

PepVax, Inc. has developed a novel DNA-based drug delivery and development platform, called SMARTmid™, to make safer and more effective autologous drugs using a "Trojan Horse" approach

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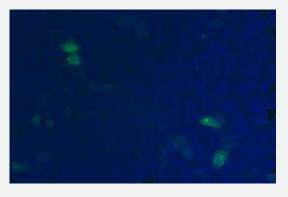
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Overview

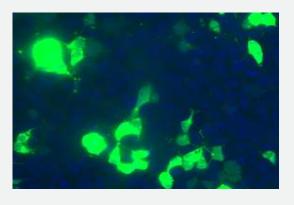
PepVax, Inc. has developed a potent drug delivery system for nucleic and amino acid-based (NAA) drugs called SMARTmidTM

- Can "manufacture" required proteins, antibodies, and DNA/RNA inside patient.
- Can boost immune response for existing and newer technologies
- Regulatory and Capital efficent

Seed: \$2 MM to finish development and establish collaborations



Commercial Plasmid



PepVax Plasmid

A comparative analysis

Current precision medicine approaches have problems.

SMARTmid™: Autologous

PepVax DNA Plasmid Technology

Desired NAAs "manufactured" inside the patient

Potential to be more effective

Broad applicability

Current approaches: Allogeneic

In Precision Medicine

Cell Therapies

Gene Therapies

Immunotherapies

Problems:

Long production timeline

High toxicities

Low efficacy rates

Narrow applicability

Our Technology: SMARTmid™ DNA

NAA

We can add sequences for any NAA of interest to "manufacture" or create immune responses for boosts, such as vaccines. 1 3

Protein Signaling

In-built Protein
Transportation Signaling
provides a holistic immune
response through both Tcells and B-cell antibodies.

Nuclear Localization

SMARTmid™ DNA plasmid technology will localize the treatment for increased effectiveness with lower doses and decreased the side effects.

Inbuilt Adjuvant

In-built adjuvant enhances the immune reaction by attracting the right tumortargeting cells.

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2

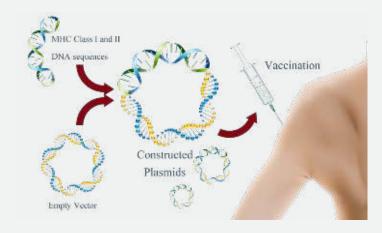
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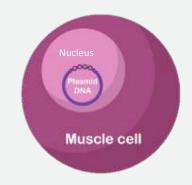
SMARTmid™ Drives Strong Expression

1



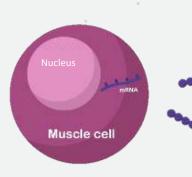
Designed SMARTmid™ DNA vectors delivered intramuscularly for systemic response or intravenously for targeted response

2



Plasmid DNA taken into cells. *Process* can stop here for delivery.

3



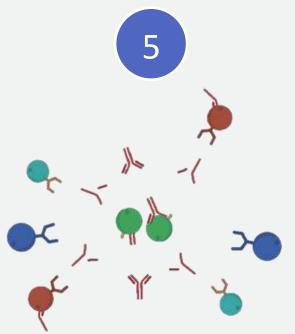
Plasmid DNA converted to target proteins for expression. *Process can stop here for "manufacturing"*

