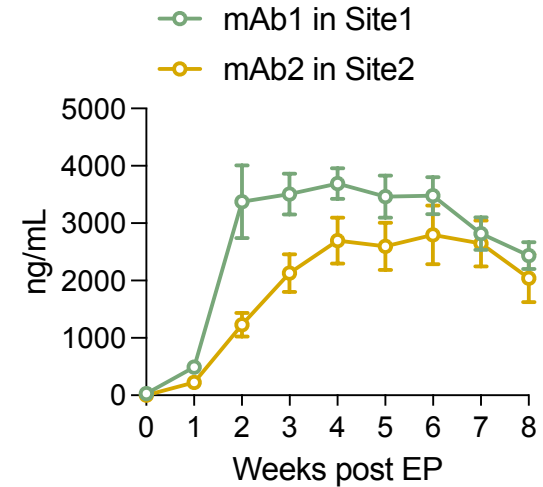
MYO Technology permits delivery of multiple molecules at different sites



Sequential delivery



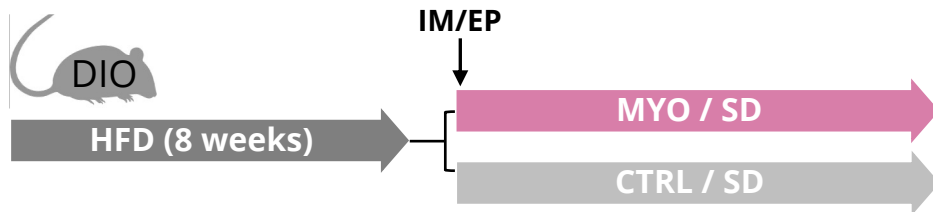
Co-delivery



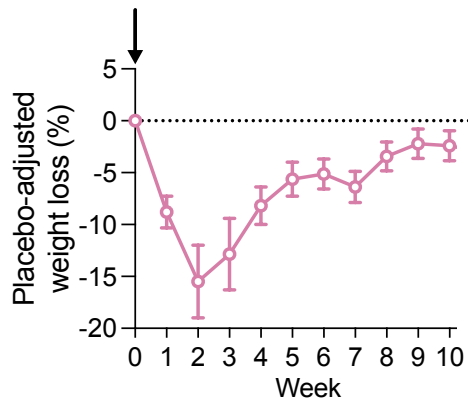
MYO Technology

Preclinical results on Incretin Receptor Agonists

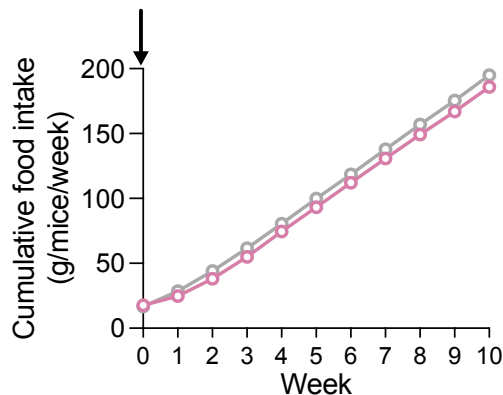
DNA-based delivery of GLP-1 peptide is efficacious in a DIO model



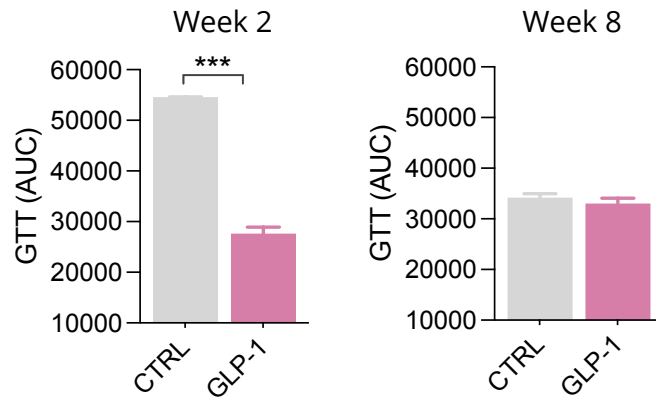
Weight Loss



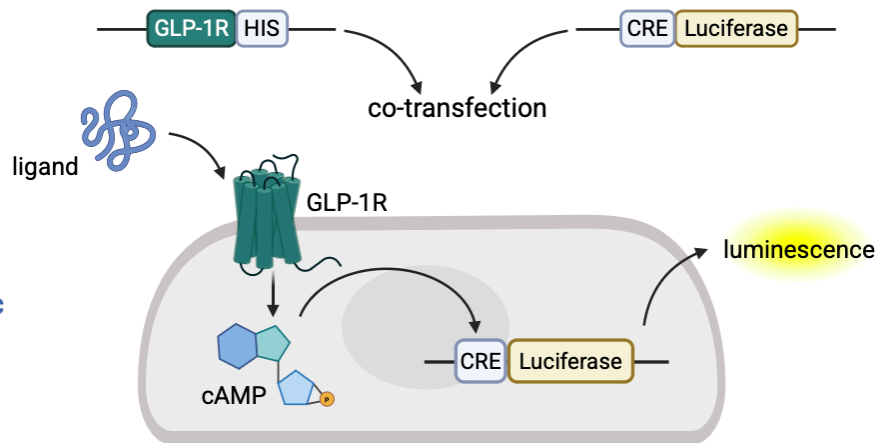
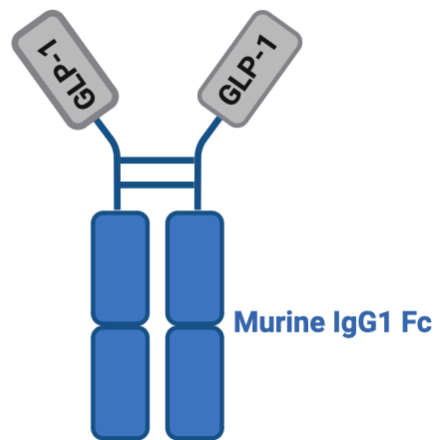
Food Intake



