



Program Announcement for the Defense Health Agency

Spinal Cord Injury Research Program Investigator-Initiated Research Award

Funding Opportunity Number: HT942526SCIRPIIRA

Pre-Application Due: August 03, 2026

Application Due: November 12, 2026

This program announcement must be read in conjunction with the General Application Instructions, version [CD26_01](#).

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Before You Begin

- **Active [SAM.gov](#), [eBRAP.org](#) and [Grants.gov](#) registrations are required for application submission.** User registration for each of these websites can take several weeks or longer. Each applicant must ensure their registrations are active and up to date prior to application preparation.
- **Read this funding opportunity announcement in the order it is written before beginning to prepare application materials.** It is the responsibility of the applicant to determine whether the proposed research meets the intent of this funding opportunity and that all parties meet eligibility requirements.
- **To support application preparation, additional resources are available** including an application process [FAQ](#), a [Guide for Intragovernmental & Intramural Applicants](#) and a [CDMRP Video Series](#) detailing the application process.

Who to Contact for Support

eBRAP Help Desk

301-682-5507
help@eBRAP.org

*Questions regarding
funding opportunity submission
requirements,
as well as technical assistance
related to pre-application or
intramural application submission.*

Grants.gov Support Center

800-518-4726
International: 1-606-545-5035
support@grants.gov

*Questions regarding
Grants.gov registration
and Workspace.*

This document uses internal links; you can go back to where you were by pressing the Alt + left arrow keys (Windows) or command + left arrow keys (Macintosh) on your keyboard.

Click  to be taken to additional guidance and instructions within the *General Application Instructions (GAI)*.

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1. Basic Information About the Funding Opportunity

Summary: The fiscal year 2026 (FY26) Spinal Cord Injury Research Program (SCIRP) Investigator-Initiated Research Award (IIRA) supports studies that have the potential to make an important contribution to spinal cord injury (SCI) research, patient care and/or quality of life. May focus on any phase of research from basic through translational science, though studies that focus solely on identifying intervention targets are discouraged.

Distinctive Features:

- This funding opportunity contains an **Early-Career Partnership Option**, which allows for two Principal Investigators (PIs). If this option is selected, at least one of the named PIs must be an early-career investigator. Only the initiating PI will submit a pre-application, but all PIs will need to submit full applications. The partnering PI's application is an abbreviated package specific to their distinct portion of the research project. If recommended for funding, each PI will be named on a separate award to the recipient organization(s). Be advised, all associated applications for a research project may be withdrawn if the initiating or partnering application is rejected or administratively withdrawn.
- ****NEW for FY26**** An investigator may be named on only **one** FY26 SCIRP IIRA application as a PI.

Funding Details: The Congressionally Directed Medical Research Programs (CDMRP) expects to allot roughly **\$4.96M** to fund approximately **6** Investigator-Initiated Research Award applications with total cost caps of **\$0.8M** for a Single PI, or **\$0.96M** for the Early-Career Partnership Option, per award. The maximum period of performance is **3** years. It is anticipated that awards made from this fiscal year 2026 (FY26) funding opportunity will be funded with FY26 funds, which will expire for use on September 30, 2032. Awards supported with FY26 funds will be made no later than September 30, 2027.

Submission and Review Dates and Times

- **Pre-Application (Preproposal) Submission Deadline:** 5:00 p.m. Eastern Time (ET), August 03, 2026
- **Invitation to Submit an Application:** September 18, 2026
- **Application Submission Deadline:** 11:59 p.m. ET, November 12, 2026
- **End of Application Verification Period:** 5:00 p.m. ET, November 16, 2026
- **Peer Review:** January 2027
- **Programmatic Review:** March 2027

Announcement Type: Initial

Funding Opportunity Number: HT942526SCIRPIIRA

Assistance Listing Number: 12.420

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2. Eligibility Information

2.1. Eligible Applicants

2.1.1. Organization

[Extramural](#) and [intramural U.S. Department of War \(DOW\)](#) organizations are eligible to apply, **including foreign and domestic organizations, for-profit and nonprofit organizations, and public or private entities.**

2.1.2. Principal Investigator

Investigators affiliated with an eligible organization are eligible to be named Principal Investigator (PI) on the application, regardless of ethnicity, nationality or citizenship status.

