



Pfizer Announces a [Research Grant RFP](#)

2022 IBD Competitive Grant Program- Using Expert Review Panel

Pfizer Global Medical Grants (GMG) 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 GMG competitive grant program involves a publicly posted Request for Proposal (RFP) that provides detail regarding a specific area of interest, sets timelines for review and approval, and uses an expert review panel (ERP) to make final grant decisions. Organizations are invited to submit an application addressing the specific gaps in research, practice or care 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.

2022 IBD Competitive Grant Program

Competitive Grant Program Eligibility

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| Geographic Scope | United States |
| Applicant Eligibility Criteria | <p>To be eligible:</p> <ul style="list-style-type: none">• The institution and principal investigator (PI) must be based in one of the eligible countries noted above.• Only organizations are eligible to receive grants, not individuals or medical practice groups.• The applicant (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.• Applicant must be affiliated with a host institution.• Both early career and experienced investigators are encouraged to apply and consideration will be given to all proposals meeting the selection criteria. |

Requirements

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| Date RFP Issued | November 3 rd , 2021 |
| Clinical Area | Inflammatory Bowel Disease |
| Area of Interest Focus | <p>The intent of the RFP is to support/fund proposals that focus on the below areas:</p> <ul style="list-style-type: none">• Real world effectiveness and safety of tofacitinib in moderately to severely active ulcerative colitis; examples would include comparative effectiveness, long term effectiveness and safety• Use of tofacitinib in certain subpopulations: examples would include extra-intestinal manifestations, pouchitis• Approaches to improving IBD management; examples would include treating to target; impact and outcomes related to healthcare disparities |

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| Expected Approximate Monetary Range of Grant Applications | <ul style="list-style-type: none">• Individual projects requesting up to \$150,000 will be considered. Pfizer anticipates awarding up to 3 grants• The amount of the grant Pfizer will be prepared to fund for any project will depend upon the expert review panel's evaluation of the proposal and costs involved, and will be stated clearly in the approval notification |
| Key Dates | <ul style="list-style-type: none">• RFP release date: November 3rd, 2021• Full Proposal Deadline: February 21st, 2022 (EXTENDED) [Please note the deadline is 23:59 Eastern Time (New York, GMT -5).]• Review of Full Proposals by ERP: April, 2022• Anticipated Full Proposal Notification Date: May, 2022 |
| How to Apply | <ul style="list-style-type: none">• Please go to www.cybergrants.com/pfizer/Research and sign in. First-time users should click "Create your password". <i>[Note: there are individual portals for each grant application type (e.g., knowledge, LOI, research full proposal, and QI full proposal). Please be sure to use the URL above.]</i> <p>Requirements for submission:</p> <ul style="list-style-type: none">• For the question "Competitive Grant?" select Yes• Select the following Competitive Grant Program Name: 2022 I&I L: IBD Competitive Grant Program• Complete all required sections of the online application. See Appendix A for additional details• If you encounter any technical difficulties with the website, please click the "Technical Questions" link at the bottom of the page |

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| Questions: | <ul style="list-style-type: none">• If you have questions regarding this RFP, please direct them in writing to the Grant Officer, Abdul Bhatti (Abdul.Bhatti@Pfizer.com), with the subject line “2022 I&L: IBD Competitive Grant Program”• Please click here to view Frequently Asked Questions regarding the Competitive Grant Program |
| Grant Agreements: | <ul style="list-style-type: none">• 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.• Pfizer has drafted the terms of these agreements to be balanced and reasonable and to further the goals of both parties. Negotiating grant agreements requires significant resources, so please ensure that your institution (including your legal department) is able and willing to abide by these terms before proceeding with submission of your application as they will need to be accepted in their entirety. |
| Review and Approval Process | <ul style="list-style-type: none">• Grant requests received in response to a specific RFP are reviewed by an expert review panel (ERP) to make final grant decisions.• The panels are comprised of professionals from the medical community with advanced degrees and expertise in particular clinical areas, or specific needs of a geographic region/learner group, or expertise in research, continuing professional development or quality improvement |
| Mechanism by which Applicants will be Notified: | <ul style="list-style-type: none">• All applicants will be notified via email by the dates noted above• Applicants may be asked for additional clarification during the review period |

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Appendix A Full Proposal/Protocol

Applications will be accepted via the online portal. Full Proposal/Protocol documents should be no longer than 10-15 pages in length (12-point font and 1-inch margins) excluding Organization Detail and References. When uploading your Full Proposal/Protocol please ensure it addresses the following:

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| Goals and Objectives | <ul style="list-style-type: none">• 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 | <ul style="list-style-type: none">• 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 | <ul style="list-style-type: none">• 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 | <ul style="list-style-type: none">• 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 | <ul style="list-style-type: none">• 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 | <ul style="list-style-type: none">• 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. All publications must follow ICH guidelines |
| Anticipated Project Timeline | <ul style="list-style-type: none">• Provide an anticipated timeline for your project including project start/end dates |
| Additional Information | <ul style="list-style-type: none">• 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 |

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| | career. |
| Organization Detail | <ul style="list-style-type: none">• 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 | <ul style="list-style-type: none">• A total amount requested is the only information needed for the LOI stage. Full Budget is not required. This amount can be adjusted at the Full Proposal stage as applicable.• The budget amount requested must be in U.S. dollars (USD).• While estimating your budget please keep the following items in mind:<ul style="list-style-type: none">○ Institutional overhead and indirect costs may be included within the grant request. Examples include human resources department costs, payroll processing and accounting costs, janitorial services, utilities, property taxes, property and liability insurance, and building maintenance as well as additional project expenses such as costs for publication, IRB / IEC review fees, software license fees, and travel. Please note: Pfizer does not provide funding for capital equipment.○ The inclusion of these costs cannot cause the amount requested to exceed the budget limit set forth in the RFP.○ It should be noted that grants awarded through GMG cannot be used to purchase therapeutic agents (prescription or non-prescription).• Pfizer maintains a company-wide, maximum allowed overhead rate of 28% for independent studies and projects |
| References | <ul style="list-style-type: none">• Bibliography of relevant references. |