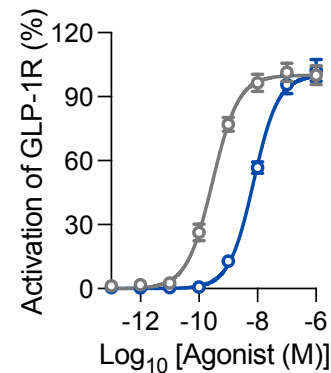
Glucose clearance



GLP-1 fused to an Fc domain exhibits therapeutic effects comparable to those of semaglutide

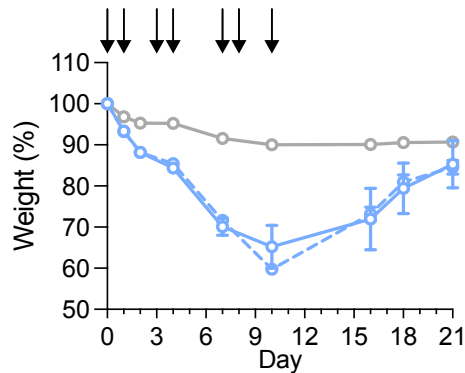


- Semaglutide - EC_{50} 0.29 nM
- GLP-1-mIgG1 - EC_{50} 7.35 nM

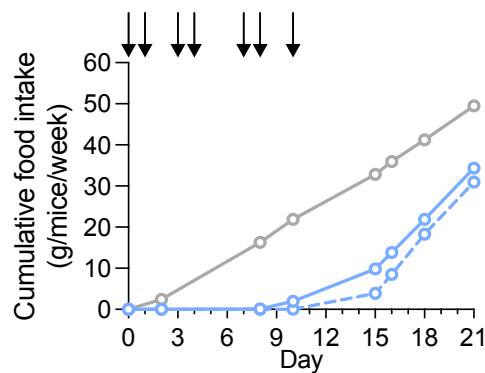


GLP-1 fused to an Fc domain exhibits therapeutic effects comparable to those of semaglutide

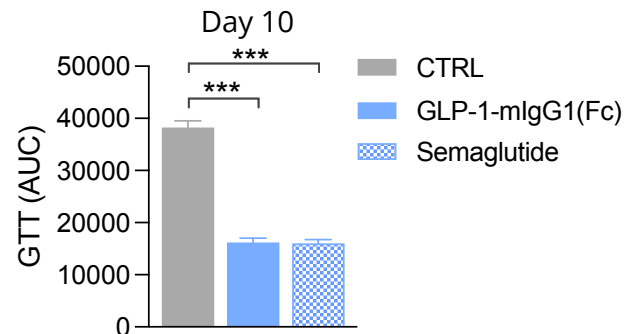
Weight Loss



Food Intake

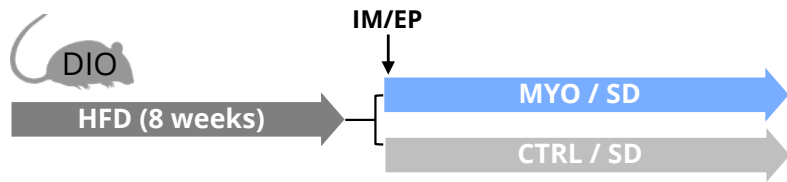


Glucose clearance

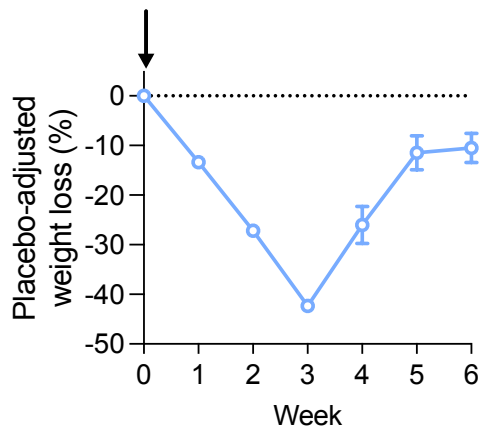


—○— CTRL —○— GLP-1-mIgG1(Fc) - - -○- - Semaglutide

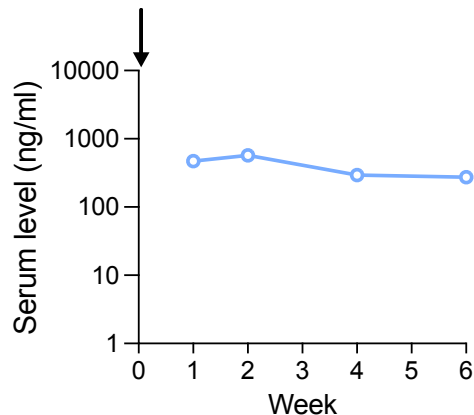
DNA-based delivery of GLP-1 Fc fusion improves therapeutic effects compared to native GLP-1



