REQUEST FOR APPLICATIONS
PILOT PROJECTS

SUMMARY: The NIH/NIGMS-funded Louisiana Center for Lung Biology and Disease (CLBD) at LSU School of Veterinary Medicine (MPI: Tammy Dugas and Stephania Cormier) invites Pilot Project Investigators (PPIs) to submit applications for pilot research projects consistent with the research goals of this Center. The thematic focus of the CLBD encompasses mechanistic studies of respiratory diseases induced by both infectious (bacterial, viral, fungal) and non-infectious (e.g., environmental, cigarette, e-cigarette, aerosols, allergens) stimuli.

RESEARCH OBJECTIVES: The COBRE program seeks to promote the initiation, development, or expansion of unique, innovative, state-of-the-art biomedical and behavioral research at institutions in IDeA-eligible states. The long-term goal of the CLBD is to establish a Center of Excellence (COE) in pulmonary diseases in Louisiana that is highly interactive and consists of a dynamic group of investigators whose research focus is on understanding the molecular and cellular immunological mechanisms associated with the pathogenesis of both infectious and non-infectious lung diseases. This COBRE program will provide novel insights into the pathogenesis of devastating lung diseases that guide future improved strategies for treating and preventing these and other lung diseases in animal and human populations.

PILOT RESEARCH PROJECTS: The pilot research project application should describe the specific aims of the research on pulmonary disease and the goals for the first year and/or second year. The design principles supporting the research or the hypotheses to be tested should be delineated. Preliminary studies are required for COBRE projects. Moreover, each pilot project should critically assess the existing knowledge and approaches that have been or are being directed in the area with an emphasis on specifically how the multi-disciplinary COBRE approach will advance the field of pulmonary diseases. Each pilot project must identify a senior mentoring team that will assist the independent career development of the PPI Investigator and describe the institutional commitment to the PPI (support letters from the mentor and Investigator’s department chair). The goal of the mentoring program is achieving independent status by the Investigators through an acquisition of a major funded grant, such as an R01.

FUNDS AVAILABLE: Budgets should not exceed $50,000 in direct costs only for the time period of October 16, 2024, through October 15, 2025. Although the initial funding is limited for one year, the applicant should present a 2-year research plan although the PI’s second year funding ($50,000) is based on the satisfactory progress of the project. It is anticipated that at least 1-3 new pilot projects will be awarded. NIH - PHS 2590 progress reports are required at 1 year.
ELIGIBLE INDIVIDUALS: Faculty investigators of all ranks are eligible, but priority will be given to Early Stage and New Investigators for leading a Pilot Project. Individuals holding postdoctoral, trainee, or similar non-independent positions are not eligible to lead a Pilot Projects. For the purpose of eligibility, a PPI is defined either as (1) an individual who does not have or has not previously had an external, peer-reviewed Research Project Grant (RPG) or Program Project Grant (PPG) from either a Federal or non-Federal source that names that investigator as the PI or (2) an established investigator who is making a significant change to his/her career. Senior, funded investigators who are not making a significant career change are ineligible.

A PPI must hold a tenure-, clinical- or research-track full-time faculty appointment at Louisiana State University Baton Rouge or any other institutions in Louisiana at the time that the award is made. Moreover, a clear commitment to support this appointment and career development of this applicant independent of the outcome of this application must be demonstrated from the institution by a letter(s) from the appropriate senior institutional official(s). Applicants must propose a project that fits into the central theme of the CLBD-COBRE, i.e., respiratory diseases.

APPLICATION SUBMISSION: Interested investigators should submit an application that conforms to the NIH Exploratory/Developmental Research Grant Program (R21) guidelines that can be found at the website: http://grants.nih.gov/grants/funding/r21.htm. The application should be written as an NIH R21 application intended for submission to the NIH, containing an explanation of the type of preliminary results that will be generated by the COBRE project funding that are needed to render this application competitive for NIH R01 funding.

In the interest of competitiveness for NIH R21 or R01 funding, it is critical for applicants to articulate the potential translational application of the proposed research to human health. The applicant should also consider the overall fit of the aims of their application for the goals of this COBRE, which are to promote excellence in research in the broad area of pulmonary diseases. The deadline for submission of a letter of intent (LOI) is August 1, 2024. The LOI should have the Specific Aim (one) page and NIH Biosketch (5 pages). If you have questions about the pilot grant application process, please contact Dr. Tammy Dugas or Dr. Stephania Cormier.

