

Truth in Testimony Disclosure Form

In accordance with Rule XI, clause 2(g)(5)* of the *Rules of the House of Representatives*, witnesses are asked to disclose the following information. Please complete this form electronically by filling in the provided blanks.

Committee: Oversight and Accountability

Subcommittee: Select Subcommittee on the Coronavirus Pandemic

Hearing Date: 07/11/23

Hearing Title :

“Investigating the Proximal Origin of a Cover Up”

Witness Name: Robert Francis Garry

Position/Title: Professor/Associate Dean, Tulane University School of Medicine

Witness Type: Governmental Non-governmental

Are you representing yourself or an organization? Self Organization

If you are representing an organization, please list what entity or entities you are representing:

FOR WITNESSES APPEARING IN A NON-GOVERNMENTAL CAPACITY

Please complete the following fields. If necessary, attach additional sheet(s) to provide more information.

Are you a fiduciary—including, but not limited to, a director, officer, advisor, or resident agent—of any organization or entity that has an interest in the subject matter of the hearing? If so, please list the name of the organization(s) or entities.

Co-founder, Zalgen Labs, LLC

Please list any federal grants or contracts (including subgrants or subcontracts) related to the hearing's subject matter that you or the organization(s) you represent have received in the past thirty-six months from the date of the hearing. Include the source and amount of each grant or contract.

Dr. Garry does not personally receive any federal grant funding, and he is appearing before the Subcommittee in his personal capacity. Nonetheless, to provide as much information to the Subcommittee as possible, please see attached, as Exhibit C, a list of coronavirus-related research that is supported by federal funding. Given the topic of the hearing, Dr. Garry has not included other grants related to research on non-coronaviruses.

Please list any contracts, grants, or payments originating with a foreign government and related to the hearing's subject that you or the organization(s) you represent have received in the past thirty-six months from the date of the hearing. Include the amount and country of origin of each contract or payment.

Dr. Garry is appearing before the Subcommittee in his personal capacity, and he does not personally receive any foreign funding. Further, Dr. Garry receives no foreign funding for coronavirus-related research. Given the topic of the hearing, he has not included other grants related to research on non-coronaviruses.

Please complete the following fields. If necessary, attach additional sheet(s) to provide more information.

- I have attached a written statement of proposed testimony.
- I have attached my curriculum vitae or biography.

*Rule XI, clause 2(g)(5), of the U.S. House of Representatives provides:

(5)(A) Each committee shall, to the greatest extent practicable, require witnesses who appear before it to submit in advance written statements of proposed testimony and to limit their initial presentations to the committee to brief summaries thereof.

(B) In the case of a witness appearing in a non-governmental capacity, a written statement of proposed testimony shall include— (i) a curriculum vitae; (ii) a disclosure of any Federal grants or contracts, or contracts, grants, or payments originating with a foreign government, received during the past 36 months by the witness or by an entity represented by the witness and related to the subject matter of the hearing; and (iii) a disclosure of whether the witness is a fiduciary (including, but not limited to, a director, officer, advisor, or resident agent) of any organization or entity that has an interest in the subject matter of the hearing.

(C) The disclosure referred to in subdivision (B)(ii) shall include— (i) the amount and source of each Federal grant (or subgrant thereof) or contract (or subcontract thereof) related to the subject matter of the hearing; and (ii) the amount and country of origin of any payment or contract related to the subject matter of the hearing originating with a foreign government.

(D) Such statements, with appropriate redactions to protect the privacy or security of the witness, shall be made publicly available in electronic form 24 hours before the witness appears to the extent practicable, but not later than one day after the witness appears.

False Statements Certification

Knowingly providing material false information to this committee/subcommittee, or knowingly concealing material information from this committee/subcommittee, is a crime (18 U.S.C. § 1001). This form will be made part of the hearing record.

Robert F. Garry

Witness signature

7/6/23

Date

Truth in Testimony Disclosure Form
Dr. Robert F. Garry, PhD

Attachment A

Written Statement of Dr. Robert F. Garry
Professor and Associate Dean, Tulane University School of Medicine

United States House of Representatives
Oversight and Accountability Committee
Select Subcommittee on the Coronavirus Pandemic
“Investigating the Proximal Origin of a Cover Up.”
Tuesday, July 11, 2023
2154 Rayburn House Office Building
Washington, DC 20515

**Written Statement of Dr. Robert F. Garry
Professor & Associate Dean, Tulane University School of Medicine**

Chairman Wenstrup, Ranking Member Ruiz, distinguished members of the Subcommittee, thank you for inviting me to testify today. For the last forty years, I have worked as a professor at Tulane University School of Medicine. I’ve devoted my life’s work to understanding emerging viruses, such as HIV, Ebola virus, Lassa virus and the first SARS virus, and helping to develop diagnostics, drugs and vaccines to help stop those viruses. At the outset, it is important to note that I make these statements in my personal capacity, and I am not speaking on behalf of Tulane University.

Although we have all lived through a very challenging viral pandemic, my personal perspective has been different than most. For nearly 20 years, I’ve worked closely with scientists and clinicians at the Kenema Government Hospital (KGH) in Sierra Leone. KGH is a major site for research on the virus that causes Lassa fever. Ten years ago, Ebola virus emerged just 50 miles from KGH.¹ Ultimately, the Ebola outbreak that occurred in Sierra Leone between 2013 and 2016 claimed 12,000 lives, including the lives of dozens of healthcare workers at KGH. Having previously lost many close colleagues to an outbreak of a deadly virus, the December 2019 reports of cases of a novel pneumonia in Wuhan, China were ominous. The reports raised the specter of a

¹ Goba A *et al.* 2016. An Outbreak of Ebola Virus Disease in the Lassa Fever Zone. *J Infect Dis* 15: S110-S121; *see also*

possible impending global disaster caused by a novel airborne virus – one I worried that the world would be ill-equipped to handle.

Shortly after the first release of the SARS-CoV-2 genetic sequence,² I participated in an in-depth molecular and phylogenetic analysis of the virus with a group of three other scientists, Dr. Kristian Andersen, Dr. Eddie Holmes, and Dr. Andrew Rambaut. The four of us, as well as a fifth co-author, Dr. Ian Lipkin, wrote a peer-reviewed publication, titled “The Proximal Origin of SARS-CoV-2.” In the paper, we concluded that it was likely that SARS-CoV-2 had evolved naturally.³

Importantly, however, we specifically did not rule out a laboratory origin. Instead, in our paper, we discussed three possible origin scenarios. The first scenario was direct spillover from a bat to a human, and the second scenario was spillover from a bat to an intermediate animal and then to a human. The third scenario we discussed in the paper was a lab origin. Specifically, we discussed the possibility that some of the SARS-CoV-2 Spike protein’s features, including a receptor binding domain (RBD) that effectively binds human angiotensin-converting enzyme 2 (ACE2) and a furin cleavage site (FCS), may have arisen during passage in a laboratory.⁴ However, because we observed these notable features in related coronaviruses, which provided a straightforward evolutionary route for SARS-CoV-2 to emerge in nature, we concluded that the natural origin scenarios were most plausible and that, based on the then-available scientific evidence, we did not believe that laboratory-based scenarios, including bioengineering, were plausible.

² Holmes E. 2020. Novel 2019 coronavirus genome, available at <https://virological.org/t/novel-2019-coronavirus-genome/319>.

³ Andersen KG *et al.* 2020. The proximal origin of SARS-CoV-2. *Nat Med* 26:450-452.

⁴ Passage is the process of growing a virus in iterations in different environments (either in cultures or animals), either to observe or achieve mutations and changes.

Based on evidence that has accumulated since we wrote *Proximal Origin*, it is my opinion that SARS-CoV-2 emerged via the wildlife trade in a market in Wuhan, China.⁵ Peer-reviewed papers provide robust evidence supporting that the virus spilled over from a bat through an intermediate animal or animals to humans, one of three origin scenarios discussed in *Proximal Origin*. First, the Huanan Market in Wuhan, China was the early epicenter of the COVID-19 outbreak.⁶ Most of the earliest diagnosed human COVID-19 cases from December 2019 lived in the immediate neighborhood around this market, including those that did not work or shop there. The ascertainment of cases in late 2019 was by local health officials, not the central Chinese Centers for Disease Control, and was not biased by linkage to the Huanan Market [6].⁷ The two campuses of the Wuhan Institute of Virology (WIV) are considered the prime suspects for a lab origin of SARS-CoV-2 because they are the sites of the most advanced research programs on coronaviruses in the city.⁸ However, there was no clustering of diagnosed cases of COVID-19 in December 2019 around either the Wuchang or Jiangxi campuses of the WIV, which are 7 and 15 miles respectively, from the Huanan Market, as would be expected if entry of SARS-CoV-2 into humans involved a laboratory accident.

Second, despite official denials from China, it was determined that Huanan Market vendors sold illegal SARS-CoV-2 susceptible wildlife in November 2019, the most likely timeframe in

⁵ Garry RF. 2022. The evidence remains clear: SARS-CoV-2 emerged via the wildlife trade. *Proc Natl Acad Sci USA*. 119:e2214427119.

⁶ Holmes EC *et al.* 2021. The origins of SARS-CoV-2: A critical review. *Cell* 184:4848-4856; Worobey M. 2021. Dissecting the early COVID-19 cases in Wuhan. *Science* 374:1202-1204; Worobey M *et al.* 2022. The Huanan Seafood Wholesale Market in Wuhan was the early epicenter of the COVID-19 pandemic. *Science* 377:951-959.