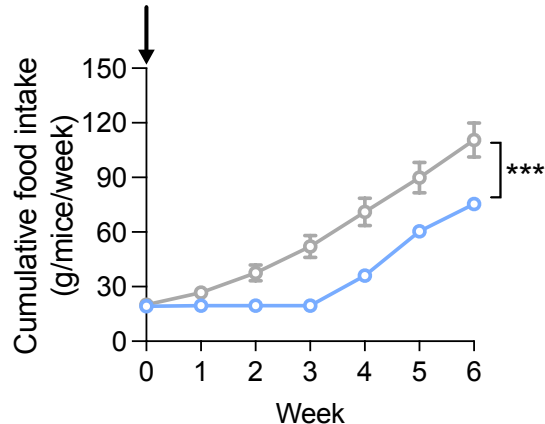
Weight Loss



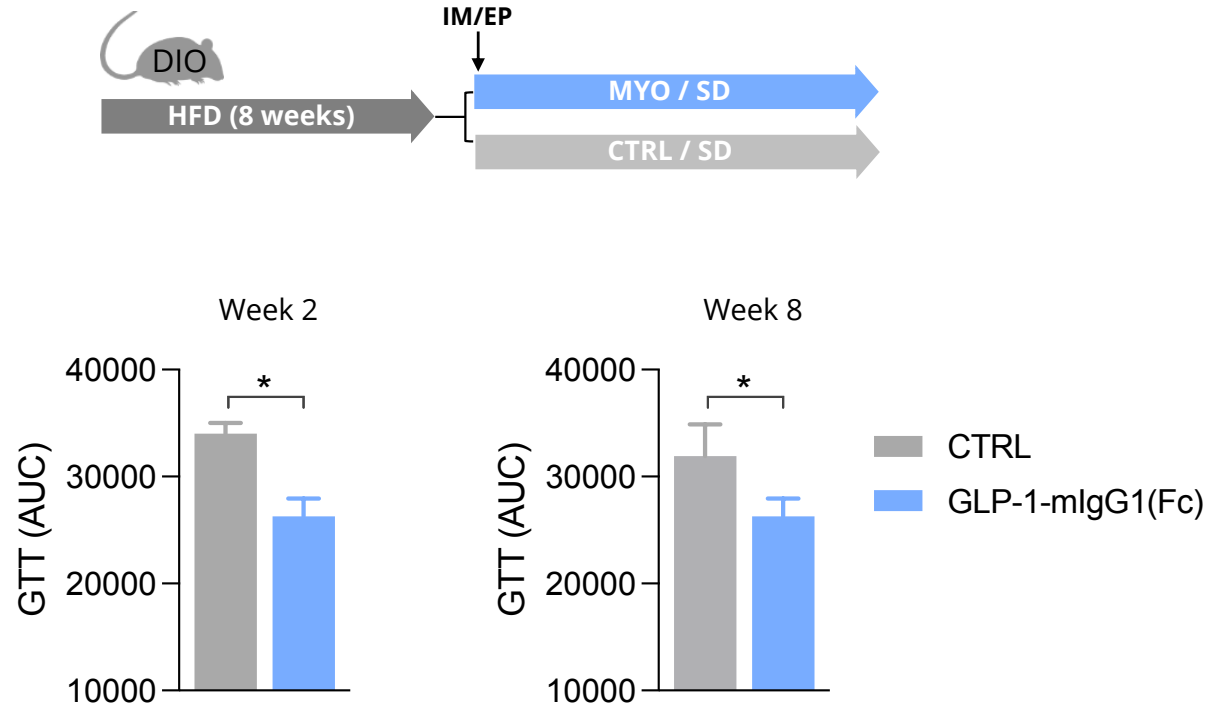
Serum level



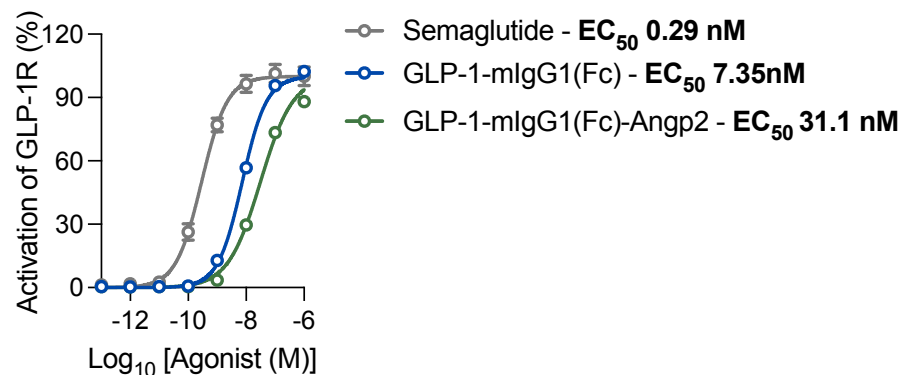
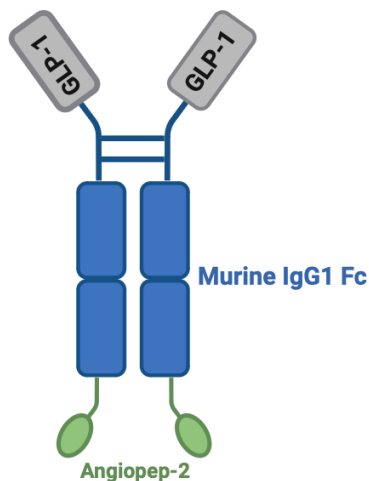
Food Intake



