

Pfizer Research Grant Request for Proposals

Competitive Grant Program – Pfizer Internal Review Process

Epidemiology & Real-World Evidence of SUL-DUR in Special Populations with CRAB Infection

Overview

This competitive research program aims to characterize the epidemiology and clinical burden of carbapenem-resistant Acinetobacter baumannii (CRAB) infections in China and to generate real-world evidence on the effectiveness and safety of sulbactam-durlobactam (SUL-DUR) in routine clinical practice[1,2]. The study will focus on critically ill adult patients with suspected or confirmed CRAB infections, including special populations such as postoperative neurosurgical patients, liver or kidney transplant recipients, burn patients. Additionally, the study will explore treatment patterns and the impact of SUL-DUR initiation timing on key clinical outcomes in these high-risk populations.

Geographic Scope

Mainland China

Project Types and Area of Interest

Potential applicants are requested to submit research proposals to study the epidemiology of CRAB infections, real-world evidence on effectiveness, safety of SUL-DUR, and influence of SUL-DUR treatment timing on clinical outcomes for CRAB infections in special adult populations:

- Postoperative neurosurgical patients
- Liver or kidney transplant recipients
- Burn patients

Key Milestones

Submission Deadline	Anticipated Grant Award Notification	Anticipated Project Start Date/Duration
3 Feb 2027	16 Apr 2027	16 Jul 2027

Funding Range and Project Length

Individual projects requesting up to $58,200 will be considered.

Maximum project length is 1 year.

I. Eligibility

Geographic Scope:

- Mainland China

Applicant Eligibility Criteria

- The institution and Principal Investigator (PI) must be based in mainland China noted above.
- Only organizations are eligible to receive grants, not individuals or medical practice groups (i.e., an independent group of physicians not affiliated with a hospital, academic institution, or professional society).
- If the project involves multiple departments within an institution and/or between different institutions / organizations / associations, all institutions must have a relevant role and the requesting organization must have a key role in the project.
- The PI must have a medical or postdoctoral degree (MD, PhD, or equivalent), an advanced nursing degree (BSN with a MS/PhD), or a degree in Pharmacy, Physiotherapy, or Social Work.
- The applicant must be the PI or an authorized designee of such individual (e.g., PI's research coordinator).
- The PI must be an employee or contractor of the requesting organization.
- Requesting organization must be legally able to receive award funding directly from Pfizer Investment Co., Ltd.. We strongly recommend that applicants confirm this with their organization or institution prior to submitting an application. Grants awarded to organizations that are subsequently found to be unable to accept funding directly from Pfizer Investment Co., Ltd. may be subject to rescission.

II. Requirements

Primary Area of Interest:

- Infectious Disease - Bacterial

General Area of Interest for this RFP:

- Projects that will be considered for Pfizer support will focus on evaluating the epidemiology, disease burden, and treatment patterns of carbapenem-resistant Acinetobacter baumannii (CRAB) infections, as well as generating real-world evidence on the effectiveness and safety of SUL-DUR. Areas of interest include clinical success, microbiological success, mortality, adverse events, and the impact of treatment timing on outcomes, with a focus on:
    - Postoperative neurosurgical patients
    - Liver or kidney transplant recipients
    - Burn patients

This RFP does not include requests for Pfizer therapeutic agents.

Expected Approximate Monetary Range of Grant Applications

IMPORTANT: Grants will be distributed following a fully executed agreement and submission of Final Protocol, Documentation of IRB/IEC Approval, Regulatory Approval (if applicable), Exemption or Waiver.

- Individual projects requesting up to $58,200 will be considered. A total of 3 qualified proposals will be awarded.
- Award amounts include direct costs, institutional overhead costs (capped at 28% per Pfizer policy), and indirect costs.

Key Dates:



IMPORTANT: Be advised applications submitted after the due date will not be reviewed.

*Please note the deadline is 23:59 Eastern Standard Time (e.g., New York, GMT -5)

How to Submit:

IMPORTANT: Please read this section carefully since applications submitted not following these instructions will not be accepted and will be cancelled.

- Click here to navigate to this Competitive Grant Program's unique Program Application Form.
- First-time users should create a Requester Portal account by clicking **Not a member?**
- OR
- Navigate to: https://pfizermarcie.my.site.com/pfzrequester and sign in.
- In the upper navigation bar, click **Grant Programs** to see all available programs.
- Select **Research**
- Click **Apply** on the "**2027 I&I CN SUL-DUR EPI RWE RES**" to open the Program Application Form.
- Requirements for submission:
    - Complete all required sections of the online application
    - **IMPORTANT**: Upload proposal (see Appendix) in the Proposal/Protocol field.

- For more information on navigating Pfizer's new grant management system (MARCIE), please refer to this helpful User Guide.

Questions:

- Please click here to view "Frequently Asked Questions" regarding the Competitive Grant Program.
- If you have questions regarding this RFP, please direct them in writing to the Grant Officer, Juan Liu (GMGChina@pfizer.com), with the subject line "2027 I&I CN SUL-DUR EPI RWE RES – 3Nov2026"

Review and Approval Process

- Grant requests received in response to a general RFP are reviewed by Pfizer to make final grant decisions.