⁷ Worobey M. 2021. Dissecting the early COVID-19 cases in Wuhan. *Science* 374:1202-1204.

⁸ Chan A and Ridley M. *Viral: The Search for the Origin of COVID-19* (News Corp: Harper-Collins, New York, 2021); Harrison NL and Sachs JD (2022), A call for an independent inquiry into the origin of the SARS-CoV-2 virus. *Proc Natl Acad Sci USA* 119(21):e2202769119.

which SARS-CoV-2 emerged.⁹ Environmental sampling was performed at the Huanan Market after it was shut down on January 1, 2020 and after illegal wildlife had been removed.¹⁰ SARS-CoV-2 positive samples clustered in the southwest corner of the Huanan Market, where live SARS-CoV-2 susceptible mammals were sold. Specifically, an iron cage, carts used to move cages and animals, and drainage from this area were positive for SARS-CoV-2.¹¹ It is difficult to reconcile this striking distribution of the SARS-CoV-2 positive environmental samples in the Huanan Market with the theory that SARS-CoV-2 originated at the WIV or another Wuhan laboratory.

Third, analyses of the environmental samples also indicated that at least two separate spillovers of SARS-CoV-2 from animals to humans occurred at the Huanan Market.¹² In this regard, the emergence of SARS-CoV-2 resembles the first outbreaks of a SARS coronavirus in China that happened between 2002 and 2004 in which there were multiple spillovers from live animals sold in the wildlife trade. There are no plausible lab origin scenarios that are compatible with two independent spillovers of SARS-CoV-2 at the same market location.

Finally, in November 2022, my co-authors and I obtained access to large files containing the DNA and RNA sequences from the environmental samples taken at the Huanan Market in early 2020. This long-suppressed data not only showed that live raccoon dogs, civet cats, and other mammals susceptible to SARS-CoV-2 infection were present, but also pin-pointed their precise

⁹ Xiao X *et al.* 2021. Animal sales from Wuhan wet markets immediately prior to the COVID-19 pandemic. *Sci. Rep.* 11:11898 (2021); Huang C *et al.* 2020. Clinical features of patients infected with 2019 novel coronavirus in Wuhan, China. *Lancet* 395: 497–506.

¹⁰ Liu WJ *et al.* 2023. Surveillance of SARS-CoV-2 at the Huanan Seafood Market. *Nature*. 2023 Apr 5. doi: 10.1038/s41586-023-06043-2.

¹¹ Liu WJ *et al.* 2023. Surveillance of SARS-CoV-2 at the Huanan Seafood Market. *Nature*. 2023 Apr 5. doi: 10.1038/s41586-023-06043-2; Crits-Christoph *et al.* 2023. Genetic evidence of susceptible wildlife in SARS-CoV-2 positive samples at the Huanan Wholesale Seafood Market, Wuhan: Analysis and interpretation of data released by the Chinese Center for Disease Control <https://doi.org/10.5281/zenodo.7754299>.

¹² Pekar JE *et al.* 2022. The molecular epidemiology of multiple zoonotic origins of SARS-CoV-2. *Science*. 377:960-966.

locations within the market. Raccoon dog and civet cat DNA and RNA were present in the wildlife stall that contained the highest numbers of SARS-CoV-2 positive samples in the market.¹³ This is equivalent to finding a smoking gun carrying the main suspect's DNA at the exact scene of the crime. All of this evidence demonstrates that the simplest explanation is that live animals were shedding SARS-CoV-2 at the Huanan Market in late 2019.

Theories of COVID-19 origin must be investigated in a transparent manner. In prior testimony, this Subcommittee has heard scientific testimony regarding SARS-CoV-2 that is not fully accurately. For example, the Subcommittee has been told that there is no FCS in any other SARS-like virus.¹⁴ However, an FCS exists in the first SARS-CoV near putative fusion peptides.¹⁵ Moreover, other SARS-like viruses are but a single mutation from having an FCS, which could evolve in a single step.¹⁶ The Subcommittee was also told that cleavage at the FCS reorients the RBD so it can specifically bind to human ACE2.¹⁷ This is inaccurate. The same witness described “human” arginines, which do not exist.

Three and a half years into the COVID-19 pandemic, it is still my opinion that there is no credible scientific evidence to support a lab-based origin for SARS-CoV-2. I support the efforts of the Subcommittee to better understand the origins of coronavirus pandemics, as understanding viral origin plays an important role in developing strong policies to help prevent the next potential

¹³ Crits-Christoph *et al.* 2023. Genetic evidence of susceptible wildlife in SARS-CoV-2 positive samples at the Huanan Wholesale Seafood Market, Wuhan: Analysis and interpretation of data released by the Chinese Center for Disease Control <https://doi.org/10.5281/zenodo.7754299>.

¹⁴ Testimony of Nicholas Wade on 8 March 2023 to the House Select Subcommittee on Coronavirus Pandemic; Testimony of Jamie Metzl, PhD on 8 March 2023 to the House Select Subcommittee on Coronavirus Pandemic.

¹⁵ Sainz B Jr *et al.* 2005. Identification and characterization of the putative fusion peptide of the severe acute respiratory syndrome-associated coronavirus spike protein. *J Virol* 79: 7195-206.

¹⁶ Sander, AL., Moreira-Soto, A., Yordanov, S. *et al.* 2022. Genomic determinants of furin cleavage in diverse European SARS-related bat coronaviruses. *Commun Biol* 5: 491.

¹⁷ Testimony of Robert Redfield, MD on 8 March 2023 to the House Select Subcommittee on Coronavirus Pandemic.

pandemic. The global community remains ill-equipped to prevent or manage the emergence of novel viruses.¹⁸ Prevention efforts for the next coronavirus emergence should focus on obtaining a greater understanding of the diversity of the *Coronaviridae* family in wild animals, increased surveillance at the animal-human interface, and stringent oversight of the wildlife and fur trade. As mentioned earlier, I've spent much of my career developing countermeasures for emerging viruses. Diagnostics, vaccines and therapeutics for potentially pandemic coronaviruses and other high-risk viruses, can and must be developed and prepositioned.¹⁹

¹⁸ Goodrum F, et al. 2023. Virology under the microscope—a call for rational discourse. *J Virol* 97:e00089-23.

¹⁹ Graham B and Corbett KS. 2020. Prototype pathogen approach for pandemic preparedness: world on fire. *J Clin Invest* 130: 3348-3349.

Truth in Testimony Disclosure Form
Dr. Robert F. Garry, PhD

Attachment B

Curriculum Vitae of Dr. Robert F. Garry

CURRICULUM VITAE

NAME: **Robert Francis Garry, Jr.** tel: 504-988-2027
fax: 504-988-1994
e-mail: rfgarry@tulane.edu

Date and place of birth: August 7, 1951. Terre Haute, Indiana.

Marital status: Married to Jeanette P. Alexander-Garry (three children).

Education:

1978 Ph.D. in Microbiology. The University of Texas at Austin, Texas.
(Marilynn R. Fairfax Waite, advisor).

1974 B.S. in Life Science. Indiana State University at Terre Haute,
Indiana.

Academic positions held:

2012-date Associate Member. Broad Institute of Harvard and MIT

2012-date Adjunct Professor. Tuskegee University

2006-date Assistant Dean for Graduate Studies in Biomedical Sciences, Tulane
University

2004-2006 Director, Interdisciplinary Program in Molecular and Cellular
Biology

1993-date Professor of Microbiology and Immunology. Tulane University School
of Medicine at New Orleans, Louisiana. 70112.

1987-1993 Associate Professor of Microbiology and Immunology
(appointment with tenure 1989) Tulane University School of Medicine
at New Orleans, Louisiana. 70112.

1991 Fogarty Visiting Professor of Molecular Biology (Gebhard Koch,
sponsor). University of Hamburg at Hamburg, Germany.

1983-1987 Assistant Professor of Microbiology and Immunology. Tulane
University School of Medicine at New Orleans, Louisiana. 70112.

1985 Visiting Professor of Pathology (Suraiya Rasheed, sponsor).
University of Southern California at Los Angeles, California.

1982-1983 Instructor of Microbiology. The University of Texas at Austin,
Texas. 78712.

1979-1983 Post-doctoral (Henry R. Bose, Jr., sponsor). The University of
Texas at Austin, Texas. 78712.

Other Appointments:

2016 Center for Viral Systems Biology. Co-Director.

2016 Tulane Center of Excellence under the Global Viral Network. Director.

2013 African Centre of Excellence for Genomics of Infectious Disease [ACEGID], which is part of the NIH/Wellcome Trust H3Africa Consortium. Founding member.

2012 Viral Immunotherapeutics Consortium. Member.

2010 Co-founder Zalgen Labs. Germantown, Maryland and Aurora, Colorado.

2008 Viral Hemorrhagic Fever Consortium [VHFC, vhfc.org]. Program Manager.

Research interests:

Investigations at the molecular level of cytopathic, immunodisruptive and carcinogenic mechanisms of human viruses. Development of countermeasures: diagnostics, therapeutics and vaccines.

Key words:

viruses, Lassa fever, Ebola, SARS/COVID-19, dengue, influenza, HIV, AIDS, hepatitis C virus, cytopathology, molecular biology, genetics, immunology.