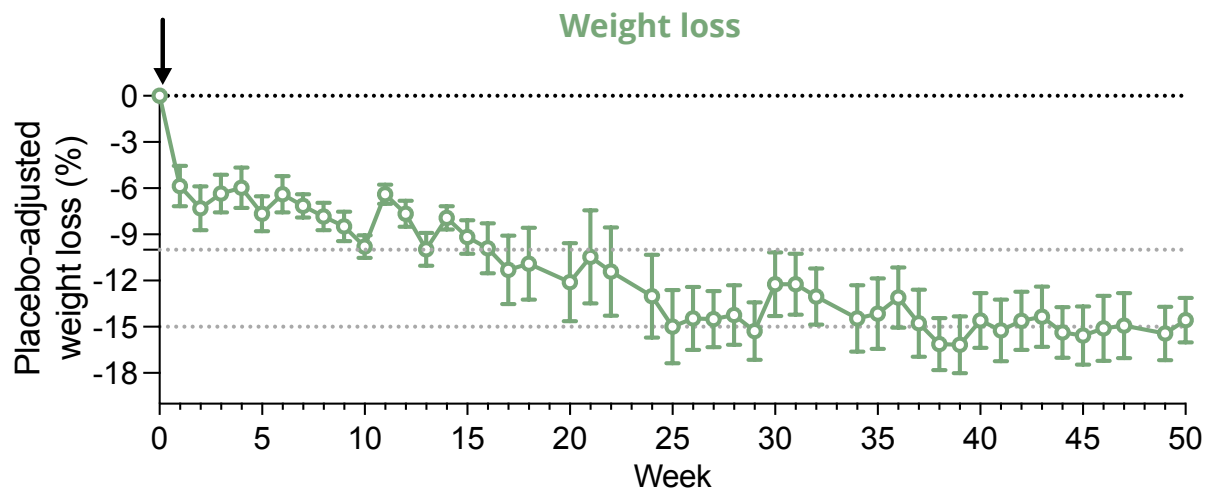
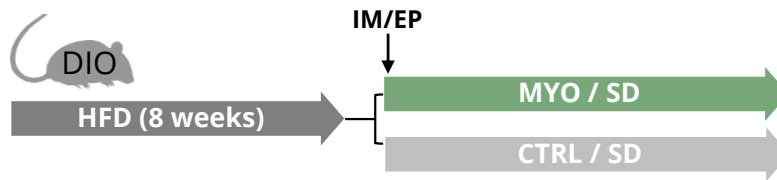
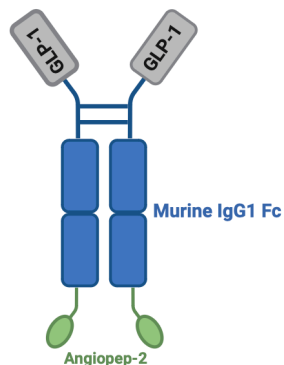
DNA-based delivery of GLP-1 Fc fusion improves therapeutic effects compared to native GLP-1



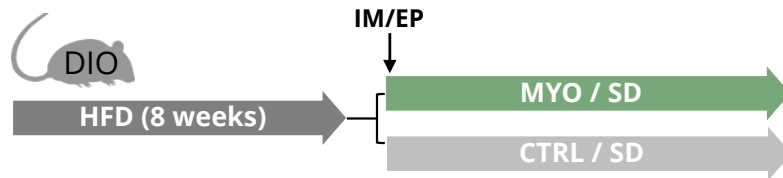
Will enhanced BBB penetration further improve efficacy?



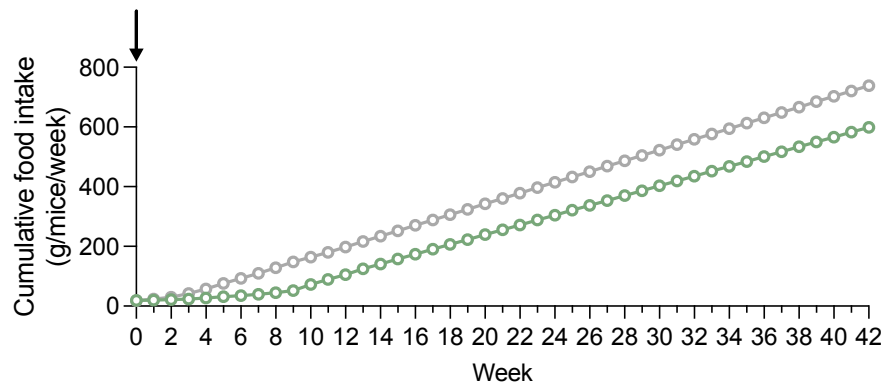
Enhanced BBB targeting dramatically improves GLP-1 Fc fusion efficacy



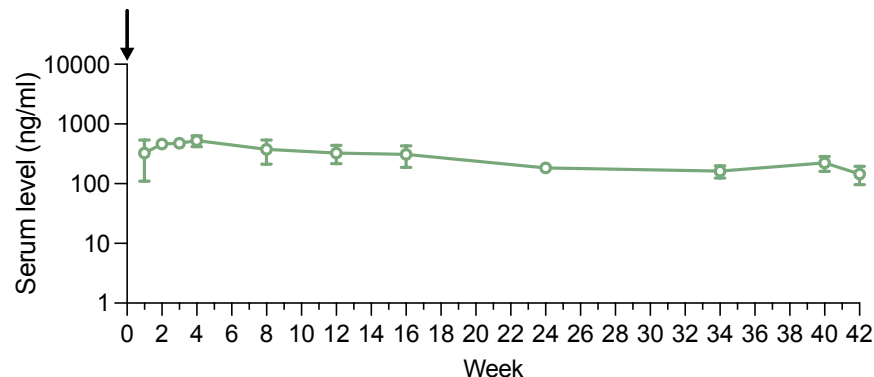
Enhanced BBB targeting dramatically improves GLP-1 Fc fusion efficacy



