

Advancing Immunotherapy & Prevention

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Certain statements in this presentation may constitute "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act. These statements are based on management's current expectations and are subject to uncertainty and changes in circumstances. Actual results may differ materially from those included in these statements due to a variety of factors, including whether: GeoVax can develop and manufacture its vaccines with the desired characteristics in a timely manner, GeoVax's vaccines will be safe for human use, GeoVax's vaccines will effectively prevent targeted infections in humans, GeoVax's vaccines will receive regulatory approvals necessary to be licensed and marketed, GeoVax raises required capital to complete vaccine development, there is development of competitive products that may be more effective or easier to use than GeoVax's products, GeoVax will be able to enter into favorable manufacturing and distribution agreements, and other factors, over which GeoVax has no control. GeoVax assumes no obligation to update these forward-looking statements, and does not intend to do so. More information about these factors is contained in GeoVax's filings with the Securities and Exchange Commission including those set forth at "Risk Factors" in GeoVax's Form 10-K.



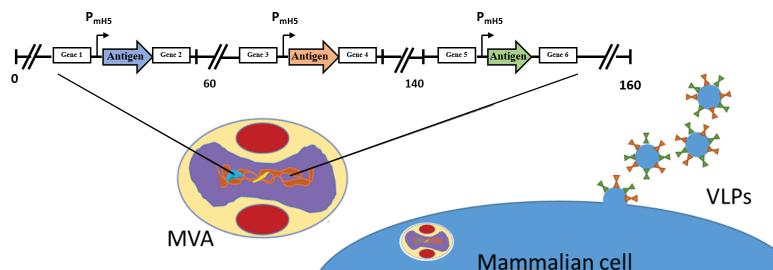
GeoVax...A Compelling Opportunity

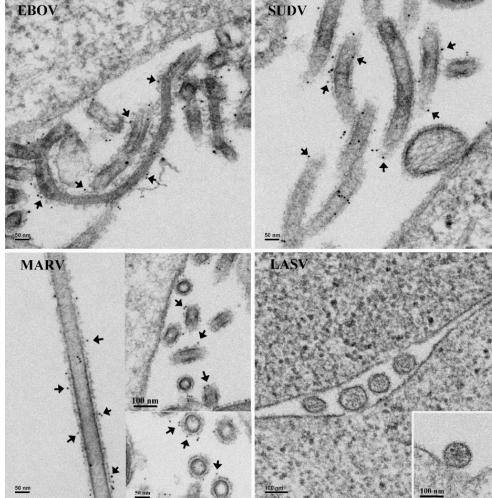
- Developing immunotherapies and infectious disease prevention based on
 - Single-dose
 - Durable immunity
 - No adjuvant
 - Extensive safety profile
 - Cost-effective manufacturing
- Leveraging internal and external expertise to minimize development risk

GeoVax

GeoVax MVA-VLP Vaccine Technology

- ► MVA Vector = Safety
 - Replication-deficient in mammalian cells → Vaccination
 - Replication competent in avian cells → Manufacturing
- ► Transgenes: Antigen = Immunogenicity
- ► Vector + Antigen = MVA-(VLP) Platform





Advantages of MVA-VLP vaccines



- ► Inherently safe
- ► Non-replicating in host
- Potential for single dose protective immunity
- ▶ Not hindered by pre-existing immunity to vector
- Self-adjuvant
- Thermostable > can be lyophilized
- Established methods for manufacturing

















Advantages of GeoVax Technology

Technology	Single-Dose Immunity	Immunogenicity	Optimized VLP Formation	Transgene Stability
GeoVax MVA-VLP	Yes	Excellent	Yes	Excellent
Competitive MVA Technologies	No 🗸	Limited 🗸	No 🗸	??Unknown?? 🗸

Cost-Effective Manufacturing and No Adjuvant Necessary



Broadly Validated Technology

		Target Identification	Preclinical Validation	Clinical Trials	
				Phase 1	Phase 2
Immuno-Oncology					
Solid Tumors					
HPV-associated Head and Neck Cancer					
Infectious Disease					
HIV (HVTN)	GOVX-B11				
HIV (functional cure, AGT)	GOVX-B01				
Lassa Fever	GEO-LM01				
Ebola, Marburg, Sudan	GEO-EM01				
Zika Virus	GEO-ZM02				
Malaria					
Hepatitis B (chronic infection)					



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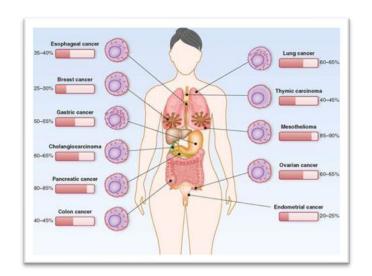


GeoVax Cancer Immunotherapy Focus

GeoVax has developed a combination cancer vaccine strategy:

- GeoVax novel Cancer Immunotherapy uses combinations of:
 - MVA-VLP cancer vaccines
 - Select proteins, peptides (e.g., MUC1; Cyclin B1),
 - Immune check-point inhibitors (e.g., anti-PD1)

For treating and potentially preventing solid tumors.