Research Support (Selected):

National Institute of Allergy and Infectious Diseases. "Consortium for Viral Systems Biology (CViSB)" (5U19AI135995). 02/01/2018 - 01/31/2028. Principal Investigator: Andersen, K. Director of subcontract.

European & Developing Countries Clinical Trials Partnership (EDCTP) "Lassa Fever Vaccine Efficacy and Prevention for West Africa" (RIA2019LV-3053) 01/01/22 - 12/31/26. Principal Investigator: Swati Gupta (International AIDS Vaccine Initiative). Director of subcontract.

National Institute of Allergy and Infectious Diseases. "West African Emerging Infectious Disease Research Center" (U01AI151812) 05/21/2020 - 04/30/2025. Multiple Principal Investigators: Garry, Kristian Andersen (Scripps Research) and Pardis Sabeti (Harvard).

Wellcome Trust Foundation (U.K.) "Incidence studies in support of Lassa fever vaccine development." (215858/Z/19/Z) 07/01/2020-02/26/2023 Principal Investigator: Gupta, S. Director of subcontract.

National Institute of Allergy and Infectious Diseases. "Consortium for Immunotherapeutics against Emerging Viral Threats" (5U19AI142790). 05/01/2019-01/30/2024. Principal Investigator: Sapphire, E. Investigator.

National Institute of Allergy and Infectious Diseases. "Preclinical evaluation of a potent Lassa Fever immunotherapeutic antibody cocktail" (1R01AI132223-. 06/26/17 - 05/31/23. Principal Investigator.

National Institute of Allergy and Infectious Diseases. "Structure-based design of novel Lassa virus glycoproteins for vaccine development" (1 R01AI132244-. 07/14/17 - 06/30/23. Principal Investigator.

National Eye Institute "Pathogenesis of Uveitis in EVD Survivors" (5R01EY029594-02A1) 07/01/2020-06/30/2024. Principal Investigator: Yeh, S. Director of subcontract.

BARDA "Recombinant vesicular stomatitis virus pseudotyped Sudan glycoprotein (rVSV pseudotyped SUDV GP) vaccine/" (BAA-18-100-SOL-00003). 01/01/2022 - 12/31/2023 Principal Investigator: Price, M. Director of subcontract.

Coalition for Epidemic Preparedness Innovations. "Lassa virus multi-country serum supply and multi-lineage recombinant antigens" (INTU1901). 1/1/19 - 12/30/22. Principal Investigator.

Coalition for Epidemic Preparedness Innovations "Lassa neutrality-Production of HIV-1 Env pseudotyped virus batches-Task Order 2" (CCIDA09212-CEPI) 07/01/2021-09/30/2022 Principal Investigator: Gupta, S. Director of subcontract.

Coalition for Epidemic Preparedness Innovations."Cross-Sectional Multi-Site Seroprevalence Study to Estimate Lassa Virus Infection in West African Countries" (ESEP1904) 07/1/2019 - 6/30/2022. Multiple Principal Investigators: Garry, Grant (Sierra Leone), Happi (Nigeria).

National Heart Lung and Blood Institute. NIH. LA-CEAL: Louisiana Community-Engagement Research Alliance Against COVID-19 in Disproportionately Affected Communities. (HL158260) 09/09/21 - 09/08/21. Principal Investigators: Krouselwood (Tulane) and Sarpong (Xavier) Investigator.

Burroughs Wellcome Foundation. "Postdoc Program for Underrepresented Minorities" 09/01/20 - 08/31/23. Principal Investigator.

Centers for Disease Prevention and Control. "Immune Response to SARS CoV-2 in Special Populations" (co75D30120C08472) 07/01/20 - 06/30/21. Principal Investigator: Fusco (Tulane). Investigator.

National Institute of Allergy and Infectious Diseases. "Consortium for Immunotherapeutics against emerging viral diseases." (U19AI109762). 03/15/2014 - 02-28-19. Principal Investigator: Erica Saphire (TRSI/LaJolla Institute for Immunology). Director of Project 3.

National Institute of Allergy and Infectious Diseases. "Identification and validation of novel human T cell epitopes in Lassa fever." (HHSN272201400048C. 03/15/2014 - 02-28-19. Principal Investigator: Michael B. A. Oldstone (TRSI). Program Manager and Director of subcontract.

National Institute of Allergy and Infectious Diseases. "International Collaboration in Infectious Disease Research on Lassa fever and Ebola" (U19 AI115589. 01/01/15 - 12/31/20. Principal Investigator.

National Institute of Human Genome Sciences. "Genomic Characterization and Surveillance of Microbial Threats in West Africa " (U54 HG007480). 06/01/17 - 05/31/22 Principal Investigator: Erica Saphire (TRSI/LaJolla Institute for Immunology). Role: Director of subcontract.

Coalition for Epidemic Preparedness and Innovations. "A Vaccine for Prevention of Lassa Fever based on a Live Vesicular Stomatitis Virus-Lassa Virus Chimera." Principal Investigator: Swati Gupta (International AIDS Vaccine Initiative). Director of subcontract.

National Institute of Allergy and Infectious Diseases. "Preclinical development of recombinant antigen diagnostics for Lassa fever" (AI082778). 04/14/09 - 05/13/14. Principal Investigator. Direct costs: \$7,073,538.

National Institute of Allergy and Infectious Diseases. Peptide inhibitors of influenza entry" (AI082778). 04/01/09 - 3/30/12. (Wilson, PI). Director of Subcontract). Direct costs: \$1,304,709 (Tulane subaward: \$672,091 - first year only, years 2-3 TBD).

National Institute of Allergy and Infectious Diseases. "Roles of protective or pathogenic B cell epitopes in human Lassa fever." (Robinson, PI). 09/15/09 - 09/14/14. Program Manager. Direct costs: \$15,280,399.

Louisiana Board of Regents. "Design, Delivery and Development of Therapeutic Peptides" (RC-0013-07) 9/1/07 - 7-31-10. Principal Investigator. Direct costs: \$5,800,000.

National Institute of Allergy and Infectious Diseases. Host Genetic Factors in Resistance to Lassa Hemorrhagic Fever 10/01/10 - 09/30/15. Co-Principal Investigator. Direct costs: \$7,698,856

National Institute of Allergy and Infectious Diseases. Recombinant antigen diagnostics for filoviruses-FAST TRACK. 04/01/2009 - 03/31/12 Investigator. Direct costs: \$1,995,760

National Institute of Allergy and Infectious Diseases. Recombinant antigen multiagent diagnostic assays for Lassa and other arenaviruses (AI067188). 7/1/05 - 6/30/08. Principal Investigator. \$3,860,593

National Institute of Diabetes and Digestive and Kidney Diseases "HIV-1 Tat modulation of HCV replication and pathogenesis (DK070551)" 4/1/05-3/31/08. Principal Investigator. Direct costs: \$275,000

National Institute of Allergy and Infectious Diseases. Development of molecular diagnostic assays for systemic autoimmune diseases (AI068221)" 12/1/05 - 11/30/06. (Kolakovsky- PI) Investigator. Direct costs: \$100,000.

National Institute of Allergy and Infectious Diseases. "Peptide drugs against influenza virus (AI068230)". 7/1/06 - 6/30/08. Principal Investigator. Direct costs: \$700,000.

National Institute of Allergy and Infectious Diseases. Peptide inhibitors of dengue virus infectivity (AI64617)" 7/1/06 - 6/30/07. Principal Investigator. Direct costs: \$300,000.

National Institute of Allergy and Infectious Diseases. "Rapid screen for Ebola virus membrane interactions/drugs." (AI54626) 4/1/03 - 3/31/06. Direct costs: \$525,000 Principal Investigator.

National Institute of Allergy and Infectious Diseases. "Rapid screen for HIV TM membrane interactions/drugs." (AI54238) 4/1/03 - 3/31/05 Direct costs: \$300,000 Principal Investigator.

Wall Fund "Upgrade to Biosafety Level Three facilities" 7/1/03-6/30/06 Direct costs: \$500,000.

National Center for Research resources. "Primate pilot study: coincident AIDS/relapsing malaria." (AI 34764) 7/1/03 - 6/30/05 Direct costs: \$816,720. Principal Investigator.

National Cancer Institute. CA08921 (Srikanta Dash, PI) Garry: Investigator Hepatitis C virus and hepatocellular carcinoma Direct costs: \$900,000 7/1/01 - 6/30/06. Investigator.

Roche Molecular Systems/Autoimmune Technologies. "Blinded trial to define the role of a HMTV in breast cancer" Direct costs: \$25,000, 7/1/02 - 8/31/02 Principal Investigator.

Department of Defense. "Role of a human endogenous retrovirus in breast cancer." (BC990847) 9/1/00 - 10/31/03. Direct costs: \$225,000. Principal Investigator.

National Institute of Allergy and Infectious Diseases. "Alterations of ion transport by HIV". (AI 34764) 12/1/93 - 11/30/02 Direct costs: \$1,785,110 Principal Investigator.

National Institute of Dental Research. "Sjogren's Xerostomia: Viral/Immunological Etiology NIH-DE10862-01 Direct costs: \$1,250,873 (\$291,007 for Garry Laboratory - subcontract from LSU to Tulane) 3/1/93 - 2/29/00 Co-principal Investigator (William R. Gallaher, Principal Investigator).