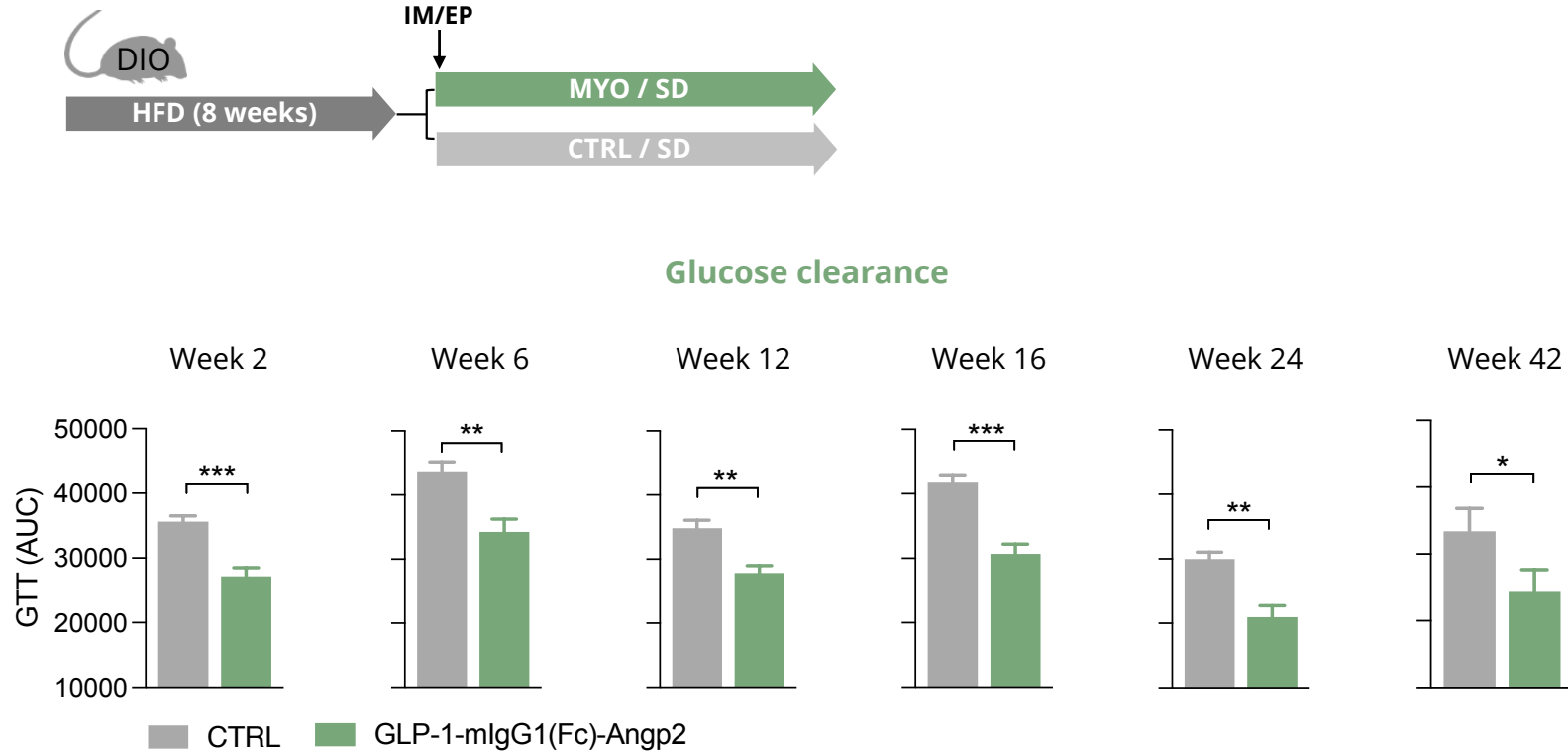
Food Intake



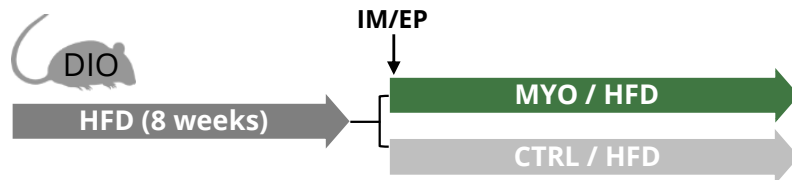
Serum level



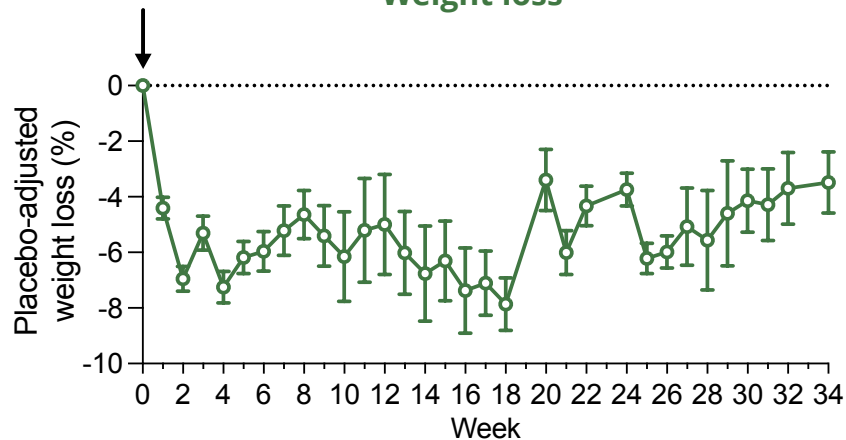
Enhanced BBB targeting dramatically improves GLP-1 Fc fusion efficacy



