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5R21AI135373-02

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Development of a subunit vaccine for MERS-CoV and other emerging coronaviruses

Contact PI/Project Leader WHITTAKER, GARY R

Awardee Organization CORNELL UNIVERSITY

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Abstract Text

Project Summary/Abstract We propose to develop a subunit vaccine for MERS-CoV, using a structure-based approach targeting conserved and functionally essential domains in the stalk region of the viral spike protein. MERS-CoV itself is a currently emerging virus in humans with strong links to camels, but an as-yet unproven animal reservoir. The most closely related viruses to MERS-CoV are found in **bats**. We expect our vaccine platform to be applicable and effective across a wide range of existing and emerging coronaviruses. As our system is based on expression in E. coli is it expected to be cost-effective, and our stalk-based approach is specifically designed to cover a range of distinct coronaviruses. However, it is important to note that our vaccine platform is highly flexible, with the antigen able to be re-engineered rapidly in the face of a novel coronavirus that may emerge, and for which the vaccine developed in this application is not effective.

Public Health Relevance Statement

Project Narrative We propose to develop a novel, cost-effective, and highly flexible vaccine platform to immunize against MERS- CoV. We will use an innovative subunit vaccine design approach, based on conserved epitopes in the coronavirus spike S2 fusion domain, express the vaccine platform in E. coli, and test both the serological and protective response in a mouse model. Due to the conserved nature of the structural region targeted, we anticipate that our stalk-based approach will be applicable to and effective across a wide range of existing and emerging coronaviruses.

NIH Spending Category

Biodefense Biotechnology Emerging Infectious Diseases Immunization

Infectious Diseases Orphan Drug Prevention Rare Diseases Vaccine Related

Project Terms

Animals Antigens Camels Chiroptera Coronavirus **Cryoelectron Microscopy** Development **Engineering Epitopes** Escherichia coli Human Immune response Link Goals **Immunize** Middle East Respiratory Syndrome Coronavirus Mus **Nature Proteins** SARS coronavirus Serological Structural Models **Structure Subunit Vaccines** System **Testing Transgenic Mice Vaccine Antigen Vaccination Vaccine Design** Viral Wild Type Mouse **Vaccines** Virus base cost effective design flexibility model building innovation monoclonal antibody production mouse model novel vaccines novel



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Project Number Contact PI/Project Leader 5R21AI135373-02 WHITTAKER, GARY R

Awardee Organization **CORNELL UNIVERSITY**

gary.whittaker@cornell.edu

Organization

Country

State Code Name Department Type **CORNELL UNIVERSITY** MICROBIOLOGY/IMMUN/VIROL(NY

City **Organization Type**

SCHOOLS OF VETERINARY ITHACA

MEDICINE

Congressional District

23

Other Information

UNITED STATES (US)

FOA Administering Institutes or **Project Start** 06-June-PA-16-161 Centers 2018 Date NATIONAL INSTITUTE OF Study Section **ALLERGY AND INFECTIOUS Project End** 31-May-**Vaccines Against Microbial DISEASES** Diseases Study
Section[VMD] 2021 Date **DUNS Number CFDA Code Budget Start** 01-June-872612445 855 **Award Notice** 2019 Date Fiscal Year Date **Budget End** 31-May-2019 24-May-2019 2021 Date

Project Funding Information for 2019

Indirect Costs Total Funding Direct Costs \$235,500 \$150,000 \$85,500

Year **Funding IC**

2019 NATIONAL INSTITUTE OF ALLERGY AND INFECTIOUS DISEASES \$235,500

Click here for more information on NIH Categorical Spending NIH Categorical Spending

Funding IC	FY Total Cost by IC	NIH Spending Category
NATIONAL INSTITUTE OF ALLERGY AND INFECTIOUS DISEASES	\$235,500	Biodefense; Biotechnology; Emerging Infectious Diseases; Immunization; Infectious Diseases; Orphan Drug; Prevention; Rare Diseases; Vaccine Related;



No Sub Projects information available for 5R21Al135373-02

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No Patents information available for 5R21Al135373-02

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 5R21Al135373-02

Clinical Studies

No Clinical Studies information available for 5R21Al135373-02

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Related News Releases

No news release information available for 5R21Al135373-02

History

No Historical information available for 5R21Al135373-02

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