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Outcomes





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## **Understanding the Risk of Bat Coronavirus Emergence**

**Project Number** 2R01Al1110964-06

**Contact PI/Project Leader** DASZAK, PETER

**Awardee Organization ECOHEALTH ALLIANCE, INC.** 



## (≡) Description

#### **Abstract Text**

Project Summary: Understanding the Risk of Bat Coronavirus Emergence Novel zoonotic, bat-origin CoVs are a significant threat to global health and food security, as the cause of SARS in China in 2002, the ongoing outbreak of MERS, and of a newly emerged Swine Acute Diarrhea Syndrome in China. In a previous R01 we found that bats in southern China harbor an extraordinary diversity of SARSr-CoVs, some of which can use human ACE2 to enter cells, infect humanized mouse models causing SARS-like illness, and evade available therapies or vaccines. We found that people living close to bat habitats are the primary risk groups for spillover, that at one site diverse SARSr-CoVs exist that contain every genetic element of the SARS-CoV genome, and identified serological evidence of human exposure among people living nearby. These findings have led to 18 published peer-reviewed papers, including two papers in Nature, and a review in Cell. Yet salient questions remain on the origin, diversity, capacity to cause illness, and risk of spillover of these viruses. In this R01 renewal we will address these issues through 3 specific aims: Aim 1. Characterize the diversity and distribution of high spillover-risk SARSr-CoVs in bats in southern China. We will use phylogeographic and viral discovery curve analyses to target additional bat sample collection and molecular CoV screening to fill in gaps in our previous sampling and fully characterize natural SARSr-CoV diversity in southern China. We will sequence receptor binding domains (spike proteins) to identify viruses with the highest potential for spillover which we will include in our experimental investigations (Aim 3). Aim 2. Community, and clinic-based syndromic, surveillance to capture SARSr-CoV spillover, routes of exposure and potential public health consequences. We will conduct biological-behavioral surveillance in high-risk populations, with known bat contact, in community and clinical settings to 1) identify risk factors for serological and PCR evidence of bat SARSr-CoVs; & 2) assess possible health effects of SARSr-CoVs infection in people. We will analyze bat-CoV serology against human-wildlife contact and exposure data to quantify risk factors and health impacts of SARSr-CoV spillover. Aim 3. In vitro and in vivo characterization of SARSr-CoV spillover risk, coupled with spatial and phylogenetic analyses to identify the regions and viruses of public health concern. We will use S protein sequence data, infectious clone technology, in vitro and in vivo infection experiments and analysis of receptor binding to test the hypothesis that % divergence thresholds in S protein sequences predict spillover potential. We will combine these data with bat host distribution, viral diversity and phylogeny, human survey of risk behaviors and illness, and serology to identify SARSr-CoV spillover risk hotspots across southern China. Together these data and analyses will be critical for the future development of public health interventions and enhanced surveillance to prevent the re-emergence of SARS or the emergence of a novel SARSr-CoV.

#### **Public Health Relevance Statement**

Program Director/Principal Investigator: Daszak, Peter Renewal: Understanding the Risk of Bat Coronavirus Emergence Project Narrative Most emerging human viruses come from wildlife, and these represent a significant threat to public health and biosecurity in the US and globally, as was demonstrated by the SARS coronavirus pandemic of 2002-03. This project seeks to understand what factors allow coronaviruses, including close relatives to SARS, to evolve and jump into the human population by studying viral diversity in their animal reservoirs (bats), surveying people that live in high-risk communities in China for evidence of bat-coronavirus infection, and conducting laboratory experiments to analyze and predict which newlydiscovered viruses pose the greatest threat to human health.

## **NIH Spending Category**

**Infectious Diseases Biodefense Biotechnology Clinical Research Emerging Infectious Diseases Rare Diseases** Lung **Pneumonia** Pneumonia & Influenza Prevention

### **Project Terms**

**Acute Diarrhea Acute Address Amino Acid Sequence Animals Behavior Behavioral Biological** Cells China Chiroptera Clinic **Clinic Visits** Clinical Communities **Data Analyses Coronavirus Coronavirus Infections** Coupled **Data Development Epithelial Cells** Genome **Disease Outbreaks Exposure to** Family suidae **Future** 

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**Geographic Distribution** 

Infection Influenza Investigation Lead Middle East Respiratory Syndrome Coronavirus Maps

Habitats

**Monoclonal Antibodies Phylogenetic Analysis** Modeling Molecular **Nature Paper Patients** 

Health

Human

In Vitro

Individual

**Read More** 

# **Details**

**Other Pls Contact PI/ Project Leader Program Official** 

Not Applicable Name Name

Geography

STEMMY, ERIK J DASZAK, PETER

Contact Title **Email not available PRESIDENT** 

Contact mark.denison@vanderbilt.edu

**Organization** 

Name Department Type State Code Unavailable NY

ECOHEALTH ALLIANCE, INC.

City Organization Type **Congressional District NEW YORK Other Domestic Non-Profits** 12

Country

**Other Information** 

**UNITED STATES (US)** 

FOA Administering Institutes or Centers Project Start 01-June-2014 **NATIONAL INSTITUTE OF ALLERGY** 

PA-18-484 Date **AND INFECTIOUS DISEASES** Study Section **Project End Date** 30-June-2026 **Clinical Research and Field Studies DUNS Number** CFDA Code of Infectious Diseases Study Section[CRFS] 077090066 855

Date Fiscal Year **Award Notice Date** 

**Budget End Date** 30-June-2022 2019 24-July-2019

**Project Funding Information for 2019** 

**Total Funding Direct Costs Indirect Costs** \$661,980 \$538,926 \$123,054

**Funding IC FY Total Cost by IC** Year

2019 NATIONAL INSTITUTE OF ALLERGY AND INFECTIOUS DISEASES \$661,980

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

**FY Total Cost by IC NIH Spending Category Funding IC** 

Research; Emerging Infectious Diseases; Infectious Diseases; Lung;

\$661,980

Pneumonia; Pneumonia & Influenza; Prevention; Rare Diseases;

Biodefense; Biotechnology; Clinical

**Budget Start** 

24-July-2019

## **品 Sub Projects**

No Sub Projects information available for 2R01Al110964-06

NATIONAL INSTITUTE OF ALLERGY AND INFECTIOUS DISEASES

## **Publications**

No Publications available for 2R01Al110964-06



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No Patents information available for 2R01Al110964-06

## 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 2R01Al110964-06

## **†** Clinical Studies

No Clinical Studies information available for 2R01Al110964-06

## News and More

#### **Related News Releases**

No news release information available for 2R01Al110964-06

# **(**□) History

No Historical information available for 2R01Al110964-06

# Similar Projects

No Similar Projects information available for 2R01Al110964-06