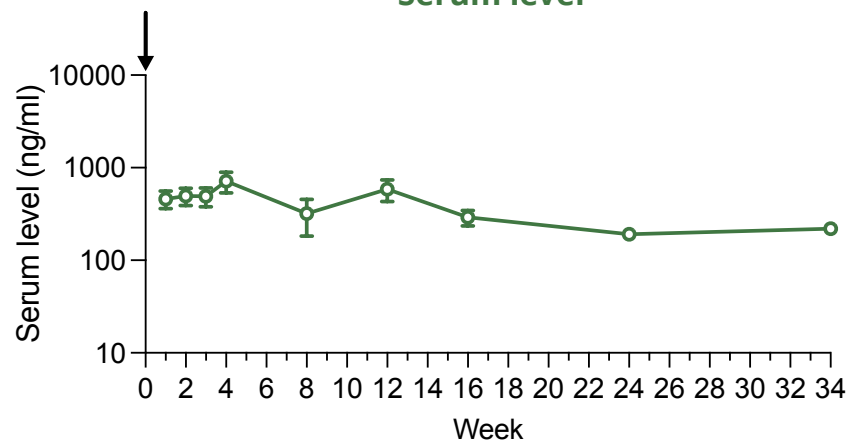
Enhanced BBB targeting dramatically improves GLP-1 Fc fusion efficacy



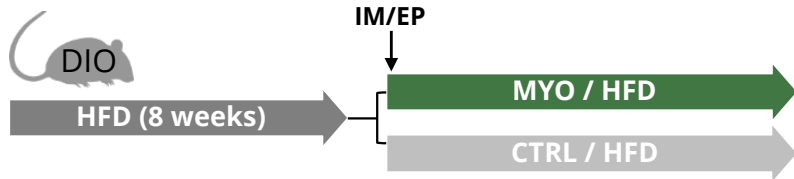
Weight loss



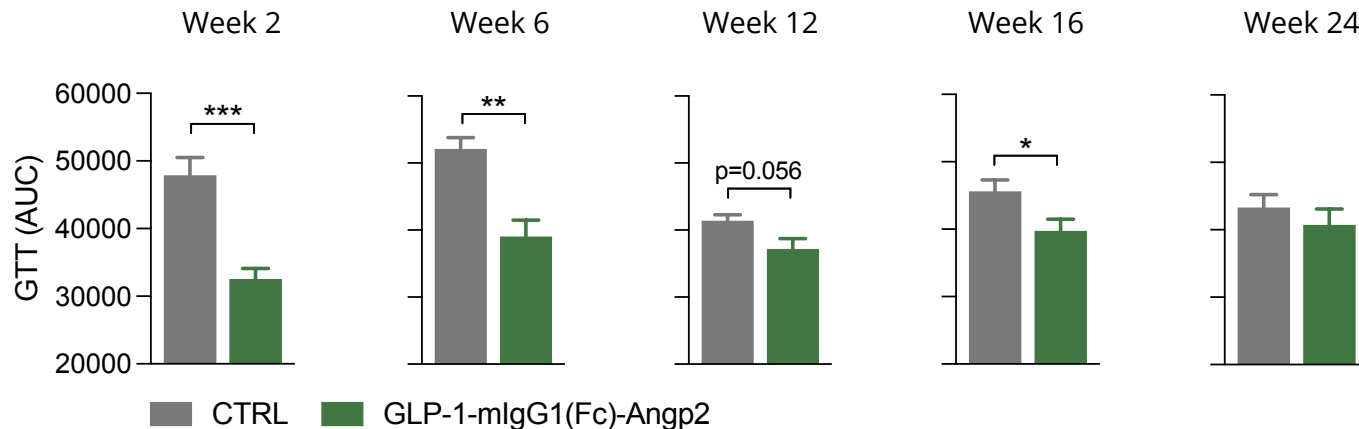
Serum level



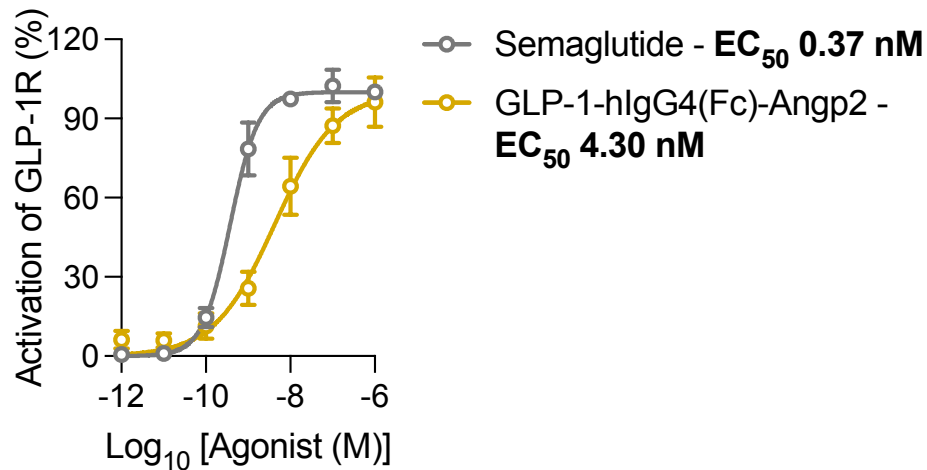
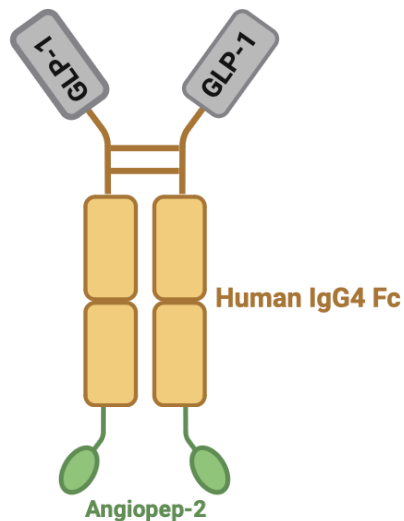
Enhanced BBB targeting dramatically improves GLP-1 Fc fusion efficacy



