











[Back to Search Results](#)

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

Omics for TB: Response to Infection and Treatment

Project Number
5U19AI135976-03

Contact PI/Project Leader
ADEREM, ALAN A[Other PIs](#)

Awardee Organization
SEATTLE CHILDREN'S HOSPITAL

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Description

Abstract Text

Abstract – Overview With about 10 million new cases of active disease and 1.8 million deaths annually, TB is a global health emergency. A distinguishing feature of TB disease is its biological heterogeneity, which manifests at the clinical level chiefly in 2 forms: disease progression and treatment response. The premise of this Program is that the heterogeneous outcomes of TB infection and treatment are determined by the interplay of competing regulatory networks between the pathogen and the host. Our primary goal is to apply systems biology approaches to elucidate the biological control underlying the variability of disease outcome and response to treatment. Our first specific aim is to define novel host regulators of TB disease progression in vivo, and the innate and adaptive networks they control. We will also seek to define novel Mtb regulators of TB treatment response, and the Mtb regulatory networks that they control. This work will allow us to produce and validate host and Mtb models of TB disease progression and treatment response. Altogether, this program addresses key unanswered questions that stymie efforts to combat the TB **pandemic**. Our team has perfected the required platforms and scientific approaches to execute this ambitious research plan in a timely and cost- effective manner. All the participating investigators have strong records of interacting productively, and of disseminating their data and reagents to the scientific community.

Public Health Relevance Statement


Project Narrative - Omics for TB: Response to Infection and Treatment Mycobacterium tuberculosis causes ~10 million new cases of active disease and 1.8 million deaths each year, and our tools to combat tuberculosis (TB) disease are universally outdated and overmatched. This project combines separate advances in systems biology and network modeling to produce experimentally grounded and verifiable systems-level models of the host and MTB regulatory networks that affect disease progression and response to treatment.

Project Terms

Address	Affect	Bacteria	Biological	Cessation of life	Clinical	Collection	
Communicable Diseases		Communities	Complex	Data	Data Set	Disease	Disease Outcome
Disease Progression	Drug-sensitive	Eicosanoids	Elements	Genetic	Transcription	Goals	
Human	Immunologic Receptors	Infection	Inflammatory Response	Infrastructure			
Machine Learning	Mass Spectrum Analysis	Methodology	Modeling	Molecular Profiling			
Mouse Strains	Multiplexed Ion Beam Imaging	Mus	Mycobacterium tuberculosis	Network-based			
Outcome	Pharmaceutical Preparations	Phenotype	Predisposition	Proteomics	Reagent		
Receptor Activation	Records	Regulator Genes	Research	Research Personnel	Resources		
Read More							

Details

Contact PI/ Project Leader

Name
[ADEREM, ALAN A](#) 

Title
PROFESSOR

Contact
Alan.Aderem@seattlechildrens.org

Other PIs

Name
[SHERMAN, DAVID R](#) 

Program Official

Name
SHABMAN, REED SOLOMON

Contact
reed.shabman@nih.gov

Organization

Name
SEATTLE CHILDREN'S HOSPITAL

City
SEATTLE


Country
UNITED STATES (US)

Department Type
Unavailable

Organization Type
Independent Hospitals

State Code
WA

Congressional District
07

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[Back to Search Results](#)

 [Description](#)

 [Details](#)

 [Sub-Projects](#)

 [Publications](#)

 [Patents](#)

 [Outcomes](#)

 [Clinical Studies](#)

 [News and More](#)

 [History](#)

 [Similar Projects](#)

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Awardee Organization
SEATTLE CHILDREN'S HOSPITAL

Fiscal Year 2019	Award Notice Date 21-February-2019	040002137	033	2020
		Budget Start Date	01-February-2019	
		Budget End Date	31-January-2020	

Project Funding Information for 2019

Total Funding \$3,359,502	Direct Costs \$2,323,147	Indirect Costs \$1,370,783
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Year	Funding IC	FY Total Cost by IC
2019	NATIONAL INSTITUTE OF ALLERGY AND INFECTIOUS DISEASES	\$3,359,502

 Sub Projects

 Export

Project Number	Sub	Principal Investigator(s)/ Project Leader(s)	Organization	Fiscal Year	Admin IC	FY Total Cost by IC
Project 1: Mechanisms of Disease Progression						
5U19AI135976-03	6321	 ADEREM, ALAN A 	SEATTLE CHILDREN'S HOSPITAL	2019	NIAID	\$1,195,053
Adminstrative Core						
5U19AI135976-03	6317	 ADEREM, ALAN A 	SEATTLE CHILDREN'S HOSPITAL	2019	NIAID	\$258,881
Technology Core						
5U19AI135976-03	6319	 AITCHISON, JOHN D. 	SEATTLE CHILDREN'S HOSPITAL	2019	NIAID	\$1,214,310
Modeling Core						
5U19AI135976-03	6320	 BALIGA, NITIN S. 	SEATTLE CHILDREN'S HOSPITAL	2019	NIAID	\$429,688
Project 2: Response to Treatment						
5U19AI135976-03	6322	 SHERMAN, DAVID R. 	SEATTLE CHILDREN'S HOSPITAL	2019	NIAID	\$452,538

 Publications

No Publications available for 5U19AI135976-03

 Patents

No Patents information available for 5U19AI135976-03


 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 5U19AI135976-03

 Clinical Studies

No Clinical Studies information available for 5U19AI135976-03

 Thank you for your feedback!

[Back to Search Results](#)

 [Description](#)

 [Details](#)

 [Sub-Projects](#)

 [Publications](#)

 [Patents](#)

 [Outcomes](#)

 [Clinical Studies](#)

 [News and More](#)

 [History](#)

 [Similar Projects](#)

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Awardee Organization
SEATTLE CHILDREN'S HOSPITAL

History

No Historical information available for 5U19AI135976-03

Similar Projects

No Similar Projects information available for 5U19AI135976-03