SMARTmid™ Drives Strong Immune Response

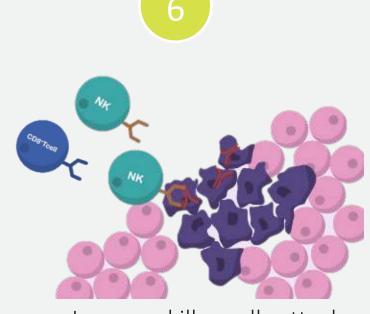
Against Disease Targets



Immune cells recognize proteins as foreign and prepare to attack it



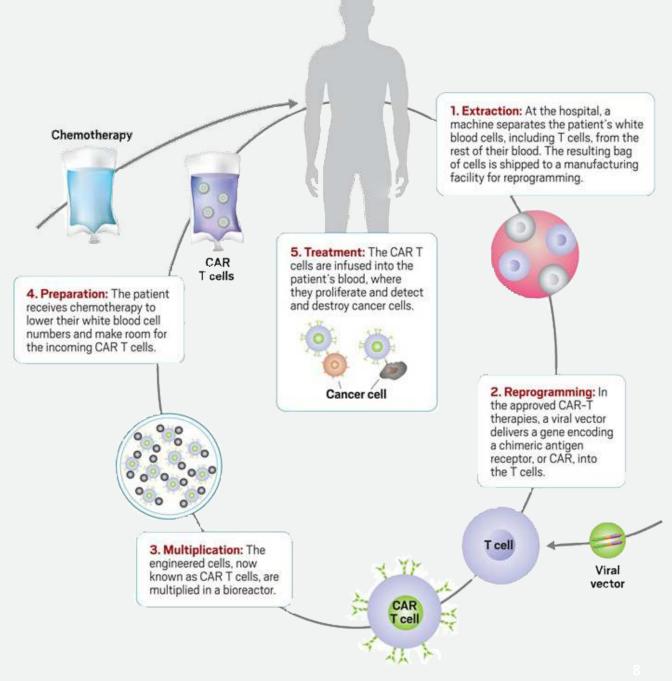
Strong immune response is generated against protein target throughout the body



Immune killer cells attack antigen-positive diseased cells and destroy them

Example: Allogeneic CAR-T

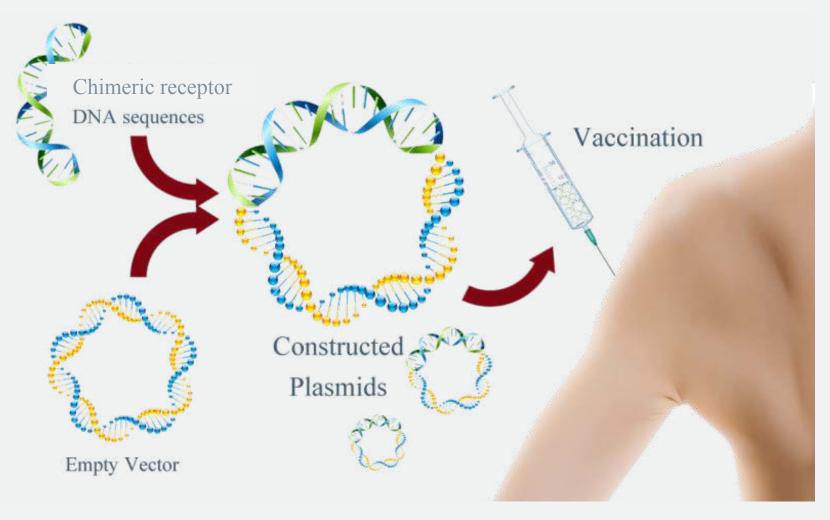
The current approach for CAR-T development is highly invasive, painful, and has a long production timeline for ex vivo engineering



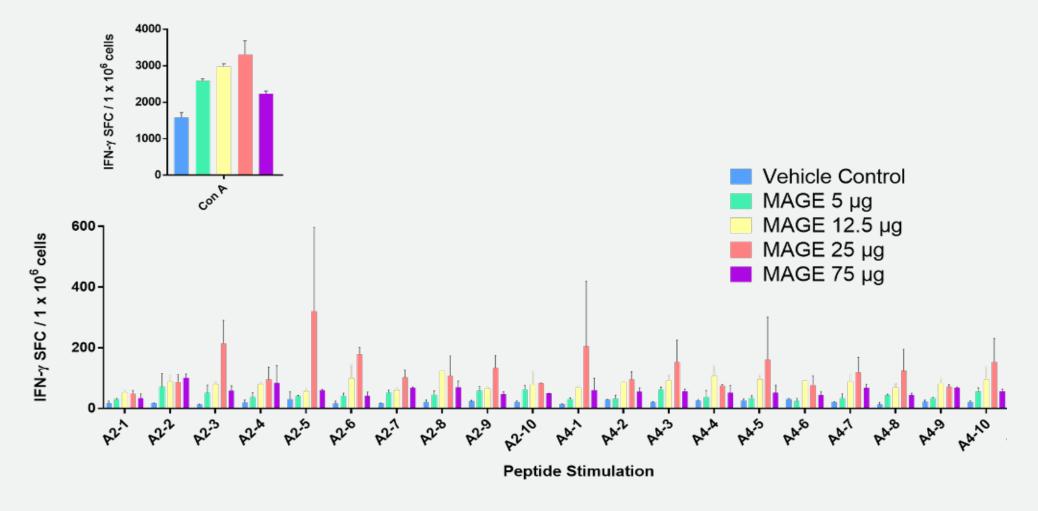
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Example: Autologous CAR-T

Instead of using viral vectors to transfect T-cells from patients, we can use SMARTmid™ to transfect T-cells directly into the patient's lymph nodes, making the process safer and more effective.