National Cancer Institute. "Molecular characterization of HIV-1 and HHV-8 from a 1968 AIDS/KS case" 11/30/96 - 12/1/99 Direct costs: \$278,491. Principal Investigator.

Teaching:

- | | |
|-----------|--|
| 2018-23 | Short course in Medical Microbiology. Johns Hopkins School of Public Health and Tropical Medicine. Lecturer. |
| 2013-date | ACEGID summer Program. Broad Institute of Harvard and MIT. |
| 2010-2019 | Tulane University Ethics course (lecture on peer review) |
| 2006-date | BMS Seminar series (course director) |
| 2005-date | Instructor/Co-coordinator of Advanced Virology course. (team taught Laura Levy, Cindy Morris, Lilia Melnik, etc. Tulane Medical School, New Orleans. |
| 2003 | Instructor in Structural Biology course (Course Director: Sam Landry). Tulane University, New Orleans |
| 1998 | Instructor in Environmental Ethics course (Course Director: William Tuscano). Tulane University, New Orleans |

1991-2001 Instructor in Molecular and Cellular Pathogenesis course. (Course director, Gil Morris). Tulane University, New Orleans.

1983-date Instructor in Medical Microbiology and Immunology course (Virology lectures and general laboratory; Course director 1998-99. Tulane Medical School, New Orleans).

1983-2005 Instructor/Co-coordinator of Advanced Virology course. (team taught with Laura Levy). Tulane Medical School, New Orleans.

1983-1990 Instructor in Cancer Biology course. (Course director, M. Mizell). Tulane University, New Orleans.

1982-1983 Instructor/Coordinator of Introductory Virology courses. University of Texas, Austin.

1974 Student Teaching Internship in Biological Science. Broadripple High School, Indianapolis, Indiana.

Invited Lectureships and Symposia:

2021 Global Viral Network - Forefront of Virology Webinar Series. Invited Speaker. (On-line meeting).

2021 Fifteenth International Conference on Molecular Epidemiology and Evolutionary Genetics of Infectious Diseases. Keynote Speaker (On-line meeting).

2020 Viral Immunotherapeutics Consortium Annual Meeting, La Jolla California. Invited Speaker.

2019 Viral Hemorrhagic Fever Consortium Annual Meeting, Boulder Colorado. Organizer.

2019 Global Virus Network. Barcelona Spain. Invited Speaker.

2019 CEPI Biological Standards and Assays workshop. Oslo Norway. Invited Speaker.

2019 H3Africa Annual Meeting, Dakar Senegal. Invited Speaker.

2019 National Institute of Allergy and Infectious Diseases. Systems Biology for Infectious Diseases Annual Meeting, Galveston Texas. Invited Speaker.

2018 Global Viral Network. Annecy, France.

2018 Institute of Human Virology. Baltimore Maryland. Invited Speaker.

2018 National Institute of Allergy and Infectious Diseases. Bethesda, MD. Invited Speaker.

2018 Antibodies as Drugs (Keystone meeting). Whistler, BC. Invited speaker.

2017 United States Embassy, Freetown Sierra Leone. Invited speaker.

2017 Global Viral Network meeting. Melbourne, Invited Speaker.

2017 Coalition for Epidemic Preparedness and Innovations. Kickoff meeting. Invited participant. Paris.

2017 PEGS Boston. Symposium Speaker/Session Chair.

2017 ASM Biothreats. Keynote address (replacing Thomas M. Countryman, Under Secretary for Arms Control and International Security) and Session Chair.

2012 Annual Meeting Division of Microbiology and Infectious Diseases, National Institute of Allergy and Infectious Diseases, NIH. Keynote address.

2012 Improving Malaria Vaccine Strategies through the Application of Immunological Principles. Division of Allergy, Infection and Transplantation, National Institute of Allergy and Infectious Diseases, NIH. Invited Workshop participant

2011 World Health Organization meeting on Lassa Fever. Freetown, Sierra Leone.

2011 HBIOMED-SL Symposium, Freetown, Sierra Leone. Invited Speaker.

2010 World Health Organization. Geneva, Switzerland, Invited Speaker.

2010 HBIOMED-SL Symposium, Freetown, Sierra Leone. Keynote Address.

2009 Global Viral Forecasting Initiative. San Francisco. Invited Speaker.

2009 Irrua Specialist Teaching Hospital, Nigeria. Invited Symposium Speaker.

2009 Division of Microbiology and Infectious Diseases (DMID), National Institutes of Health. Rockville, MD. Invited Speaker.

2008 XIV International Congress of Virology. Istanbul, Turkey. Invited Speaker.

2008 International Symposium on Filoviruses. Libreville, Gabon. Participant.

2008 Synexis Symposium on Hepatitis C Virus. Research Triangle Park, North Carolina. Invited Speaker.

2007 Biomedical Research Symposium. Tuskegee University. Keynote Speaker.

2006 International Research in Infectious Diseases Annual Meeting. Washington, DC. Speaker.

2005 Novartis. Boston, MA. Invited Speaker.

2004 11th International Symposium on Hepatitis C Virus and Related viruses. Heidelberg, Germany. Invited Speaker.

2004 XVI International AIDS Congress. Bangkok, Thailand. Participant.

2004 International Seminar on Viruses and Neoplasias. Keynote Speaker. Santiago Caballeros, Dominican Republic.

2003 International workshop on hepatitis C virus. Kyoto, Japan. Speaker.

2003 Workshop on receptors and entry of oncogenic viruses. Park City, Utah. Invited speaker.

2003 American Society for Virology. Speaker

2002 NIH symposium of the role of viruses in human breast cancer. Bethesda, Maryland. Invited speaker.

2002 Roche Molecular Systems, Alameda, California. Invited speaker.

2001 International Congress of the Hellenic Society for Breast Cancer Research, Athens, Greece. Invited Keynote Speaker.

2000 Viruses and Human Cancer: New Associations. Fred Hutchinson Cancer Center. Seattle, Washington. Invited Symposium Speaker.

2000 Viral Cause of Human Breast Cancer. North Shore University Hospital, Manhasset, New York. Invited Symposium Speaker.

1999 International Congress of the Hellenic Society for Breast Cancer Research, Herakleion, Crete. Invited Plenary Speaker.

1999 Duke University School of Medicine, Durham, North Carolina, Invited Speaker.

1999 International Congress of Virology, Sydney, Australia, Invited speaker.

1998 Symposium on Ion Channels and Disease. Kelso, Scotland. Invited Symposium Speaker.

1998 British Society for Nutritional and Environmental Medicine. Oxford, England. Invited Symposium Speaker.

1998 University of Tennessee School of Medicine, Memphis Tennessee, Invited Speaker.

1997 American Society for Virology Annual National Meeting. Hamilton, Montana. Invited Symposium Chair.

1996 Southern Association of Clinical Microbiologists Annual Meeting, Nashville, Tennessee. Invited Symposium Speaker

1996 American Society for Microbiology Annual National Meeting. New Orleans, Louisiana. Invited Symposium Speaker.

1996 Baylor University School of Medicine, Houston, Texas. Invited

Speaker.

1996 National Institutes of Health, National Cancer Institute Bethesda, Maryland. Invited Speaker.

1995 American Society for Virology Annual National Meeting. Austin, Texas. Invited Symposium Chair.

1995 American Society for Microbiology Annual Regional Meeting. Invited Symposium Chair.

1995 Calypte Biomedical, Berkeley, California. Invited Speaker.

1995 Rheumatology on the Beach. Destin, Florida. Keynote Speaker.

1995 Laboratory of Tumor Cell Biology Annual Meeting. Washington, D.C. Invited Symposium Speaker.

1994 Baylor University School of Medicine, Houston, Texas. Invited Speaker.

1993 National Sjögren's Syndrome Foundation Annual Meeting. New Orleans, Louisiana. Invited symposium speaker.

1992 Smith-Kline Beecham, Inc. King of Prussia, Pennsylvania. Invited speaker.

1992 Louisiana State University, Baton Rouge, Louisiana. Lectureship in Molecular Biology.

1991 University of Bern, Bern, Switzerland. Invited speaker.

1991 Robert Koch Institute, Berlin, Germany. Invited speaker.

1991 Max Plank Institute, Martinsreid (Munich), Germany. Invited speaker.

1991 Conference on Translational Controls in Virus-infected Animal Cells. Sigüenza, Spain. Invited symposium speaker.

1991 St. Luke's/Roosevelt Hospital at Columbia University. New York, New York. Invited speaker.

1991 Guthrie Research Foundation. Sayre, Pennsylvania. Invited speaker.

1991 Boehringer Ingelheim, Inc. Ridgefield, Connecticut. Invited speaker.

1991 New York University School of Medicine. New York, New York. Invited speaker.

1988-89 American Academy of Dermatology National Meetings. Washington D.C. and San Francisco. Invited symposium speaker.

- 1988 St. George's University Medical School. London, England. Invited speaker.
- 1988 University of Eppendorf-Krankenhaus, Hamburg, Federal Republic of Germany. Invited speaker.
- 1987 Conference on Membrane-mediated Controls. Hamburg, Federal Republic of Germany. Invited participant/session chairman.
- 1987 University of Hamburg. Hamburg, Federal Republic of Germany. Invited speaker.
- 1987 National Medical Association National Meeting. New Orleans. Invited symposium speaker.
- 1986 International Symposium on Immunological Adjuvants and Nonspecific Resistance to Microbial Infections. Columbia, MD. Invited speaker.
- 1984 Alberta Heritage Foundation Lecture. University of Edmondton, Alberta, Canada. Invited speaker.
- 1982 Conference on Membrane-mediated Controls. Hamburg, Federal Republic of Germany. Invited speaker.