An investigator may be named on only **one** fiscal year 2026 (FY26) Spinal Cord Injury Research Program (SCIRP) Investigator-Initiated Research Award (IIRA) application as a PI.

Single PI Option: Independent investigators at all career levels may be named by the organization as the PI on the application.

Early-Career Partnership Option: If this option is selected, **at least one** of the named PIs (Initiating or Partnering) must be an investigator with **at least three** years' research experience (independent or non-independent) beyond a terminal degree, but **no more than seven** years within their first faculty appointment or an equivalent independent research position (excluding time spent on family medical leave). Lapses in research time or appointments as denoted in the biographical sketch should be explained in the application. **The other PI** (Initiating or Partnering) may be an independent investigator at any career level.

2.2. Cost Sharing

Cost sharing is not an eligibility requirement.

2.3. Other

Awards are made to eligible **organizations**, not to individuals. Refer to the GAI for additional [recipient qualification requirements](#).

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3. Program Description

The Defense Health Agency Contracting Activity (DHACA) is soliciting applications to this funding opportunity using delegated authority provided by United States Code, Title 10, Section 4001 (10 USC 4001). The CDMRP is the program office managing this FY26 funding opportunity as part of the Spinal Cord Injury Research Program (SCIRP). The CDMRP is located within the Defense Health Agency Research and Development (DHA R&D), which is a part of the Department of Defense, DOD, herein referred to using the secondary title Department of War, DOW. Congress initiated the SCIRP in 2009 to provide support for traumatic spinal cord injury (SCI)-related research of exceptional scientific merit that has the potential to make a significant impact on improving the health and well-being of military Service Members, Veterans, and other individuals living with SCI. Appropriations for the SCIRP from FY09 through FY25 totaled \$477.85 million (M). The FY26 appropriation is \$33M.

Mission of the SCIRP: To fund research and encourage interdisciplinary collaborations for the development and translation of more effective strategies to improve the health and well-being of Service Members, Veterans, and other individuals with spinal cord injury

Vision of the SCIRP: Advance the treatment and management of spinal cord injury and ameliorate its consequences

The FY26 SCIRP challenges the scientific community to design research that will advance the development, translation and ultimate adoption of health care solutions for people living with SCI, their families and/or care partners. The SCIRP encourages impactful and well-reasoned research across the continuum of care, at time of injury and throughout the lifespan. Additionally, to ensure alignment with current program priorities, applicants to SCIRP funding opportunities should consider the following:

- Innovative research that investigates new directions or addresses neglected issues in the field of traumatic SCI is supported, although studies focused exclusively on identifying intervention targets are discouraged.
- Mechanism-focused studies must be specific to SCI and demonstrate a clear path from increased understanding to advancing treatments.
- Applications proposing interventions must demonstrate a clinically feasible window for treatment and more than incremental improvement over existing therapies.
- Studies addressing critical questions essential for advancing research toward clinical use, such as study population selection, dosing, and safety, are allowable.
- Applications from investigators within the DOW and applications involving multidisciplinary collaborations among academia, industry, the DOW, the U.S. Department of Veterans Affairs (VA) and other federal government agencies are highly encouraged. These relationships can leverage knowledge, infrastructure and access to unique clinical populations that the collaborators bring to the research effort, ultimately advancing research that is significant to Service Members, Veterans, their Families and the American Public.

3.1. Program Priority Areas With Programmatic Goals

For FY26, the SCIRP identified near-term and long-term programmatic goals for program-supported research outcomes and products, which align to four key priority areas within the SCI field. ***Research supported by the FY26 SCIRP must address at least one of the program priority areas and undertake an associated near-term programmatic goal listed below.***

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Additionally, applicants will be asked to describe the steps the research team will take to further advance the project's outcome(s)/product(s) in accordance with the long-term programmatic goal of the same priority area.

Priority Area: Acute Injury Intervention

- **Near-Term Goal:** Develop and/or test interventions, e.g., surgical, device, pharmacologic, that protect spinal cord tissue from further damage shortly after injury and can be administered on a timeline suitable for acute trauma care in both battlefield and civilian settings.
- **Long-Term Goal:** Adoption of management/stabilization practices into the standard of care for acute spinal cord injury, leading to meaningfully improved outcomes within the prehospital, hospital, or deployed environments.