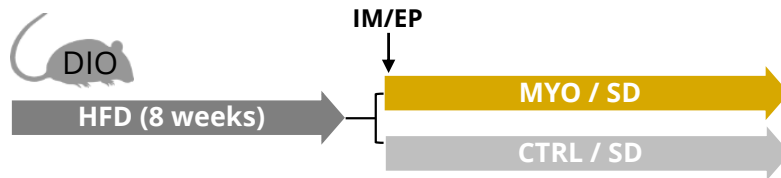
Glucose clearance



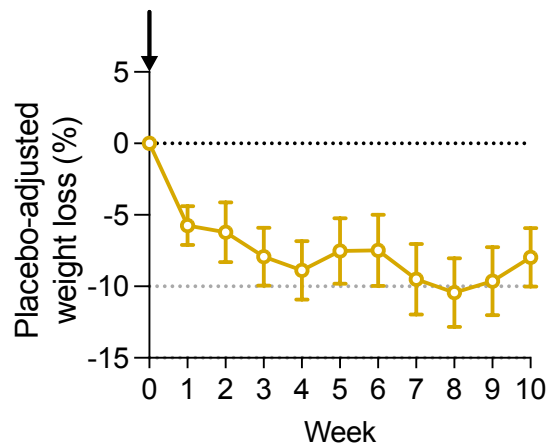
Human Fc fusion is also efficacious in a DIO model



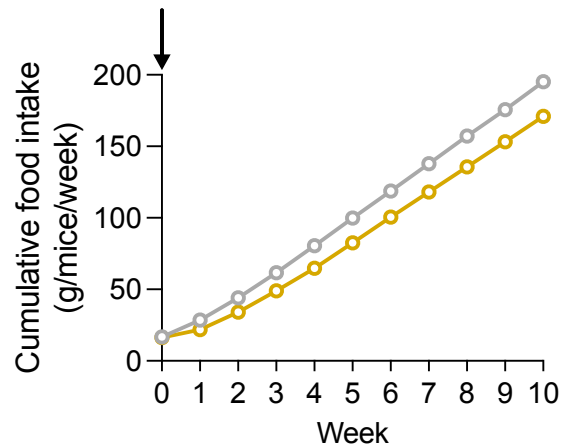
Human Fc fusion is also efficacious in a DIO model