Honors, Awards and Fellowships:

- 2020 Health Care Hero (*City Business*)
- 2015 100 Global Thinkers (*Foreign Policy*)
- 2015 Health Care Hero (*City Business*)
- 2003-2019 Research Scholar Awards (10 awards) - Tulane University
- 1996 Dean's Distinguished Faculty Forum
- 1994 Louisiana Innovator of the Year
- 1991 Fogarty Foundation Senior International Fellowship
- 1991 Alexander von Humboldt Award
- 1990 Outstanding Young Investigator Award (Tulane University)
- 1987 Owl Club Teaching Award.
- 1979-1981 NIH Individual Postdoctoral Research Fellowship.
- 1978 Margaret Jane McKinney Lewis Fellowship.
- 1977-1979 University Fellow Award.
- 1974 Sigma Xi Undergraduate Research Award.
- 1969-1974 Indiana Scholarship Commission Fellowship.

Professional Societies:

American Association for the Advancement of Science

American Society for Microbiology.

American Society for Virology.

Public Service:

2005-2007 President, South Central Branch of the American Society for Microbiology.

2004-2012 Founding Editor-in-Chief of *Virology Journal* (BioMed Central).

1995-date All the Virology on the World Wide Web (www.virology.net) with Dr. David Sander.

National and International Committees, Official Appointments and Directorships:

2022 Chair, National Institute of Allergy and Infectious Diseases. NIH. *Biocontainment Facility Improvements and Building System Upgrades to Support Pandemic Preparedness Study Section*

2021 Chair, and Infectious Diseases. NIH. Infectious diseases and Immunology Study Section

2020 Chair, NIAID COVID-19 Review meeting

2018-20 Chair, 40-year review of the US-Israel Binational Agricultural Research and Development Fund.

2020 Deutsche Forschungsgemeinschaft (DFG - German Research Foundation), ad hoc reviewer

2020 National Institute of Allergy and Infectious Diseases. NIH. Member, HIV AIDS Clinical Trials Units Review Meeting

2020 National Institute of Allergy and Infectious Diseases. NIH. Chair, Advanced Development of Vaccine Candidates for Biodefense and Emerging Infectious Diseases.

2020 National Institute of Allergy and Infectious Diseases. NIH. Member, Targeted Prevention for Tick-borne Diseases.

2019 National Institute of Allergy and Infectious Diseases. NIH. Member, NIH Director's Early Independence Awards (DP5).

2018 Member, World Health Organization Lassa fever Taskforce.

2018 Member, Executive Committee, Global Viral Network.

2018 National Institute of Allergy and Infectious Diseases. NIH. Member, Cellular and Molecular Immunology-B Study Section (Special SEP, Member conflicts).

2018 World Health Organization Ebola and Lassa fever Roadmap Meetings. London. January 16-19, 2018. Invited consultant/speaker.

2012-2018 National Institute of Allergy and Infectious Diseases. NIH. Member, AIDS Study Section (AIDSRRRC).

2016,2017 National Institute of Allergy and Infectious Diseases. Member or Chair, 3 Special ZIKA virus Study sections.

2009, 2010 National Institute of Allergy and Infectious Diseases. NIH. Chair, AIDS Vaccine Study Section.

2009 National Institute of Allergy and Infectious Diseases. NIH. Chair. Calicivirus Study Section.

2009 National Institute of Allergy and Infectious Diseases. NIH. Co-Chair, Regional Centers of Excellence for Biodefense Study Sections.

2008-2011 National Institute of Allergy and Infectious Diseases. NIH. Member, NIAID Conferences Study Section.

2007-2010 National Institutes of Health. Member. "Bugs and Drugs" Study Section.

2003-2007 National Institute of Allergy and Infectious Diseases. Chair, SBIR Biodefense Study Section.

2003 National Institute of Allergy and Infectious Diseases. Chair, Vaccines Adjuvants Therapeutics and Drugs for Biodefense Study Section.

2003 National Institutes of Health. Member, SBIR Biodefense Study Section.

2002 National Heart, Lung and Blood Institute, NIH. Member, Special AIDS Study Section.

2002 National Heart, Lung and Blood Institute, NIH. Member, Special AIDS Study Section.

2001 National Institute of Allergy and Infectious Diseases, NIH. Member of 2 Special AIDS Molecular Biology Study Sections.

2000 Institute of Women's Health. National Institutes of Health. Invited participant in research hearings.

1998 National Institute of Allergy and Infectious Diseases. NIH. Chair, Special AIDS Molecular Biology Study Section.

1998 National Institute of Diabetes and Digestive and Kidney Diseases. NIH. Invited participant in special panel to evaluate "Environmental etiology of type I diabetes: viruses and other factors."

1998 North Carolina Institute of Technology, External reviewer.

1997 British Medical Devices Agency, Invited participant in research hearings.

1997-2008 National Institute of Allergy and Infectious Diseases. Member, Centers for AIDS Research Study Section.

1997 National Institute of Allergy and Infectious Diseases, NIH. Member, or Chair of 3 Special AIDS Molecular Biology Study Sections.

1996 National Institute of Allergy and Infectious Diseases, NIH. Member, AIDS Molecular Biology Study Section.

1995 National Institute of Allergy and Infectious Diseases, NIH. Member, AIDS Molecular Biology Study Section.

1995 National Institute of Allergy and Infectious Diseases, NIH. Member, four Special AIDS Study Sections.

1995 Research Council of British Columbia. Member site visit committee.

1994 National Institute of Allergy and Infectious Diseases, NIH. Member, AIDS Molecular Biology Study Section.

1994 National Institute of Allergy and Infectious Diseases, NIH. Member, two Special AIDS Study Sections.

1994 National Heart, Lung and Blood Institute, NIH. Member of two Special AIDS Study Sections.

1994 National Institute of General Medical Sciences, NIH. Member, General Clinical Research Center Study Section.

1993-1997 NIH, Member Reviewers Reserve.

1993 National Institute of Allergy and Infectious Diseases, NIH. Member, Chronic Fatigue Syndrome Study Section.

1988-1993 National Institute of Allergy and Infectious Diseases, NIH. Member, AIDS Molecular Biology Study Section (ARRC, Chartered 1990).

1992 World Health Organization, Member, Advisory panel on unexplained immunodeficiency without evidence of HIV infection.

1992 Public Health Service, Member, Centers for Disease Control Advisory panel on idiopathic CD4+ lymphocytopenia.

1992 National Institute of General Medical Sciences, NIH. Member, MBRS Study Section.

1987-date National Research Council of Canada. Grant reviewer.

1989, 1992 American Society for Microbiology Annual Meeting Committee

1988 Presidential Commission on the Human Immunodeficiency Virus Epidemic. Invited participant in research hearings.

Committees (Intramural):

2006-date	Emergency Preparedness Committee (member 2006-11, Faculty Chair 2012-date)
1994-2000 2011-2015	Personnel and Honors Committee.
1988-date	Institutional Biosafety Committee (Vice-Chair, 1988-95; Chair, 1995-2002).
2006-date	Biomedical Sciences Steering Committee (Ex officio)
2006-2011	Graduate Council
1999-2006	University Senate (elected 1999; re-elected 2002)
2000 - date	Department of Microbiology and Immunology Promotion and Tenure Committee (Chair or Member)
2001	Molecular and Cellular Biology Program Review (Chair)
2000	University Senate Governance Review Committee
1998-date	Dean's Scholars Selection Committee
1998	Chancellor's Teaching Scholar Award Committee
1998	Search Committee for Chair of Pathology
1997-1999	Faculty Advisory Committee (elected 1997)
1997	Information Technology Committee (Chair Research subcommittee)
1997	Faculty Incentive Committee
1996-1999	University Senate Budget Advisory Committee
1995	Tulane Center for Molecular Medicine Organizing Committee
1995	Tulane Center for Infectious Diseases Organizing Committee
1994-2000	Molecular and Cellular Biology Steering Committee (Program Co-Director 94-95)
1990-date	Tulane: LSU General Clinical Research Center Steering Committee
1988-1999	Student Services Committee.
1991-1993	Molecular and Cellular Biology Curriculum Committee
1988, 1995	Shaffer Award Competition Committee.

1987-1989 Graduate School Curriculum Committee.

1986-1993 American Cancer Society Institutional Grant Committee.

