



Pfizer Announces a [Research Grant RFP](#)

Pre-clinical & Translational Research in Multiple Myeloma

Competitive Grant Program - internal Pfizer review process

I. Background

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 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.

II. Eligibility

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| Geographic Scope: | United States and Canada |
| Applicant Eligibility Criteria | <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. Applicant must be affiliated with a host institution |

III. Requirements

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| Date RFP Issued | February 25, 2021 |
| Clinical Area | Oncology – Hematology – Multiple Myeloma |
| General Area of Interest for this RFP: | <p>Only pre-clinical projects with a focus on these areas of interest will be considered for Pfizer support at this time:</p> <ul style="list-style-type: none"> Correlation between soluble B-cell Maturation Antigen (sBCMA) at baseline with FLC, M-spike, deepness of overall response, MRD-negativity, and response to prior therapy Understanding the kinetics of sBCMA vs long-term responses (e.g., DoR, PFS, OS) Mechanisms of resistance (e.g., role of sBCMA in resistance/relapse; downregulation of membrane bound BCMA; T-cell exhaustion) Correlation between Soluble BCMA and BCMA expression on myeloma cells Impact of prior treatment with BCMA-targeting agents on BCMA expression and sBCMA levels Mechanistic understanding of the impact of anti-myeloma agents (e.g., IMiDs, anti-CD38, proteasome inhibitors, dexamethasone) on elranatamab-mediated T-cell activation and anti-myeloma activity Exploring in-vitro or in-vivo novel combinations with elranatamab (PF-06863135) |

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| | <p>IMPORTANT NOTES:</p> <ul style="list-style-type: none">• Studies can be in-vitro or in-vivo or use existing patient samples (e.g., blood, bone marrow aspirates, etc.)• A limited amount of the Pfizer asset elranatamab (PF-06863135) is available and can be requested through this RFP for non-clinical studies• The clinical ISR (Investigator Sponsored Research) program for elranatamab is expected to launch at a later date. |
| Expected Approximate Monetary Range of Grant Applications: | <ul style="list-style-type: none">• Individual projects requesting up to \$150,000 will be considered. The estimated total available budget related to this RFP is \$500,000.• Compound-only requests (with no funding requested) may also come through this RFP• Proposals may be drug-only or funding-only or drug plus funding |
| Key Dates: | <ul style="list-style-type: none">• RFP release date: February 25, 2021• Full Proposal Grant Application due date: May 17, 2021 Please note the deadline is 23:59 Eastern Standard Time (e.g., New York, GMT -5).• Anticipated Grant Award Notification Date: June 15, 2021• Grants will be distributed following a fully executed agreement.• Anticipated/Approximate Project Start and End Dates: July 2021 to July 2023 (study timelines may be shorter but no longer than 2 years) |
| How to Submit: | <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”.• In the application:<ul style="list-style-type: none">◦ For the question “Are you replying to a Request for Proposal as part of the Competitive Grant Program?” select Yes◦ Select the following Competitive Grant Program Name: 2021 Oncology: Pre-clinical & Translational Research in Multiple Myeloma◦ Select the following Primary Area of Interest: Oncology – Hematology – Multiple Myeloma |

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| | <ul style="list-style-type: none">• Requirements for submission: Complete all required sections of the online application and upload your project proposal (see Appendix) in the Proposal/Protocol field.• If you encounter any technical difficulties with the website, please click the “Technical Questions” link at the bottom of the page. <p>IMPORTANT: Be advised applications submitted after the due date will not be reviewed under this RFP.</p> |
| Questions: | <ul style="list-style-type: none">• If you have questions regarding this RFP, please direct them in writing to the Grant Officer, [Jacqueline Waldrop (Jacqueline.Waldrop@pfizer.com)], with the subject line “Pre-clinical & Translational Research in Multiple Myeloma.”• 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 general RFP are reviewed by Pfizer to make final grant decisions. |
| 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. |

Appendix A

General RFP Submission Requirements

Project Proposals/Protocols should be single-spaced using Calibri 12-point font and 1-inch margins. Please include 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 career. |

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| 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 |
| References | <ul style="list-style-type: none">• Bibliography of relevant references. |