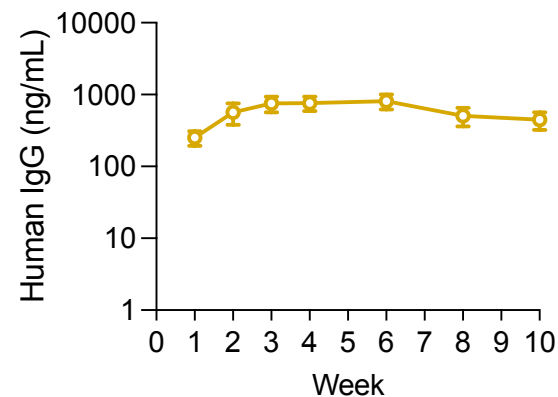
Weight Loss



Food Intake

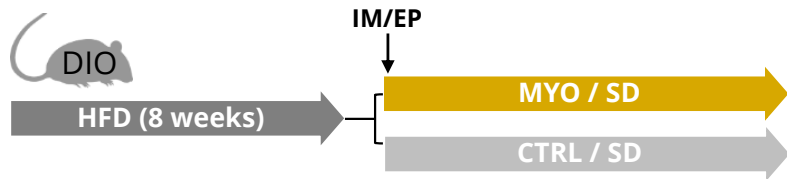


Serum level

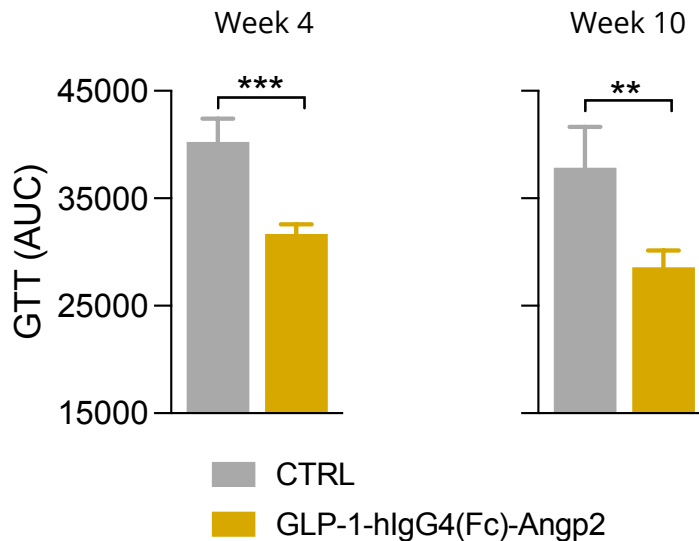


—○— CTRL —○— GLP-1-hIgG4(Fc)-Angp2

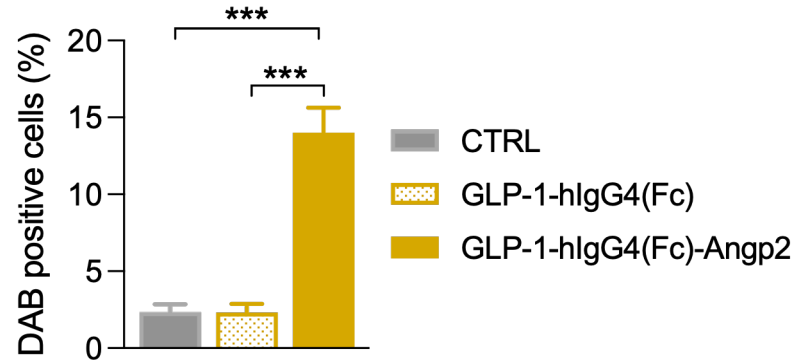
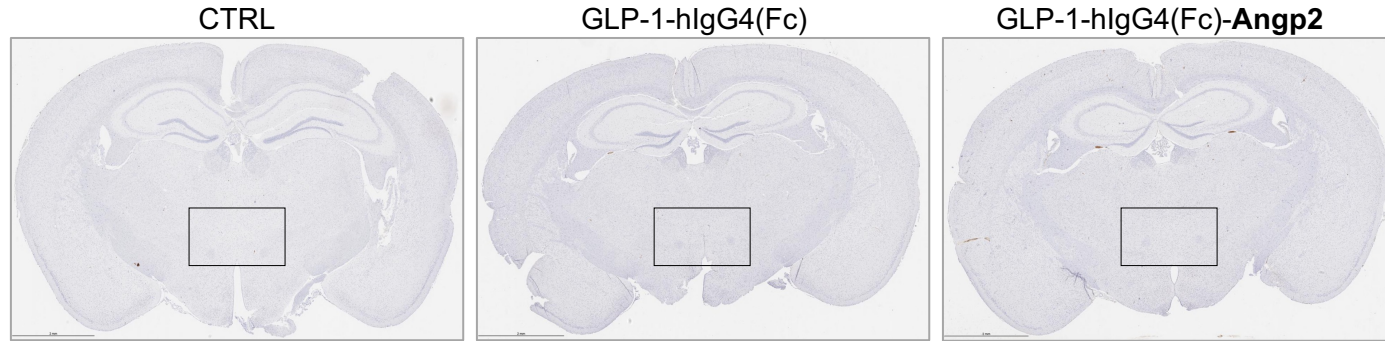
Human Fc fusion demonstrates similar effects in a DIO model



Glucose clearance



Angiopep-2 enhances BBB penetration of GLP-1 Fc fusion



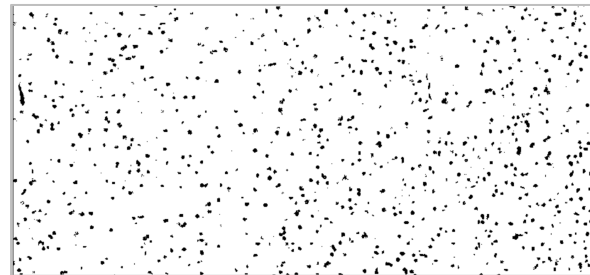
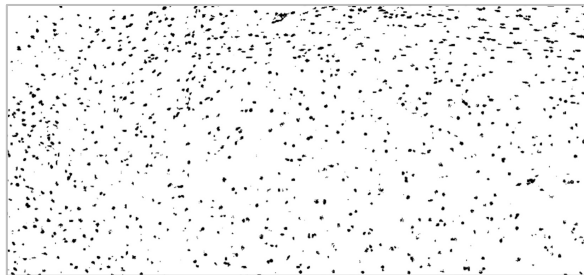
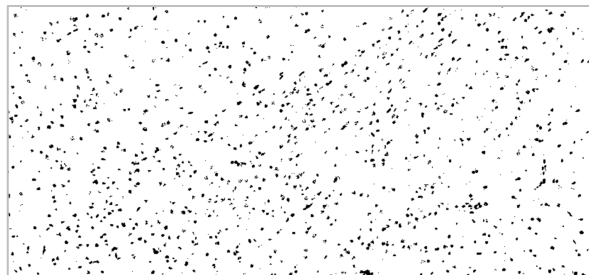
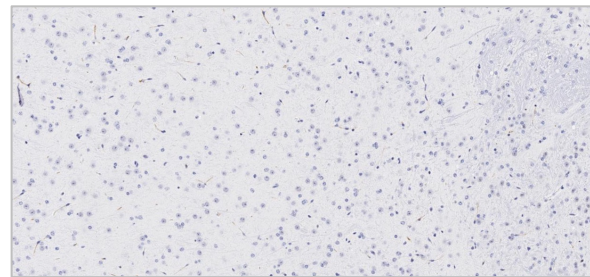
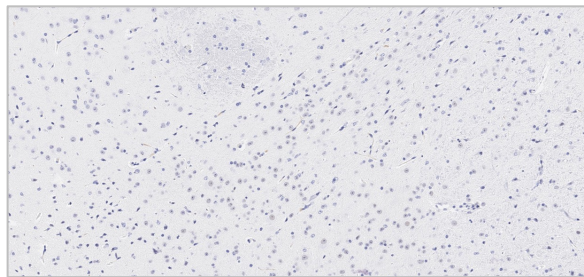
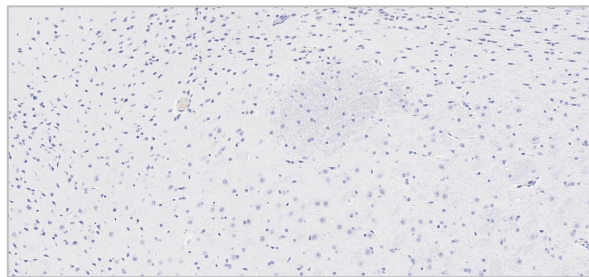
CTRL

GLP-1-hIgG4(Fc)

GLP-1-hIgG4(Fc)-Angp2

Hematoxylin

DAB



DNA-Encoded Incretin Receptor Agonists

- Durability:
 - Ultra long-term production of IRA and efficacy
- Flexibility:
 - Simultaneous or sequential delivery of multiple molecules
- Scale:
 - DNA manufacturing and less frequent administrations



Acknowledgements



Linda Sasset, PhD

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**Andy Thompson*

**Robert Miller*

**Delcora Campbell, MS, MBA*



BILL & MELINDA
GATES foundation



Empire State
Development

Thank you