GeoVax Concept to Cancer Immunotherapy





STIMULATE

MVA-VLP with TAA to provoke immune system (POC completed)



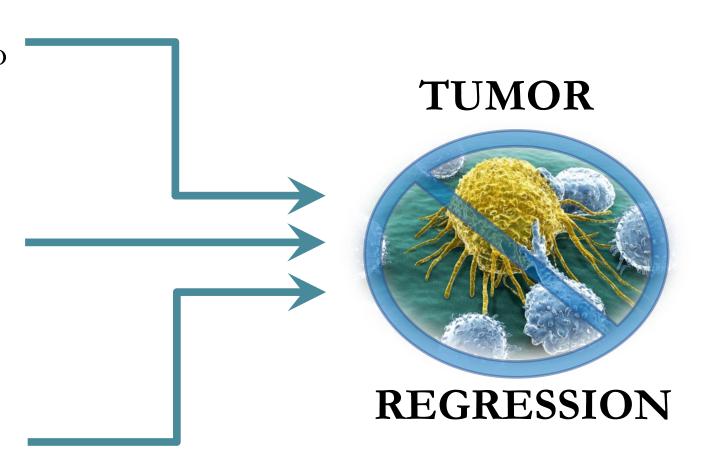
BLOCK

checkpoint inhibitor to reverse immune tolerance (POC completed)



KILL

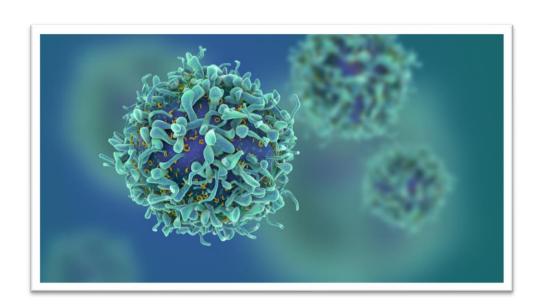
achieve oncolysis using armed vaccinia virus



GeoVax Cancer Vaccines



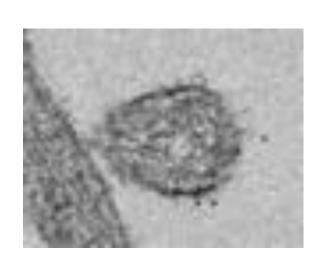
- ► MUC1 aberrantly glycosylated in many cancers
 - In collaboration with ViaMune, University of North Carolina at Charlotte & University of Pittsburgh
- CyclinB1 overexpressed in certain cancers
 - In collaboration with Vaxeal
- ► HPV head and neck cancers
 - Internal
 - E6/E7 antigens

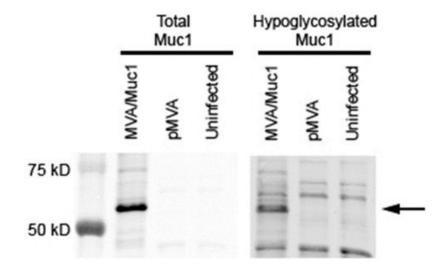


MVA-VLP-MUC1



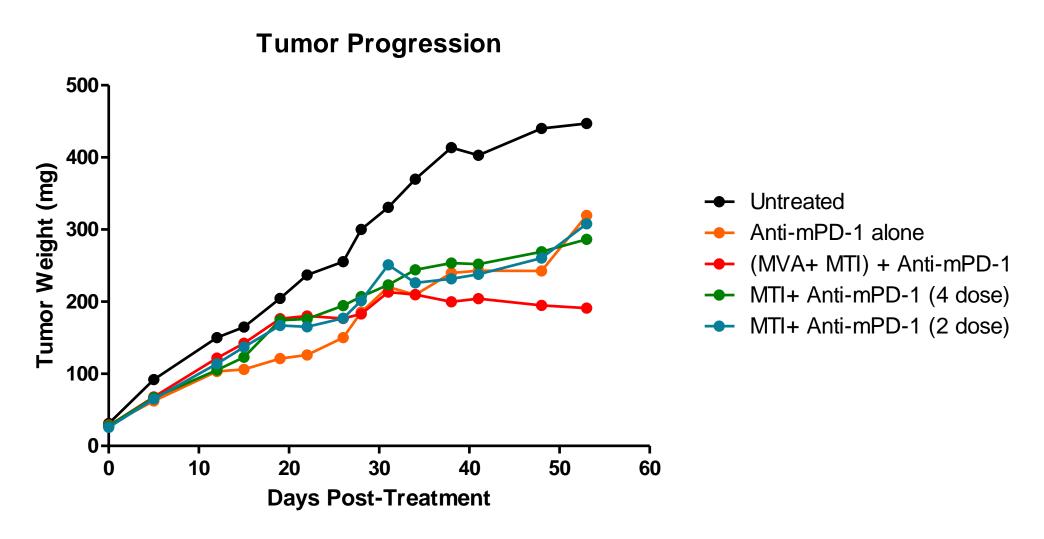
- ▶ MUC1 engineered to be displayed on VLPs produced in vivo
- ▶ 5 repeats of the VNTR region of MUC1
- Overexpression of MUC1 by MVA leads to accumulation of hypoglycosylated protein
- ► Potentially useful against all MUC1+ cancers
 - Pancreatic
 - Colorectal
 - Prostate
 - Lung
 - Breast
 - Others