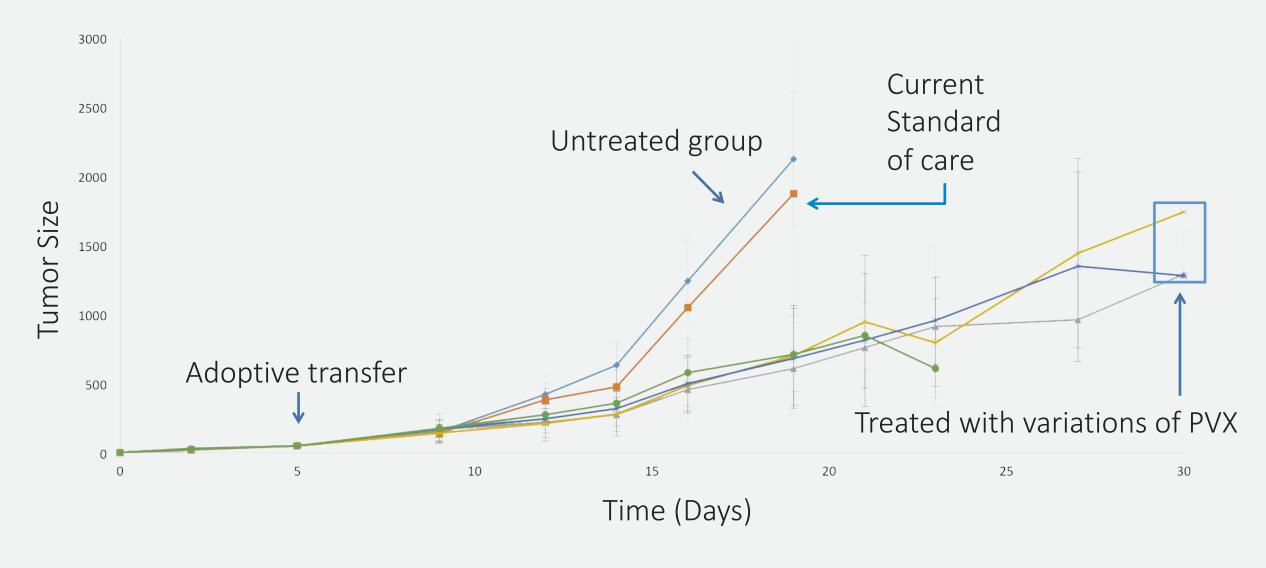


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Proof of Concept accomplished: Strong CAR-T cell Production using SMARTmid™