Dissertation/Thesis Committees:

Biomedical Sciences - Tulane University

Phil Ferro, Ph.D. (2007, Committee Chair, currently Director, Special Projects, ASPR, Director, Office of Portfolio Management, Biomedical Advanced Research and Development Authority, Assistant Secretary for Preparedness and Response (ASPR), Department of Health and Human Services).
David Nielson (Committee Chair)
Lilia Melnick (2011, Committee Chair, LEQSF Fellowship)
Nathalia Holt, Ph.D. (Acting Committee Chair, Major Advisor Dr. Paula Cannon)
Racheal Yenni (2016, Committee Chair)
Matt Boisen (2014 Committee Chair)
Luis Branco (2012, Committee Chair, Currently CSO, Zalgen, LLC)
Eric Mucker (2014, Acting Committee Chair)
Jessica Grove, Ph.D (2016, Committee Chair, LEQSF Fellowship)
Jennifer Spence, Ph.D (2015, Committee Chair, LEQSF Fellowship)
Trevor Gale, . Ph.D. (2017, Committee Chair; currently >>>)
Brandon Beddingfield, PhD (2020, Committee Chair; currently post-doctoral fellow Chad Roy lab TNPRC)
Andrew Hoffman, Ph.D. (2020 Committee Chair; currently post-doctoral fellow Garry lab)
Allison Smither, Ph.D. (2021, Committee Chair; currently post-doctoral fellow University of Texas Medical Branch at Galveston)
Antoinette Bell-Kareem (Committee Chair)
Alex Berry (Committee Chair)
Joe Corey Berta (Committee Chair)

Bioinnovation Program

Kaylynn Genemaris (Committee Chair (2023; currently Food and Drug Administration)
Karissa Chao (Committee Chair)

Microbiology and Immunology-Tulane Medical School

Lynn Bonham, Ph.D. (1987)
Karin Levesque, Ph.D. (1990)
M. Nasar Qureshi, M.D., Ph.D. (1990, Committee Chair; currently Mt. Sinai Medical School, Associate Professor)
Lucy Freytag, Ph.D. (1991)
Darren Hart, Ph.D. (1993, Committee Chair n; currently LSUHSC)
Scott A. Tenenbaum, M.S., Ph.D. (1994, Committee Chair; postdoctoral fellow Duke University, currently Albany Medical College, Tenured Associate Professor)
Thomas G. Voss, Ph.D. (1994, Committee Chair; currently Vanderbilt University Professor)
Paul Gatti, Ph.D. (1998, Committee Chair; currently CSO-founder, Biocompare)
Bongkun Choi, Ph.D. (1998, Committee Chair; currently Samsung Corp., Korea)
Angelique Habis, Ph.D. (2000)
Joshua M. Costin, Ph.D. (2005, Committee Chair, currently post-doctoral fellow Florida Gulf Coast University)

Bruno Sainz, Ph.D. (2005, Committee Chair; NIH individual predoctoral fellowship, post-doctoral fellow at Scripps; currently: University of Chicago)

Yancey Hrobowski, Ph.D. (2006, Committee Chair, currently Director Virology Laboratory for the District of Columbia)

Mark Soboleski. Ph.D.

Molecular and Cellular Biology Program - Tulane University

Douglas R. Plymale, M.B.A., J.D., Ph.D. (1996; Committee Chair; postdoctoral fellow Harvard University; Tulane Law School; currently in private patent law practice)

Sara Sobel, M.D., Ph.D. (Committee Chair; currently University of Texas at Houston Assistant Professor)

David Sander, Ph.D. (1997, Committee Chair; LEQSF Fellowship; currently CSO Sander and Associates)

Heather Jaspan M.D./Ph.D. (1999, Committee Chair; NIH Individual Predoctoral Fellowship Awardee, Residency Univ. Washington and University, Johns Hopkins University faculty, currently University of Cape Town/University of Washington faculty)

Sonja Sumac, Ph.D. (1998)

Jing-Zhou Hou, Ph.D. (1998)

Laura Simpson, Ph.D. (1998; Tulane Regional Primate Center)

Anita M. Trichel, DVM, Ph.D. (1998; Tulane Regional Primate Center)

Sangeeta L. Peshori, Ph.D. (1999)

Debra Sullivan, Ph.D. (1999)

Hong Du, Ph.D. (1999)

Karen Ruli, Ph.D. (2000)

Raj Kalkeri Ph.D. (Committee Chair; 2000; currently Project Director, Vertex Pharmaceuticals, Cambridge, MA)

Micheal Bolton, Ph.D., M.D. (Committee Chair; 2001; LEQSF Fellowship; University of California Medical School, M.D. 2005, currently: Seattle Veteran's Hospital)

Sarah Nangle, Ph.D. (2006, Committee Chair)

Baiking Pei, Ph.D.

Chandtip Chandhasin, Ph.D.

Samantha Finstad, Ph.D.

Kristin Ruff, Ph.D.

Claudia Copeland, Ph.D.

Guy Odum, Ph.D.

Harris McFerrin, Ph.D.

Ali Shibahi, Ph.D. (2009, Committee Chair)

Biochemistry- Tulane Medical School

Rana Khan

Nilufar Inamdar

Geetha Kothandaraman

Pharmacology- Tulane Medical School

Claire Moss

Issac Rondon, Ph.D. (1993)

Khatidja Ali, M.S. (2000)

Christopher Williams, Ph.D. (2003)

Physiology- Tulane Medical School

Steve Antrobus (M.S., 1991)

Chemistry- Tulane University

Cheng-Feng Chiang (Ph.D. 1995, Committee Co-Chair; currently Clarke Medical School, Atlanta)

Microbiology, Immunology and Parasitology- LSU Medical School
Angela Martin, M.S. (1988); Ph.D. (1992)
Al Mock, M.S. (1999)

Biochemistry and Molecular Biology - LSU Medical School
James B. Chen, Ph.D. (1990)
Carol B. Hovda (M.S. (1994)

University of British Columbia. External reviewer. Dissertation committee of Paul E. Kowalski, Ph.D. (1998, Major advisor: Dr. Dixie Mager).

University of Minho in Porto, Portugal. PhD Thesis committee
Tiago Mota, Ph.D. (2019, Major Advisor: Dr. Matthias Schnell).

Post-doctoral students

Tam David-West, Ph.D.
Linda L. Moore, Ph.D.
Yan Cao, M.D.
Scott A. Tenenbaum, Ph.D. (Judith Graham Poole Postdoctoral Fellowship)
Ramesh Prabhu, Ph.D.
Suzanne Tomchuck, Ph.D.
Lilia Melnik, Ph.D.
Jessica N. Hartnett, Ph.D.
Lina M Moses, Ph.D.
Andrew Hoffmann, Ph.D.

Publications in Referred Journals:

Straub, S.X., Garry, R.F., and Magee, W.E. Interferon induction by poly I: poly C enclosed in phospholipid particles. *Infection and Immunity* 10:783-792, 1974.

Bishop, J.M., Maldonado, R.L., Garry, R.F., Allen, P.T., Bose, H.R. (Jr.), and Waite (Fairfax), M.R.F. Effect of medium of lowered NaCl concentration on virus release and protein synthesis in cells infected with reticuloendotheliosis virus. *Journal of Virology* 17:446-452, 1976.

Moore, R.N., Berry, L.J., Garry, R.F., and Waite (Fairfax), M.R.F. Effect of Sindbis virus infection on hydrocortizone-induced hepatic enzymes in mice. *Proceedings Society for Experimental Biology and Medicine* 157:125-128, 1978.

Bell, J.W., Garry, R.F., and Waite (Fairfax), M.R.F. Effect of low NaCl medium on the envelope glycoproteins of Sindbis virus. *Journal of Virology* 25:764-769, 1978.

Garry, R.F., Bishop, J.M., Parker, S., Westbrooke, K., Lewis, G., and Waite (Fairfax), M.R.F. Na⁺ and K⁺ and the regulation of protein synthesis in Sindbis virus-infected chick cells. *Virology* 96:108-120, 1979.

Garry, R.F. and Waite (Fairfax), M.R.F. Na⁺ and K⁺ and the regulation of the interferon system in chick cells. *Virology* 96:121-128, 1979.

King, C.C., King, M.W., Garry, R.F., Wan, K.M., Ulug, E.T., and Waite (Fairfax), M.R.F. Effect of incubation time on the generation of defective interfering particles during serial undiluted passage of Sindbis virus in Aedes albopictus cells. *Virology* 96:229-238, 1979.

Garry, R.F., Westbrooke, K., and Waite (Fairfax), M.R.F. Differential effects of ouabain on host- and Sindbis virus-specified protein synthesis. *Virology* 99:179-182, 1979.

Moore, R.N., Shackelford, G.M., Garry, R.F., and Berry, L.J. Effect of Sindbis virus infection on survival of mice in the cold. *Journal of Applied Physiology* 47:923-926, 1979.

Sanders, B.G., Wan, K.M., Kline, K., Garry, R.F., and Bose, H.R., Jr. Chicken fetal antigens: Role as cell surface receptors for Sindbis virus hemagglutinin. *Virology* 106:183-186, 1981.

Garry, R.F., Moyer, M.P., Bishop, J.M., Moyer, R.C., and Waite (Fairfax), M.R.F. Transformation parameters induced in chick cells by incubation in media of altered NaCl concentrations. *Virology* 111:427-439, 1981.

Lewis, R.B., McClure, J., Rup, B.J., Niesel, D.N., Garry, R.F., Hoelzer (Pierce), J., Nazerian, K., and Bose, H.R., Jr. Avian reticuloendotheliosis virus: Identification of the hematopoietic target cell for transformation. *Cell* 25:421-432, 1981.

Garry, R.F. and Bose, H.R., Jr. Secretion of a virus-regulated factor by clonal variants of reticuloendotheliosis virus-transformed cells. *Virology* 113:403-407, 1981.

Garry, R.F., Ulug, E.T., and Bose, H.R., Jr. Induction of stress proteins in Sindbis virus- and vesicular stomatitis virus-infected cells. *Virology* 129:319-332, 1983.