Therapeutic Experiment Results

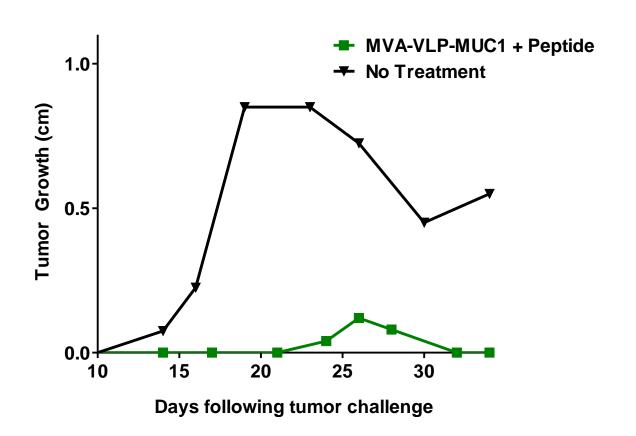


As in the initial experiment, MVA+MTI+CPI arrested tumor growth and shrank tumors

Prevention Experiment Results



- MUC1 Tumor Associated Antigen (TAA):
 - Combination therapy:
 - MVA-VLP-MUC1 and MUC1 peptide
 - GeoVax MVA-VLP combination achieved tumor prevention
- ► Indications:
 - Lung, colorectal, breast, gastric, prostate, colon, pancreas, etc.





GeoVax Cancer Immunotherapy Next Steps

- Further validate GeoVax novel cancer vaccine approach using other target antigens & novel check-point inhibitors that can be expressed by the vector and not administered separately
- Complete POC related to using oncolytic virus to achieve tumor cell lysis (e.g. KILLing the tumor)
- ▶ Proceed to clinical trials and technology validation



GeoVax Developments in Infectious Disease Vaccines

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GeoVax HIV Preventive Vaccine Advancing in Human Trials

- ▶ Excellent safety and immunogenicity: Phase 1 & 2a human clinical trials
- Superior antibody profile and durability compared to Sanofi Thailand
 Phase 3 clinical trial (RV144)
- Preventive vaccine: HVTN 132 Scheduled start H2 2019
- ► Therapeutic vaccine: Phase 1 trial to begin H2-2019
 - Collaboration with American Gene Technologies for "functional cure" effort;









Hemorrhagic Fever Vaccines

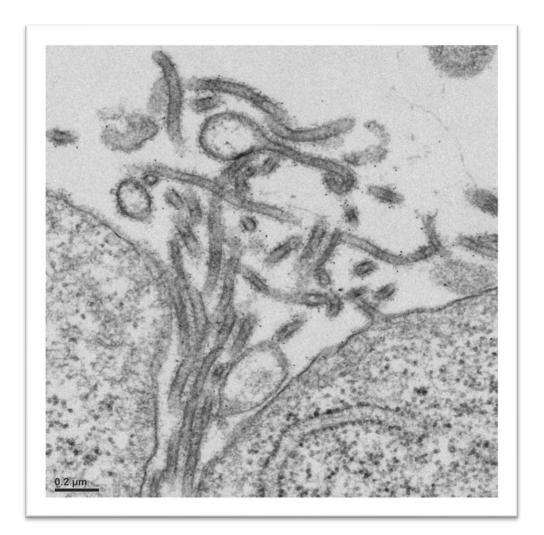
- Four vaccines covering all major hemorrhagic fevers:
 - Ebola
 - Sudan
 - Marburg
 - Lassa

Zika Vaccine (GOVX-ZM01)