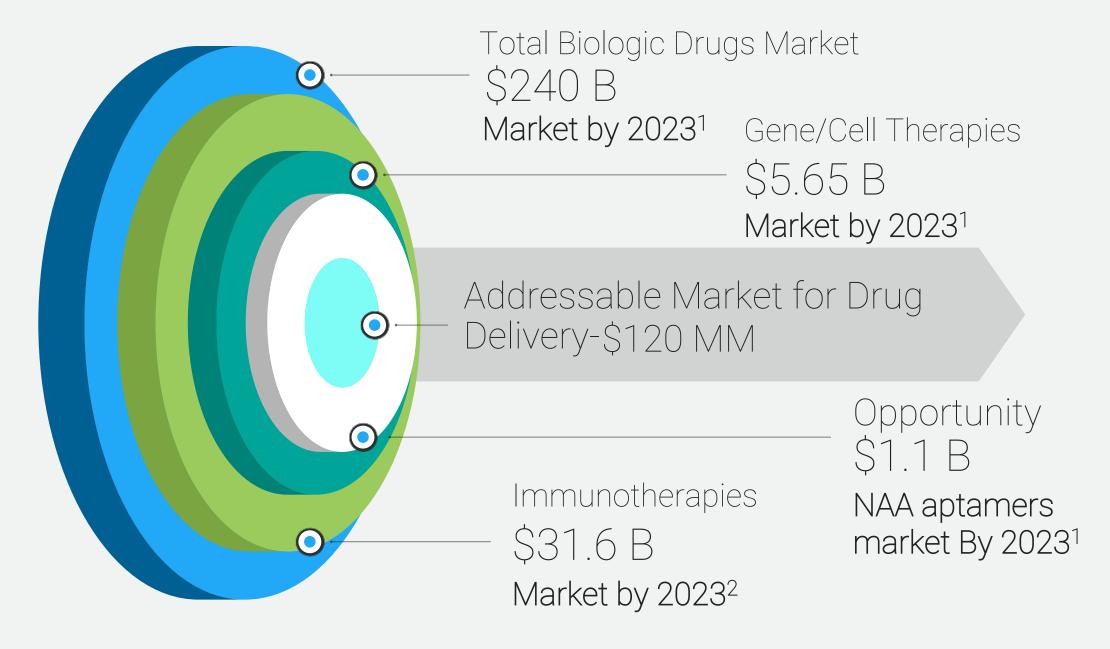
CAR-T production for MAGE A took only 2 weeks instead of 2 months using autologous developmental methods.



Proof of Concept: Anti-tumor activity using SMARTmid™ In aggressive triple-negative breast cancer, we showed a 45% reduction in tumor growth using autologous CAR-T cells from previous experiment

Advantages

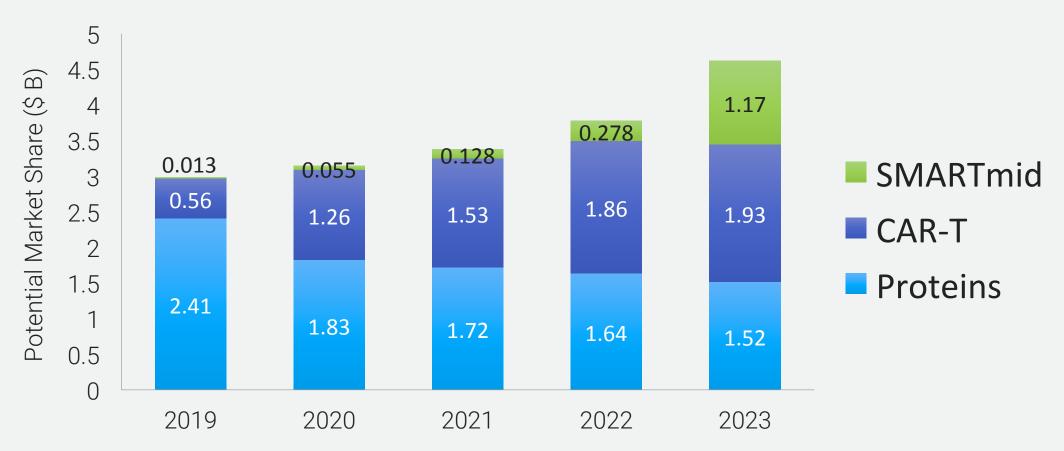
Vectors	Expression	Safety	Stability	Adjuvant	Versatility
viral		X	X	X	X
mRNA			X		X
Current DNA				X	X
SMARTmid™					



¹Coherent Market Insights Reports (2018)

²Immune Checkpoint Inhibitors Market: Allied Market Research(2018)