Mechanism by which Applicants will be Notified:

- All applicants will be notified via email by the dates noted above.
- Applicants may be asked for additional clarification during the review period.

Grant Agreements:

- If your grant is approved, your institution will be required to enter into a written grant agreement with Pfizer. Please click here to view the core terms of the agreement.
- Under Pfizer's competitive grant program, modifications to grant agreements will not be reviewed unless a genuine conflict exists as between applicable law and the terms of the relevant grant agreement. Applicant is encouraged to share the core terms with counsel for approval prior to submitting an application.
- Except where prohibited by applicable law and, in any case, subject to review by Pfizer Legal, payment of grant funding may only be paid to the grantee organization.

References

1. Kaye KS et al. Lancet Infect Dis. 2023 Sep;23(9):1072-1084.2.
2. National Medical Products Administration (NMPA). Xacduro® [Sulbactam Sodium for Injection/Durlobactam Sodium for Injection(co-packaged)] Prescribing Information. December 1, 2025.

About Pfizer Grants

Pfizer supports the global healthcare community's independent initiatives (e.g., research, quality improvement or education) to improve patient outcomes in areas of unmet medical need that are aligned with Pfizer's medical and/or scientific strategies.

Pfizer's competitive grant program involves a publicly posted general Request for Proposal (RFP) that provides detail regarding a general area of interest, sets timelines for review and approval, and uses an internal Pfizer review process to make final grant decisions. Organizations are invited to submit an application addressing the research gaps as outlined in the specific RFP.

For all Investigator Sponsored Research (ISRs) and general research grants, the grant requester (and ultimately the grantee) is responsible for the design, implementation, sponsorship, and conduct of the

independent initiative supported by the grant, including compliance with any regulatory requirements. Pfizer must not be involved in any aspect of study protocol or project development, nor the conduct or monitoring of the research program. An ISR grant request cannot be submitted for a study that has already commenced and was not originally supported by Pfizer.

Appendix

IMPORTANT: RFP Submission Requirements

Applications will be accepted via the online portal listed in the How to Submit section. Project Proposals/Protocols should be single-spaced using Calibri 12-point font and 1-inch margins. Note there is a 15-page limit exclusive of references. When uploading your Full Proposal please ensure it addresses the following sections:

Goals and Objectives

- Provide the main goal of the study and the study population (if applicable). Provide a detailed definition that is directly linked to the primary objective.

Assessment of Need for the Project

- This should reflect your study rationale. Provide a brief description of the medical/scientific question and the rationale of how this trial or study addresses the question.

Target Audience

- Describe the primary audience(s) targeted for this project. For Investigator Sponsored Clinical Trials, please specify the age, gender and other demographic information for trial population.
- Also indicate whom you believe will directly benefit from the project outcomes. Describe the overall population size as well as the size of your sample population.

Project Design and Methods

- Describe concisely the research design and methods for achieving the stated goals. For a clinical interventional study, include inclusion/exclusion criteria, treatment plan and statistical plan.

Innovation

- Explain what measures you have taken to assure that this project idea is original and does not duplicate other projects. Describe how this project builds upon existing work, pilot projects, or ongoing projects developed either by your institution or other institutions related to this project.

Evaluation and Outcomes

- Specify type and frequency of safety, efficacy, and/or outcome measures. Also indicate the method(s) used to assess measures.
- Provide a publication plan describing intended submission of abstracts to (a) congress(es) or intended submission of (a) publication(s) to peer-reviewed journals.

Anticipated Project Timeline

- Provide an anticipated timeline for your project including project start/end dates.
- An ISR grant request cannot be submitted for a study that has already commenced and was not originally supported by Pfizer.

Additional Information

- If there is any additional information you feel Pfizer should be aware of concerning the importance of this project, please summarize here.
- Early-career applicants: Letter(s) of support from mentor(s) and collaborators describing how the award will advance the applicant's career.

Organization Detail

- This information is used to assess the capability of the organizational resources available to perform the effort proposed. Identify the facilities to be used [laboratory, animal, clinical and "other"]. If appropriate, indicate their capacities, pertinent capabilities, relative proximity and extent of availability to the project.

Budget Detail

- The budget amount requested must be in Chinese Yuan (CNY).
- While estimating your budget please keep the following items in mind:
- General organizational running costs such as legal fees, insurance, heating, and lighting etc. should be included in an Institutional Overhead (if required). These costs are not specific to a grant request and therefore, should not appear as line items in budgets. However, costs that are specific to the study (e.g., some countries require insurance to be taken out on a per-study basis for clinical research) would be acceptable to be included as line items.
- The inclusion of these costs cannot cause the amount requested to exceed the budget limit set forth in the RFP.
- Pfizer does not provide funding for capital purchases (infrastructure expenses such as equipment, purchases of software or software licenses, technology or bricks and mortar). Equipment hire/leasing is acceptable and may be included in project budget.
- It should be noted that grants awarded through GMGP cannot be used to purchase Pfizer therapeutic agents (prescription or non-prescription).
- Pfizer maintains a company-wide, maximum allowed overhead rate of 28% for independent studies and projects. Please click here for details.

Required Documents

- Project Plan or Proposal
- Initial Study Protocol