The completed SF424 application (all sections) should be compiled as a PDF document and submitted electronically to:

Merilyn Jennings
Email: mjennings@lsu.edu

**Timeline**

- **Release Date:** July 15, 2024
- **Letter of Intent (open) submission deadline:** August 1, 2024
- **Proposal submission (by invitation only) deadline:** August 30, 2024
- **Peer Review Date(s):** September 16, 2024
- **Funding Start Date:** October 16, 2024 (earliest)

**ADDITIONAL REQUIREMENTS:** All applicants selected to receive CLBD project grant funding are required to submit an NIH R21 or R01 application and/or show that they are making significant progress towards submitting a competitive NIH application. Successful applicants are expected to attend the CLBD monthly meetings and present project progress reports as appropriate. Recipients will also be expected to actively participate in the CLBD seminar and work-in-progress series. A final report is required in the form of an NIH 2590 progress report. These reports should
state the original objectives of the project and indicate which of the objectives were addressed during the allotted 12-month time period. Appropriate tables and figures should be included to help clarify these issues. The final report must also include the title of the resulting grant proposal submitted to external federal agencies for funding, the date the proposal was submitted or will be submitted, and a list of any resulting articles submitted or published. A timetable should be provided that outlines a plan for seeking subsequent or supplemental extramural funding.

**BUDGET SPECIFICATIONS:** The total budget requested for each project is not to exceed $50,000 for the 12 months (April-December) or $100,000 for 24 months. If additional funds are needed, a thorough justification is needed which is subject to advisory committee (AC) approval. **Expenditures allowed:** 1) Technical staff salary support, 2) Research supplies and animal maintenance, 3) Equipment costing less than $10,000, 4) Supplies, 5) Domestic or foreign travel, and 6) Publication costs, including reprints. **Expenditures NOT allowed:** 1) Office equipment and supplies, 2) Computers or honoraria and travel expenses for visiting lecturers, 3) Dues and membership fees in scientific societies, 4) Rental of office or laboratory space or construction or building maintenance. We want to encourage the use of the COBRE scientific cores. Therefore, each project will receive an additional $2,500 credit for SVM-to cover the use of core facilities and services including the Pulmonary Immunopathology (PIPC), Molecular and Cell Biology (MCBC) and Inhalation and Infection Core (IIC) Cores. These funds will be expended to reduce by 25% actual costs of project investigators utilizing these Cores.

**UTILIZATION OF CORES:** The use of CLBD Cores, such as PIP, MCBC, and IIC should be included in the overall experimental design.

**FORMAT:** The NIH R21 format should be followed. The grant application should be compiled as a single PDF document after filling the SF424 form and emailed to Merilyn Jennings.

Furthermore, as per NIH requirement, for each of the projects that are selected for funding, please provide the following PHS 398 Forms:
- Face Page
- Project Summary
- Research Strategy Section
- Biographical Sketch
- Human Subjects and Clinical Trials Information Form (if applicable)
  - An approved IRB Protocol letter will be needed should your proposal is selected. If Human Subjects is part of the project, documentation of the human subject’s education is required.
- Vertebrate Animal Section (if applicable).
  - An approved IACUC Protocol letter will be needed should your proposal be selected.

**SELECTION:** LOIs will be screened initially by the MPIs for thematic fit and eligibility and if acceptable will be invited for submission of a full application. The AC of the CLBD-COBRE will review selected applications for merit and fit within the scope of the CLBD. Applications selected for funding by the AC will be forwarded to NIH for approval. NIH will then issue a notice of award pending just-in-time information and after approval. It is anticipated that funding will begin **October 16, 2024.** Applications are required to have approved IACUC and IRB protocols prior to receipt of funding. AC will use the NIH scale to review applications.

**MAJOR REVIEW CRITERIA:** The goals of this NIH-supported research are to advance our understanding of pulmonary diseases, to improve the prevention and control of pulmonary
diseases, and to enhance human health throughout the lifespan. Diversification of the biomedical research workforce and development of the next generation of independent investigators are also important objectives of the NIH extramural research portfolio and a specific goal of the NIH/NIGMS COBRE funding mechanism. A single-digit score and a bulleted list of strengths and weaknesses for each of the six review categories, as well as an overall priority score, using the following **NIH-based scoring scale** will be used (Note that an application does not need to be strong in all listed categories to be judged).

**Significance:** Does this study address an important health problem? If the aims of the application are achieved, how will scientific knowledge or clinical practice be advanced? What will be the effect of the study on the concepts, methods, technologies, treatments, services, or preventive interventions that drive this field?

**Innovation:** Is the project original and innovative? For example: Does the project challenge existing paradigms or clinical practice or address an innovative hypothesis or critical barrier to progress in the field? Does the project develop or employ novel concepts, approaches, methodologies, tools, or technologies?

**Investigator:** Is the investigator appropriately trained and well suited to conduct the proposed study? Is the proposed research appropriate to the experience level of the principal investigator and collaborators? If the principal investigator is a junior faculty member, has the applicant designated a senior mentor and a brief description of a mentoring plan? Does the investigative team bring complementary expertise to the project?

**Approach:** Are the conceptual or clinical framework, design, methods, and analyses adequately developed, well-integrated, well-justified, and appropriate to the aims of the project? Does the applicant acknowledge potential problems and propose alternative strategies?

**Environment, Collaborations, and Partnerships:** Does the scientific environment in which the study will be performed contribute to the probability of success? Does the proposed study benefit from unique features of the scientific environment or subject populations, or employ useful collaborative arrangements? Will there be collaborations to address the scientific questions in the proposed research area? Will the research fit into the Department and Institutional strategic plans and research?