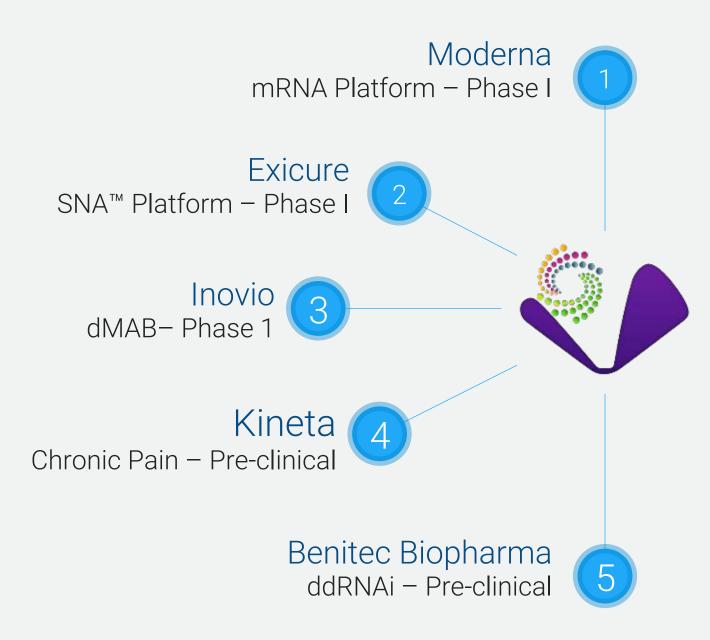
Expected NAA Market Share Growth



As we continue to grow and work with more companies to develop future applications, the overall research market share of immunotherapy and gene therapy will not only expand to approximately \$5 billion, but our own market potential will continue to grow to over \$1.15 billion by 2023.

¹Coherent Market Insights Reports (2018)

²Immune Checkpoint Inhibitors Market: Allied Market Research(2018)



Competitive Landscape

Key Differentiator: SMARTmid™ takes the complications of manufacturing and delivery and brings it into one simple system that has the versatility to be used in gene-, cell- and immunotherapy.

Protection: 1 patent-pending, 2 provisional patents in-house

Go-to-Market Strategy

Pricing and Services

Collaborations

with Universities

Partnerships

With Startups and smaller firms

Licensing

to large pharma

\$10-20K

- Grant submission
- Early research
- O Build IP Portfolio
- Out-license

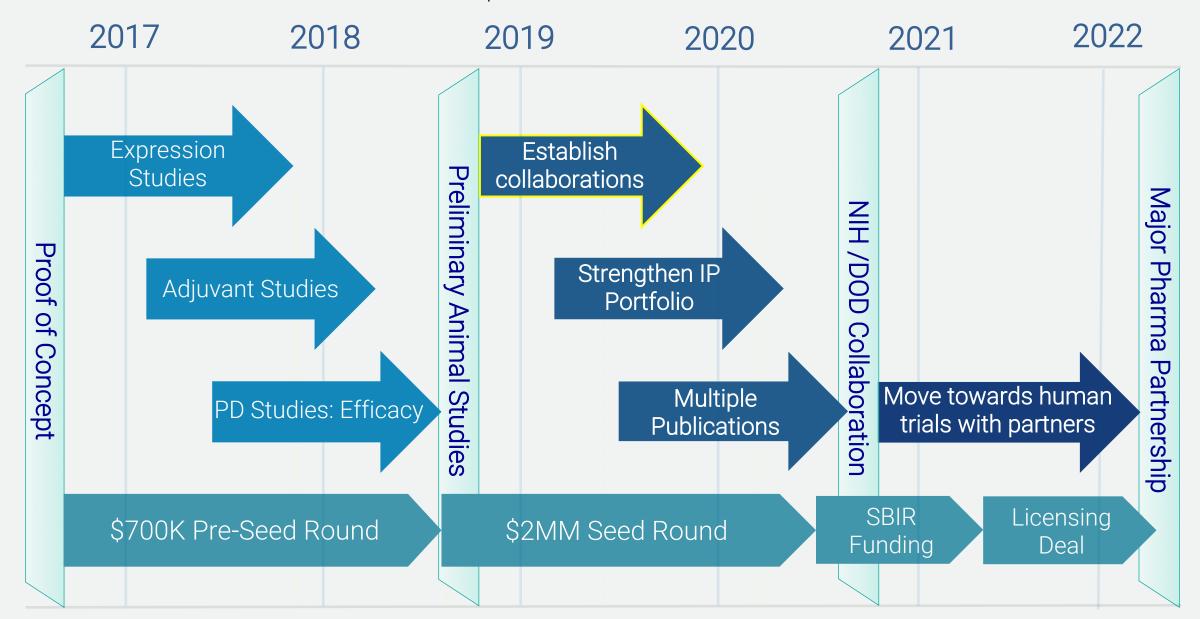
\$25-100K

- O Collaborations
- O Animal studies
- O IND Submission
- O Milestone payments
- O Royalties