 A novel vaccine candidate that removes risk of ADE

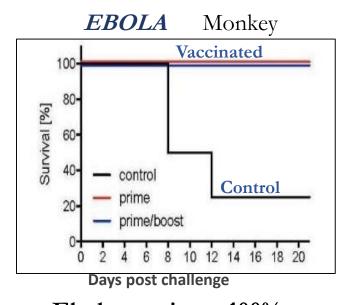
Malaria Program

• Vaccine antigens covering all 3 stages of malaria infection

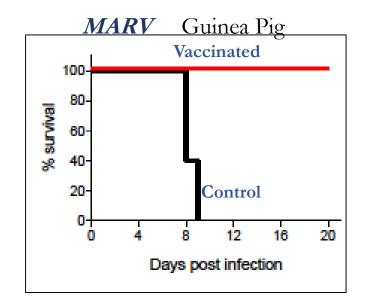


Hemorrhagic Fever Vaccines-Unparalleled Success

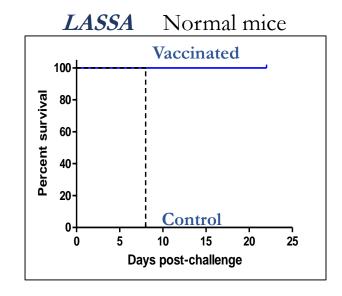




- Ebola vaccine 100%Protection,Single-Dose
 - In 2 rodent models and Nonhuman Primates
 - Anticipating govt support (NIH, BARDA)



- Marburg vaccine –100% Protection,
 - In Guinea Pig model
 - Anticipating govt support (NIH, BARDA)



- Lassa vaccine 100%Protection,Single-Dose
 - In a lethal (intracerebral inoculation) mouse model
 - Ongoing Govt. Support
 - Fast-Track Phase I / II
 SBIR grant from NIH
 - CDMRP grant from DoD for production of Master Seed Virus

Sudan Vaccine, awaiting animal testing

Potential to develop a single tetravalent vaccine \rightarrow Significant commercial opportunity

Zika Vaccine Program - Progressing to Clinic



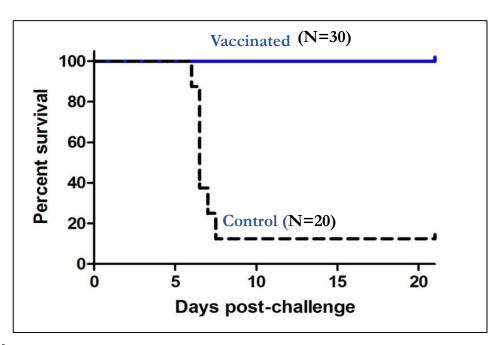
► Novel vaccine design

- Competitive Advantage
 - No risk of Antibody Dependent Enhancement (ADE) of infection against Dengue or other flaviviruses

Excellent preclinical data

- Immunocompetent outbred mice vaccinated IM and challenged IC (very rigorous model)
- 100% single-dose survival in vaccinated mice
- 90-100% death in saline controls within 1 week
- Effective clearance no virus recovered from brains of vaccinated and Zika-challenged mice

 Completed Immunogenicity and Efficacy studies in Nonhuman Primates with NIH grant support (Nature's Scientific Reports 2017)



Strong IP Protection



Patent Portfolio

- Owner of Record Patents
 - − U.S. − 11 applications pending
 - − Non-U.S. − 17 application pending
- Licensed Patents
 - − U.S. − 10 issued, 2 applications pending
 - − Non-U.S. − 14 issued, 2 applications pending



GeoVax Value → Proprietary Expertise in Vaccine Vector Development

GeoVax Scientific Advisors



Thomas Monath, MD (Chairman)

- Managing Director & CSO, Crozet BioPharma
- Acambis, Hookipa, PaxVax, NewLink Genetics, PerkinElmer

Olja Finn, PhD

Prof. Immunology and Surgery, Univ. of Pittsburgh

Barney Graham, MD, PhD

• Sr. Investigator, VRC, NIAID/NIH

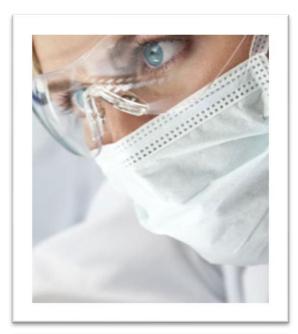
Stanley Plotkin, MD

- Vaxconsult, LLC
- Sanofi Pasteur

Scott Weaver, PhD

Director UTMB Institute for Human Infections and Immunity







A Compelling Opportunity

- ✓ Developing novel, patented vaccines and immunotherapies based on a unique technology & expertise
 - *Robust, durable immunity, single-dose, no adjuvant, costeffective manufacturing, safe vaccines
- ✓ Focused on critical major medical needs with differentiated advantages
- ✓ Translational programs underway to pursue clinical and regulatory development



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