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Generation and characterization of broadly neutralizing antibodies against diverse Henipaviruses

Project Number	Contact PI/Project Leader	Awardee Organization
5F31AI133943-02	AZARM, KRISTOPHER	ICAHN SCHOOL OF MEDICINE AT MOUNT SINAI

Description

Abstract Text

PROJECT SUMMARY The recent global emergence of Henipaviruses (HNVs) into human populations changes the risk calculus associated with the species. We see a specific need for a vaccine strategy and passive immunotherapies that will protect not only against HNVs that have previously emerged, but also diverse HNVs should zoonotic spillover and viral emergence occur from one the many divergent HNVs. The objectives of this proposal are to leverage our expertise on the functional and antigenic diversity of the HNV surface glycoproteins in designing a rational vaccine strategy against HNVs and to better understand the antibody response to the HNV glycoproteins. We do so by 1) performing a vaccination strategy that will exploit the conserved mode of host receptor ephrinB2 interaction amongst the diverse spectrum of HNV receptor attachment glycoproteins in order to generate broadly neutralizing antibodies and by 2) characterizing the conserved, vulnerable epitopes these antibodies target in the surface glycoproteins. We will use an array of virological, immunological, structural, and biochemical methods to address these aims. Combined, these approaches will generate novel broadly neutralizing antibodies against the ever-expanding spectrum of HNVs and reveal conserved regions of the HNV glycoproteins that can be exploited in future vaccination strategies.

Public Health Relevance Statement

PROJECT NARRATIVE Henipaviruses are a rising global threat, as evidenced by the World Health Organization's inclusion of Nipah virus on a 2015 list of the top emerging diseases likely to cause major epidemics. The potential for Henipavirus spillover into human populations underscores the need for broadly neutralizing antibodies that can be used in passive immunotherapy and for vaccine strategies that will elicit broad protection in high-risk areas, e.g. rural areas of Southeast Asia, Australia, Africa, etc. Through the studies proposed in this application, we will perform a rational vaccine design strategy to generate broadly neutralizing antibodies against Henipaviruses to fulfill this specific need.

NIH Spending Category

Biodefense	Biotechnology	Emerging Infectious Diseases	Immunization
Infectious Diseases	Prevention	Vaccine Related	

Project Terms










Address	Africa	African	Animals	Antibodies	Antibody Response	
Antigenic Diversity		Area	Australia	Binding	Binding Sites	Biochemical
Biological Assay	Calculi	Cell Separation	Cell surface	Cells	Chiroptera	
Complex	Crystallization	DNA	Development	Disease	Distal	
Electroporation	Epidemic	Epitopes	Escape Mutant	Exposure to	Family	
Fruit	Future	GTP-Binding Proteins	Generations	Ghana	Glycoproteins	
Head	Hendra Virus	Henipavirus	Human	Immune	Immune response	
Immunize	Immunologics	Inbred BALB C Mice	Lead	Maps	Membrane	

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Organization

Name ICAHN SCHOOL OF MEDICINE AT MOUNT SINAI	Department Type MICROBIOLOGY/IMMUN/VIROLO	State Code NY
City NEW YORK	Organization Type SCHOOLS OF MEDICINE	Congressional District 13
Country UNITED STATES (US)		

Other Information

FOA PA-16-309	Administering Institutes or Centers NATIONAL INSTITUTE OF ALLERGY AND INFECTIOUS DISEASES	Project Start Date 01-April-2018
Study Section Special Emphasis Panel[ZRG1-F13-Z(20)L]	DUNS Number 078861598	Project End Date 31-March-2020
Award Notice Date 21-March-2019	CFDA Code 855	Budget Start Date 01-April-2019
		Budget End Date 31-March-2020

Project Funding Information for 2019

Total Funding \$45,016	Direct Costs \$45,016	Indirect Costs \$0
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Year	Funding IC
2019	NATIONAL INSTITUTE OF ALLERGY AND INFECTIOUS DISEASES

NIH Categorical Spending


[Click here for more information on NIH Categorical Spending](#)


Funding IC	FY Total Cost by IC	NIH Spending Category
NATIONAL INSTITUTE OF ALLERGY AND INFECTIOUS DISEASES	\$45,016	Biodefense; Biotechnology; Emerging Infectious Diseases; Immunization; Infectious Diseases; Prevention; Vaccine Related;


Sub Projects


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Patents

No Patents information available for 5F31AI133943-02



Outcomes

The Project Outcomes shown here are displayed verbatim as submitted by the Principal Investigator (PI) for this award. Any opinions, findings, and conclusions or recommendations expressed are those of the PI and do not necessarily reflect the views of the National Institutes of Health. NIH has not endorsed the content below.

No Outcomes available for 5F31AI133943-02



Clinical Studies

No Clinical Studies information available for 5F31AI133943-02



News and More

Related News Releases

No news release information available for 5F31AI133943-02



History

No Historical information available for 5F31AI133943-02



Similar Projects

No Similar Projects information available for 5F31AI133943-02

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