Ulug, E.T., Garry, R.F., and Bose, H.R., Jr. Alterations in monovalent cation transport in Sindbis virus-infected chick cells. *Virology* 132:118-130, 1984.

Garry, R.F., Shackelford, G.M., Berry, L.J. and Bose, H.R., Jr. Inhibition of hepatic phosphoenolpyruvate carboxykinase by avian reticuloendotheliosis viruses. *Cancer Research* 45:5020-5026, 1985.

Garry, R.F., Bostick, D., Schram, R., and Waite (Fairfax), M.R.F. The ratio of plasma membrane cholesterol to phospholipid and the inhibition of Sindbis virus maturation by low NaCl medium. *Journal of General Virology* 66:1171-1177, 1985.

Garry, R.F. and Bostick, D.A. Intracellular K⁺ and the expression of transformation parameters by chick cells transformed with the Bryan strain of Rous sarcoma virus. *Virology* 150:439-450, 1986.

Rasheed, S., A. A. Gottlieb, and R.F. Garry. Cell killing by UV-inactivated human immunodeficiency virus. *Virology* 154:395-400, 1986.

Garry, R.F., Bostick, D.A., and Ulug, E.T. Sindbis virus increases hexose transport in quiescent cells. *Virology* 155:378-391, 1986.

Garry, R.F. and Bostick, D.A. Induction of the stress response: Alterations in membrane-associated transport systems and protein modification in heat shocked or Sindbis virus-infected cells. *Virus Research* 8:245-259, 1987.

Garry, R.F., and Bose, H.R., Jr. Autogenous growth factor production by reticuloendotheliosis virus-transformed hematopoietic cells. *J. Cellular Biochemistry* 37:327-338, 1988.

Henderson, L.A., Qureshi, M.N., Rasheed, S., and Garry, R.F. HIV induced cytotoxicity for CD8⁺ cells from some normal donors and virus specific induction of a suppressor factor. *Clinical Immunol. Immunopath.* 48:174-186, 1988.

Moore, L.L., Bostick, D.A., and Garry, R.F. Sindbis virus infection decreases intracellular pH: Alkaline medium inhibits processing of Sindbis virus polyproteins. *Virology* 166:1-9, 1988.

Garry, R.F., Witte, M., Gottlieb, A.A., Elvin-Lewis, M., Gottlieb, M., Witte, C., Alexander, S.S., Cole, W.R., Drake, W.L., Jr. Documentation of an AIDS virus infection in 1968. *Journal of the American Medical Association* 260:2085-2087, 1988.

Garry, R.F. Poliovirus protease 2A is required for interference with vesicular stomatitis virus-induced protein synthesis. *Arch. Virology* 103:133-137, 1988.

Garry, R.F., Witte, M., Gottlieb, A.A., Elvin-Lewis, M., Gottlieb, M., Witte, C., Alexander, S.S., Cole, W.R., Drake, W.L., Jr. HIV infection in 1968 (letter). *Journal of the American Medical Association* 261:2199, 1989.

Garry, R.F. Alteration of intracellular monovalent cation concentrations accounts for inhibition of protein synthesis by a poliovirus mutant which encodes a defective 2A protease. *Virus Research*, 13:129-142, 1989.

Gallagher, W.R., Ball, J.M., Garry, R.F., Griffin, M.C., and Montelaro, R.C. A general model for the transmembrane proteins of HIV and other retroviruses. *AIDS Research and Human Retroviruses* 5:431-440, 1989.

Ulug, E.T., Garry, R.F., and Bose, H.R., Jr. The role of monovalent cation transport in Sindbis virus maturation and release. *Virology* 172:42-50, 1989.

Talal, N., Dauphinee, M.J., Dang, H., Alexander, S.S., Hart, D.J., and Garry, R.F. Evidence suggesting a retroviral etiology for autoimmune diseases. *Progress in Immunology* 7:837-841, 1989.

Qureshi, M.N, Coy, D.H., Garry, R.F., and Henderson, L.A. Characterization of a putative cellular receptor for the HIV-1 TM glycoprotein using synthetic peptides. *AIDS* 4:553-558, 1990.

Garry, R.F., and Witte, M. Early case of AIDS in the United States (letter). *Nature* 347:509, 1990.

Talal, N., Dauphinee, M.J., Dang, H., Alexander, S.S., Hart, D.J., and Garry, R.F. Detection of serum antibodies to retroviral proteins in patients with

primary Sjögren's syndrome (autoimmune exocrinopathy). *Arthritis and Rheumatism* 33:774-781, 1990.

Talal, N., Garry, R.F., Alexander, S.S., Dauphinee, M.J., Ballester, A., Takei, M., and Dang, H. A conserved idiotypic and antibodies to retroviral proteins in systemic lupus erythematosus. *J. Clinical Invest.* 85:1866-1871, 1990.

Garry, R.F. Extensive antigenic mimicry by retrovirus capsid proteins. *AIDS Research and Human Retroviruses* 6:1361-1362, 1990.

Garry, R.F., Fermin, C.D., Hart, D.J., Alexander, S.S., Donehower, L.A., and Luo-Zhang, H. Detection of a human intracisternal A-type retroviral particle antigenically-related to HIV. *Science* 250:1127-1129, 1990.

Miller, M.A., Garry, R.F., Jaynes, J.M., and R.C. Montelaro. A structural correlation between lentivirus transmembrane proteins and natural cytolytic peptides. *AIDS Research and Human Retroviruses* 7:511-519, 1991.

Garry, R.F., Kort, J.J., Koch-Nolte, F., and Koch, G. Similarities of viral proteins and toxins that interact with monovalent cation channels. *AIDS* 5:1381-1384, 1991

Stransky, G., Garry, R.F., and Gay, S. Detection of p24 in HIV-infected cells embedded in LR White and Lowicryl K4M. *Histochemical Journal*, 23:381-384, 1991.

Dang, H., Feghali, C. A., Dauphinee, M.J., Talal, N., Garry, R.F., Seibold, J.R., Medsger, T.A., Jr., Alexander, S. Serum antibody to retroviral proteins in systemic sclerosis. *Arthritis and Rheumatism* 34:1336-1337, 1991.

Garry, R.F. Intracisternal A-type retroviruses and immune dysfunctions. *The Lancet* 340: 787-788, 1992.

Garry, R.F., Hart, D.J., Tenenbaum, S.A., Luo-Zhang, H., Breeding, S.A.L., and Alexander, S.S. Sjögren's Syndrome and retroviral infection. *Arthritis and Rheumatism* 35, 1405, 1992.

Garry, R.F. and Koch, G. Tat contains a sequence related to snake neurotoxins. *AIDS* 6, 1541-1542, 1992.

Fermin, C.D., and Garry, R.F. Cytopathic effects linked to interactions of human immunodeficiency virus with the cell surface. *Virology* 191, 941-946, 1992.

Garry, R.F., and Fermin, C.D. Viral burden in AIDS. *Nature* 365, 301-302, 1993.

Tenenbaum, S.A., Leissinger, C.A., and Garry, R.F. Seroreversion in AIDS. *Journal of the American Medical Association*. 270:2178, 1993.

Garry, R.F. Sindbis virus-induced inhibition of protein synthesis is partially reversed by medium containing an elevated potassium concentration. *J. General Virology* 75, 411-415, 1994.

Tenenbaum, S.A. Voss, T.G., Gallaher, W.R. and Garry, R.F. Sequence similarities between retroviral proteins and components of the spliceosome. *AIDS Res. Human Retro.* 10, 521-522. 1994.

Gallaher, W.R., Ball, J.M., Garry, R.F., Martin, A.M., and Montelaro, R.C. A general model for the surface glycoproteins of HIV and other retroviruses. *AIDS Research and Human Retroviruses* 11, 191-201, 1995.

Garry, R.F. Sequence similarities between latency membrane protein LMP-1 of Epstein-Barr virus, integral membrane protein p12^I of human T-cell leukemia/lymphotropic virus type 1, E5 transformation protein of bovine papilloma virus, and the transmembrane proteins of oncoviruses. 11, 431-432, 1995.

Cuéllar, M.L., Scopelitis, E., Tenenbaum, S.A., Garry, R.F., Silveira, L.H., Cabrera, G., and Espinoza, L.R. Serum antinuclear antibodies in women with silicone breast implants. *J. Rheumatology* 22, 236-240, 1995.

