











[Back to Search Results](#)

-  [Description](#)
-  [Details](#)
-  [Sub-Projects](#)
-  [Publications](#)
-  [Patents](#)
-  [Outcomes](#)
-  [Clinical Studies](#)
-  [News and More](#)
-  [History](#)
-  [Similar Projects](#)

Project 3: Nucleoside-modified mRNA-LNP vaccine platform

Parent Project Number	Sub-Project ID	Contact	Awardee
1P01AI158571-01A1	6382	PI/Project Leader HAYNES, BARTON F.	Organization DUKE UNIVERSITY

Description

Abstract Text

ABSTRACT - Project 3 Coronaviruses have the potential to cause significant morbidity and mortality as demonstrated by the ongoing SARS-CoV-2 pandemic. The purpose of this program project is to develop safe and broadly-protective group 2b and 2c betacoronavirus (panbetaCoV) vaccines capable of inducing protective immune responses and evaluate them in animal challenge models. The fact that there has been 3 major CoV outbreaks (SARS-CoV-1, MERS and SARS-CoV-2) in less than 20 years strongly supports the idea of generation of broadly protective panbetaCoV vaccines that can significantly contribute to global pandemic preparedness against future CoV epidemics and pandemics. Coronaviruses (CoVs) have significant pandemic potential, as illustrated by the outbreaks of SARS-CoV-1, MERS and SARS-CoV-2 in less than 20 years. The outbreak of a novel CoV, SARS- CoV-2, has resulted in at over 85 million infections and 1.8 million deaths. Thus, development of panbetaCoV vaccines is essential to preventing a future outbreaks due to an emerging new zoonotic CoV. Messenger RNA/LNP-based vaccines have proved to be highly effective against cancer and infectious diseases and one of the most effective platforms comprises nucleoside-modified **mRNA** (mod **mRNA**) encapsulated in LNPs. Two of the leading COVID-19 vaccines in phase 3 clinical trials by Moderna and Pfizer/BioNTech use our nucleoside-modified **mRNA-LNP vaccine** platform and are 95% protective in Phase 3 trials. Besides potency, **mRNA**/LNPs can undergo rapid, scalable production and induced durable immune responses. In Project 3, we propose to develop cross-protective and safe mod **mRNA**-LNP vaccines against animal and human betaCoVs and evaluate their immunogenicity and protective efficacy in preclinical studies. We hypothesize that mod **mRNA**-LNP vaccines encoding CoV immunogens capable of inducing broadly protective and broadly cross-protective B and T cell responses will effectively provide protection against future outbreaks of zoonotic CoVs.. We propose the following Specific Aims: Aim 1) Development of neutralizing antibody panbetaCoV vaccines using mod **mRNA**-LNP. Aim 2) Development of T cell vaccines using mod **mRNA**-LNP. In summary, this proposal aims to develop panbetaCoV vaccines that are safe, easy-to-produce and can induce protective immune responses in animal challenge models. The data generated will be capable of moving this panbetaCoV **vaccine** approach to clinical development.

Public Health Relevance Statement











Data not available.

Project Terms

2019-nCoV	Address	Adjuvant	Adverse event	Age	Animal Model
Animals	Antibody Response	Antigens	B-Lymphocytes	COVID-19 pandemic	
COVID-19 vaccine	Cell Surface Proteins	Cessation of life	Collaborations		
Communicable Diseases	Coronavirus	Coronavirus Infections	Country	Data	
Development	Disease	Disease Outbreaks	Ebola virus	Elderly	
Encapsulated	Epidemic	Epithelial Cells	Evaluation	Ferritin	Future
Generations	Glycoproteins	Goals	Helper-Inducer T-Lymphocyte	Human	
Human Herpesvirus 2	Immune	Immune response	Infection	Length	
Read More					

Thank you for your feedback!

[Back to Search Results](#)

-  [Description](#)
-  [Details](#)
-  [Sub-Projects](#)
-  [Publications](#)
-  [Patents](#)
-  [Outcomes](#)
-  [Clinical Studies](#)
-  [News and More](#)
-  [History](#)
-  [Similar Projects](#)

Project 3: Nucleoside-modified mRNA-LNP vaccine platform

Parent Project Number	Sub-Project ID	Contact	Awardee
1P01AI158571-01A1	6382	PI/Project Leader HAYNES, BARTON F.	Organization DUKE UNIVERSITY
Title			not available
FREDERIC M HANES PROF OF MED			
Contact			
HAYNE002@MC.DUKE.EDU			
Organization			
Name	Department Type	State Code	
DUKE UNIVERSITY	Unavailable	NC	
City	Organization Type	Congressional District	
DURHAM	Domestic Higher Education	04	
Country			
UNITED STATES (US)			

Other Information

FOA	Administering Institutes or Centers	Project Start Date	16-September-2021
PAR-20-072	NATIONAL INSTITUTE OF ALLERGY AND INFECTIOUS DISEASES	Project End Date	31-August-2024
Study Section	DUNS Number	Budget Start Date	21-September-2021
Special Emphasis Panel[ZAI1JP-W(S1)]	044387793 CFDA Code	Budget End Date	31-August-2024
Award Notice Date			
Fiscal Year			
2021			
16-September-2021			

Project Funding Information for 2021

Total Funding	Direct Costs	Indirect Costs
\$1,904,954	\$1,859,349	\$45,605
Year	Funding IC	
2021	NATIONAL INSTITUTE OF ALLERGY AND INFECTIOUS DISEASES	\$1,904,954

Sub Projects

No Sub Projects information available for 1P01AI158571-01A1 6382











Publications

No Publications available for 1P01AI158571-01A1 6382

Patents

Thank you for your feedback!

[Back to Search Results](#)

-  [Description](#)
-  [Details](#)
-  [Sub-Projects](#)
-  [Publications](#)
-  [Patents](#)
-  [Outcomes](#)
-  [Clinical Studies](#)
-  [News and More](#)
-  [History](#)
-  [Similar Projects](#)

Project 3: Nucleoside-modified mRNA-LNP vaccine platform

Parent Project Number	Sub-Project ID	Contact	Awardee
1P01AI158571-01A1	6382	PI/Project Leader HAYNES, BARTON F.	Organization DUKE UNIVERSITY
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 1P01AI158571-01A1 6382

Clinical Studies

No Clinical Studies information available for 1P01AI158571-01A1 6382

News and More

Related News Releases

No news release information available for 1P01AI158571-01A1 6382

History

No Historical information available for 1P01AI158571-01A1 6382

Similar Projects

No Similar Projects information available for 1P01AI158571-01A1 6382

Thank you for your feedback!