**Terms of the Center for Lung Biology and Disease (CLBD)-COBRE award:**

- Awardees (PPIs) will be required to prepare a “personal mentoring or career development plan” with their mentor.
- Awardees will be required to attend CLBD meetings, such as grant proposal development, work in progress, journal club, seminars, and give an oral presentation at the Annual CLBD Research Retreat. Awardees’ lab members are also required to participate in CLBD activities.
- Awardees’ progress will be evaluated annually by the members of the AC).
- Awardees must acknowledge the COBRE grant support as well as COBRE Research Core support (if applicable) in publications, presentations, press releases, or other documents regarding research. A disclaimer should be included such as "Research reported in this publication was supported by the National Institute of General Medical Sciences of the National Institutes of Health under P20GM130555. The content is solely the responsibility of the authors and does not necessarily represent the official views of the National Institutes of Health."
Awardees must annually provide information on their publications, presentations, honors, grants, other recognitions, and service in study sections for the NIH progress report during the tenure of the COBRE grant.
Appendix: Services provided by CLBD-COBRE Research Cores

Pulmonary Immunopathology Core (PIPC) (Core lead: Masami Yoshimura, PhD, Co-I, Tomislav Jelesijevic, DVM, PhD)

1. Mouse colony management and breeding strategies for maintaining colonies.
2. Genotyping of mouse tail DNA via PCR.
3. Intranasal/intratracheal infection of mice.
4. Post-infection monitoring of mice.
5. Pulmonary function tests.
6. Mouse dissections and tissue collections.
8. Intratracheal instillation of reagents, such as LPS, bacteria, antibodies, and compounds.
9. Support for experimental design related to pathology sampling.
10. Tissue processing for paraffin-embedded tissue sections and stains.
11. Immunohistochemistry tissue staining.
12. Histological scoring of lungs and extrapulmonary tissues.
13. Support for flow cytometry and cell sorting.
14. Support for standard tests of immune function, including tests of cell mediated immunity (intracellular cytokine staining utilizing flow cytometry), in vitro assays, cytokine gene expression, or cytokine protein expression via ELISA.
15. Support for statistical analysis (parametric and non-parametric).

Molecular and Cell Biology Core (MCBC) (Core Lead: Konstantin Kousoulas, PhD, Co-I, Vladamir Chouljenko, PhD)

1. Overexpression vectors for transgene expression.
2. Site-directed mutagenesis.
4. Support for transfection using adenovirus and lentivirus vectors.
5. Real-time PCR gene quantification.
6. DNA microarray and RNAseq analyses
7. Support for evaluations using immunofluorescence and confocal microscopy, and interpretation of images obtained by scanning and transmission electron microscopy.
8. Consultation on molecular biology aspects to the COBRE investigators and to help develop or adopt new techniques as needed.

Inhalation and Infection Core (IIC) (Core Lead: Alexandra Noël, PhD; Co-I, Tirumalai Rangasamy, PhD)

1. Development of bacterial pneumonia in mice: The IIC will provide a BSL-2 research area to develop bacterial pneumonia, using *Staphylococcus aureus* [wild type *Staphylococcus aureus*, Δmsr S. aureus, S. aureus Newman strain and Methicillin-resistant *Staphylococcus aureus* (MRSA) strain USA300 LAC], *Streptococcus pneumonia*, *Klebsiella pneumoniae*, and Carbapenem-resistant *Klebsiella pneumoniae* induced respiratory infections in mice.
2. Bacterial preparation.
3. Bacterial inoculation – by oropharyngeal, intratracheal, and intranasal routes.
4. Bronchoalveolar lavage fluid collection and inflammatory cell phenotyping (total white blood cells, macrophages, neutrophils, lymphocytes, and eosinophils).
5. Enumeration of bacterial burden in the lungs and extrapulmonary organs of mice.
7. Isolation and characterization alveolar type II epithelial cells from mice.
8. Extracellular killing assays by infecting murine/human cells/lines with different bacterial pathogen.
9. Generation of macrophages from bone marrow cells of mice.
10. Purification of neutrophils from bone marrow cells of mice using EasySep neutrophil enrichment kit.
11. Measurement of lung injury [mean linear intercept (distance between alveolar septa), alveolar diameter, alveolar perimeter, septal thickness, surface-to-volume ratios, and total number of alveoli/field] in mice using Metamorph software program.

In years 3 to 5 of this grant, the IIC will provide a BSL-2 research environment with fully integrated inhalation exposure systems for PM$_{2.5}$, second-hand smoke (SHS), and electronic-cigarette (e-cig) aerosols, to support *in vitro* air-liquid interface (ALI) and *in vivo* experiments in the field of respiratory infectious diseases and lung biology.

12. PM$_{2.5}$ inhalation exposure system.
13. SHS inhalation exposure system.
14. E-cig aerosol inhalation exposure system.
15. ALI *in vitro* exposure system.
16. Lung function testing in mice within a BSL-2 research environment.