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Patents

Arenavirus monoclonal antibodies and use

Patent number: 11,198,723

Date of Patent: December 14, 2021

Assignees: The Administrators of the Tulane Educational Fund, Zalgen Labs, LLC

Inventors: Luis M. Branco, Robert F. Garry, James E. Robinson, Erica Ollmann Sapphire, Kathryn M. Hastie, Thomas W., Geisbert

Compositions and methods for measles virus inhibition

Patent number: 9725487

Date of Patent: August 8, 2017

Assignees: The Administrators of the Tulane Educational Fund, Autoimmune Technologies, LLC

Inventors: Robert F. Garry, Russell B. Wilson

Antiviral rift valley fever virus peptides and methods of use

Patent number: 9556237

Date of Patent: January 31, 2017

Assignee: The United States of America, as represented by the Secretary of the Army, on behalf of the U.S. Army Medical Research Institute of Infectious Diseases
Inventors: Connie Schmaljohn, Robert F. Garry, Jeffrey W. Koehler, Mary Guttieri

[Peptide compositions and methods for inhibiting herpesvirus infection](#)

Patent number: 9434769
Date of Patent: September 6, 2016
Assignee: The Administrators of the Tulane Educational Fund
Inventors: Lilia I. Melnik, Robert F. Garry, Cindy A. Morris

[Influenza inhibiting compositions and methods](#)

Patent number: 9353157
Date of Patent: May 31, 2016
Assignees: The Administrators of the Tulane Educational Fund, Autoimmune Technologies, LLC
Inventors: Robert F. Garry, Russell B. Wilson

[Compositions and methods for coronavirus inhibition](#)

Patent number: 9056900
Date of Patent: June 16, 2015
Assignees: The Administrators of the Tulane Educational Fund, Autoimmune Technologies, LLC.
Inventors: Robert F. Garry, Russell B. Wilson

[Arenavirus inhibiting peptides and uses therefor](#)

Patent number: 8999925
Date of Patent: April 7, 2015
Assignee: The Administrators of the Tulane Educational Fund
Inventors: Jennifer S. Spence, Robert F. Garry

[Peptide compositions and methods for inhibiting herpesvirus infection](#)

Patent number: 8802106
Date of Patent: August 12, 2014
Assignee: The Administrators of the Tulane Educational Fund
Inventors: Lilia I. Melnik, Robert F. Garry, Cindy A. Morris

[Optimized dengue virus entry inhibitory peptide \(DN81\)](#)

Patent number: 8541377
Date of Patent: September 24, 2013
Assignee: Florida Gulf Coast University.
Inventors: Scott F. Michael, Sharon Isern, Robert F. Garry, Ram Samudrala, Joshua Costin, Ekachai Jenwitheesuk.

[Influenza inhibiting compositions and methods](#)

Patent number: 8604165
Date of Patent: December 10, 2013
Assignees: The Administrators of the Tulane Educational Fund, Autoimmune Technologies, LLC
Inventors: Robert F. Garry, Russell B. Wilson

[Treatment of influenza virus infection](#)

Patent number: 8598116
Date of Patent: December 3, 2013
Assignees: The Administrators of the Tulane, Educational Fund and Autoimmune Technologies, LLC

Inventors: Robert F. Garry, Russell B. Wilson

[Influenza inhibiting compositions and methods](#)

Patent number: 8222204

Date of Patent: July 17, 2012

Assignee: The Administrators of the Tulane Educational Fund and Autoimmune Technologies, LLC

Inventors: Robert F. Garry, Russell B. Wilson

[Flavivirus fusion inhibitors](#)

Patent number: 8153360

Date of Patent: April 10, 2012

Assignees: The Administrators of the Tulane Educational Fund, The Rockefeller University

Inventors: Robert F. Garry, Srikanta Dash, David H. Coy, Jane A. McKeating

[Flavivirus fusion inhibitors](#)

Patent number: 7854937

Date of Patent: December 21, 2010

Assignees: The Administrators of the Tulane Educational Fund, The Rockefeller University

Inventors: Robert F. Garry, Srikanta Dash, David H. Coy, Jane A. McKeating

[Influenza virus inhibiting peptides](#)

Patent number: 7491793

Date of Patent: February 17, 2009

Assignee: The Administrators of the Tulane Educational Fund

Inventors: Robert F. Garry, Russell Wilson

[Flavivirus fusion inhibitors](#)

Patent number: 7416733

Date of Patent: August 26, 2008

Assignees: The Administrators of the Tulane Educational Fund, The Rockefeller University

Inventors: Robert F. Garry, Srikanta Dash, David H. Coy, Jane A. McKeating

[Human endogenous retrovirus in breast cancer](#)

Patent number: 6670466

Date of Patent: December 30, 2003

Assignee: The Administrators of the Tulane Educational Fund

Inventor: Robert F. Garry

[Method for detecting anti-squalene antibodies](#)

Patent number: 6214566

Date of Patent: April 10, 2001

Assignee: The Administrators of the Tulane Educational Fund

Inventors: Pamela B. Asa, Robert F. Garry

[Method for detecting antipolymer antibodies and diagnosing silicone related disease \(SRD\) fibromyalgia and chronic fatigue syndrome \(CFS\)](#)

Date of Patent: November 10, 1998

Assignee: The Administrators of the Tulane Educational Fund

Inventors: Robert F. Garry, Scott A. Tenenbaum, Douglas R. Plymale

[Method to aid in the diagnosis of silicone related disease](#)

Patent number: 5620859
Date of Patent: April 15, 1997
Assignee: Administrators of the Tulane Educational Fund
Inventors: Robert F. Garry, Scott A. Tenenbaum, Douglas R. Plymale

Association between a novel human intracisternal A-type retroviral particle-type II (HIAP-II) and idiopathic CD4+ T-lymphocytopenia (ICL)

Patent number: 5580772
Date of Patent: December 3, 1996
Assignee: The Administrators of the Tulane Educational Fund
Inventor: Robert F. Garry, Jr.

The cellular receptor for the CS3 peptide of human immunodeficiency virus

Patent number: 5567805
Date of Patent: October 22, 1996
Assignee: Administrators of the Tulane Educational Fund
Inventors: Lee A. Henderson, David H. Coy, Robert F. Garry, Jr.

Methods of supporting a diagnosis of systemic lupus erythematosus

Patent number: 5364757
Date of Patent: November 15, 1994
Assignee: Administrators of the Tulane Educational Fund
Inventors: Robert F. Garry, Jr., Cesar D. Fermin, Steve S. Alexander, Jr.

Human intracisternal A-type retroviral particles associated with sjogren's syndrome

Patent number: 5344774
Date of Patent: September 6, 1994
Assignee: The Administrators of the Tulane Educational Fund
Inventors: Robert F. Garry, Jr., Cesar D. Fermin, Steve S. Alexander, Jr.

Methods and compositions for identifying and characterizing individuals having autoimmune rheumatic diseases

Patent number: 5320940
Date of Patent: June 14, 1994
Assignee: Board of Regents, The University of Texas System
Inventors: Norman Talal, Robert F. Garry

Truth in Testimony Disclosure Form
Dr. Robert F. Garry, PhD

Attachment C

Federal Grants or Contracts Related to Hearing's Subject Matter

Garry, Robert F
RFGARRY

Title: West African Emerging Infectious Disease Research Center

Major Goals: To develop the infrastructure, diagnostic assays, reagents, and computational tools to formulate effective strategies for discovering, mitigating, and preventing the emergence of human infectious diseases.

Project Number: 5U01AI151812-03

Name of PD/PI: (MPI) Andersen, K.G. (contact); Garry, R.F.; Sabeti, P.C.

Source of Support: Subaward from The Scripps Research Institute; prime source: NIAID

Primary Place of Performance: Scripps Research Institute

Project/Proposal Start and End Date: 05/01/2020-04/30/2025

Total Award Amount: 2021: \$1,757,660; 2022: \$1,848,854; 2023: \$1,741,300

Title: Immune Response to SARS-CoV-2 in Special Populations

Major Goals: This study will allow real time comparison of clinical course, virus sequence, and sero-response, collectively referred to as ClinSeqSer response, in a cohort of subjects, for the year following recruitment. The ClinSeqSer data gathered in this study will help us identify whether variants in host or virus have contributed to high mortality seen in Louisiana and will be useful for broader predictions of covid-19 response moving forward.

Project Number: 75D30120C08472

Name of PD/PI: Fusco, D.

Source of Support: CDC

Primary Place of Performance: Tulane University

Project/Proposal Start and End Date: 07/01/2020-06/30/2021

Total Award Amount: \$700,000

Person Months per budget period

Title: LA-CEAL: Louisiana Community-Engagement Research Alliance Against COVID-19 in Disproportionately Affected Communities

Major Goals: The goal is to utilize existing trusted partnerships with multiple stakeholders to leverage extant community resources and capabilities to develop and execute a rapid community-engaged action plan that will focus on the vulnerable populations in Louisiana, hardest hit by the COVID-19 pandemic.

Project Number: 1OT2HL158260

Name of PD/PI: (MPI) Krousel-Wood, M.A.(contact); Sarpong, D.F.

Source of Support: NHLBI

Primary Place of Performance: Tulane University

Project/Proposal Start and End Date: 09/01/2020-03/31/2022

Total Award Amount: \$2,400,000

Title: Tulane University COVID Antibody and Immunity Network (TUCAIN)

Major Goals: The goal is to mechanistically define the immune response to COVID-19 to determine how it protects individuals from infection and how long the protection lasts.

Project Number: 1U54CA260581

Name of PD/PI: Robinson, J.E.

Source of Support: NIH/NCI

Primary Place of Performance: Tulane University

Project/Proposal Start and End Date: 09/30/2020-09/29/2022

Total Award Amount: \$7,873,887

Title: Genomic Characterization and Surveillance of Microbial Threats in West Africa

Major Goals: Goal is to characterize and survey microbial threats across West Africa, such as Ebola, Lassa fever, and malaria, by building on a long-term collaborative effort among established academic, research and clinical centers in Nigeria, Senegal, Sierra Leone and partners in the US.

Project Number: 2U54HG007480-07

Name of PD/PI: Happi, C.

Source of Support: subaward from Redeemer's University (Nigeria); Prime source: NHGRI

Primary Place of Performance: Redeemer's University

Project/Proposal Start and End Date: 07/01/2017-06/30/2022

Total Award Amount: \$405,512