\$5-10MM

- Human trials
- O NDA submission
- Commercialization
- Milestone payments
- Royalties

Development Timeline



STARTING POINT

Speratum

Collaboration and licensing for siRNA delivery

Loma Linda University

Collaboration for gene and siRNA delivery

Howard University

Collaboration for small molecule/nanoparticle delivery

FUTURE PLAN

Novartis

Collaboration and licensing for biologics delivery

Souzhou Ribo Life Sciences

Licensing for nucleic acid and RNAi delivery

Chimeron Bio

Licensing for personalized cancer gene therapy

Go-to-Market Efficiency

Regulatory and Capital

FDA	Reimbursement	Capital	
Drug-Device Combination	Bundled Payments	\$700K To date	
 Through partners Focus on CMC Aid in IND-filing	 CPT codes: 96413 or 96365 Coverage for breakthrough and replacing technologies 	 Completed POC animals studies Identified Regulatory pathway Generating revenue 	

Financial Data

FY*	2019	2020	2021	2022	2023
Revenue	\$132,168	\$545,352	\$1,159,473	\$2,677,730	\$11,572,976
COGS	\$26,026	\$31,231	\$37,477	\$44,973	\$53,968
Expenditure	\$571,521	\$731,640	\$813,740	\$1,060,434	\$1,294,871
EBITA	(\$465,379)	(\$217,519)	\$308,255	\$1,572,324	\$10,224,138

^{*}PepVax' fiscal year begins in September of previous year (FY 2019 starts in Sept. 2018)
FY 2019 assumes the start of GMP manufacturing and collaboration research costs. FY 2020-2022 assumes regulatory operation, increase in sales force, along with upfront cash flow with licensing deal for SMARTmid™ delivery system.

¹FY 2023 revenue reflects recent licensing done by similar stage company at pre-clinical (Kineta/Genetech Deal worth up to \$359 MM in 2018).

Our Team

40+ years of combined experience in biotechnology



Mahesh Narayanan, MS Research and Business, FTE







Anton
Dormer, MD, MS
R&D, FTE







Elton Norman, Esq., CPA Finance, PTE







Daniel Achinko, PhD R&D, FTE





Our Advisors

25+ IND Filings, 6 NDAs and multiple commercialized products

Advisory Board

PepVax advisory team brings the experience to develop our technology past the regulatory approval process into commercialization



Sunil Joshi Business Development



Stephen Popielarski Business Development



George Moonsammy Regulatory



Angela Lynch Toxicology



Salvator DeSena Medical Affairs





















Ask

Founder + Family + Friends: \$700K

Current Round: Seed

\$2 MM

Convertible Note / Equity

Proceeds: \$750K towards current operations; \$1.25 MM towards future development of platform, expand intellectual property, and establish strong collaborations

Contact Us





















Mahesh Narayanan Primary contact



angel.co/pepvax



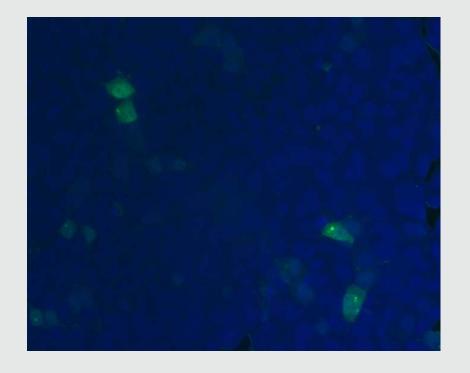
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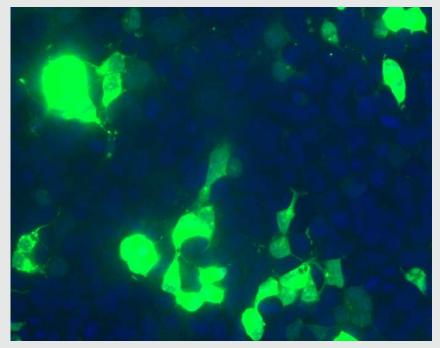


(856) 343-2804



Appendix





Green Florescent Protein (GFP) expression using best commercially available plasmid.

GFP expression using SMARTmid™ DNA shows much greater expression at same concentration of DNA

High protein expression using SMARTmid™