Priority Area: Secondary Health Effects

- **Near-Term Goal:** Develop and/or test interventions to prevent or manage the secondary health effects of spinal cord injury that impact injury survivors throughout their lifespan.
- **Long-Term Goal:** Adoption of interventions into the standard of care that significantly reduce the burden of secondary health effects and improve the lifelong health of SCI survivors.

Priority Area: Psychosocial Well-Being

- **Near-Term Goal:** Develop and/or test psychosocial interventions that are specifically tailored to the unique needs of individuals with SCI, their families, and/or care partners.
- **Long-Term Goal:** Adoption of interventions into the standard of care that promote psychosocial well-being for people with SCI, their families, and/or care partners.

Priority Area: Rehabilitation and Regeneration

- **Near-Term Goal:** Develop and/or test rehabilitation, regenerative, or neuroplastic interventions for individuals with spinal cord injury that can be implemented at any stage of life post-injury.
- **Long-Term Goal:** Adoption of validated rehabilitation and regenerative therapies into the standard of care that meaningfully improve function, independence, and quality of life after SCI.

3.2. Award History

The SCIRP Investigator-Initiated Research Award (IIRA) mechanism was first offered in FY09. Since then, 981 Investigator-Initiated Research Award applications were received, and 161 were recommended for funding.

3.3. Intent of the Investigator-Initiated Research Award

The SCIRP IIRA is intended to support studies that have the potential to make an important contribution to SCI research, patient care, and/or quality of life.

IIRA applications may focus on any phase of research from basic through translational, though studies focused exclusively on target identification are discouraged. Permitted research includes preclinical studies in animal models, research with human subjects or human anatomical substances, as well as ancillary studies associated with an existing clinical trial.

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3.3.1. Key Elements for the IIRA

- **Impact:** Applications must articulate how the proposed research will advance at least one of the FY26 SCIRP priority areas and help achieve its associated near-term programmatic goal identified in [Section 3.1](#)
- **Community-Informed Research:** Research funded by the FY26 SCIRP IIRA must be responsive to the needs of people with SCI, their families and/or their care partners. To this end, applicants are encouraged to leverage patient perspectives research to inform the proposed research question or study design.
- **Early-Career Partnership Option:** To promote enhanced research capacity within the SCI field, the FY26 IIRA includes an option specifically for partnership with or among early-career investigators. The Partnership Option is structured to accommodate two PIs who will work together on a single research project. One PI will be identified as the Initiating PI and will be responsible for the majority of the administrative tasks associated with application submission. The other PI will be identified as a Partnering PI. At least one of the PIs (Initiating or Partnering) must be an [early-career investigator](#). Both PIs should contribute significantly to the development and execution of the proposed research project. If recommended for funding, each PI will be named on a separate award to the recipient organization(s). Each award will be subject to separate reporting, regulatory, and administrative requirements. For individual submission requirements for the Initiating and Partnering PIs, refer to [Section 5.3, Submission Instructions](#).

3.3.2. Other Important Considerations for the IIRA

Allowable Research: [Clinical research](#) is allowed within this funding opportunity, but [clinical trials](#) are disallowed. Applications requesting funding for a clinical trial or combination of preclinical research and clinical trial may be more suited to one of the other [FY26 SCIRP funding opportunities](#) being offered.

In accordance with the National Defense Authorization Act for Fiscal Year 2026, Section 732, CDMRP does not support the conduct of painful research (U.S. Department of Agriculture pain category D or E) involving domestic cats or dogs, except for studies relating to military or service animals.

If the proposed research relies on access to unique resources or databases, the application must describe the access at the time of submission and include a plan for maintaining access as needed throughout the proposed research.

All projects should adhere to a core set of standards for rigorous study design and reporting to maximize the reproducibility and translational potential of clinical and preclinical research, such as those described in the [STROBE](#) and [ARRIVE 2.0](#) guidelines.

Use of Common Data Elements (CDEs): Use of the National Institute of Neurological Disorders and Stroke [SCI CDEs](#) is strongly encouraged for all human subjects research.

Relevance to Military Health: To help elucidate the realities of treating and managing SCIs while in a deployed environment, a resource document is now available that outlines [Spinal Cord Injury Management within the Military Health System \(MHS\)](#). Applicants are encouraged to read and consider this document before preparing applications. Furthermore, applicants are encouraged to integrate and/or align their research projects with DOW and/or VA research laboratories and programs. Collaboration with the DOW and/or VA is also encouraged. A list of websites that may be useful in identifying additional information about ongoing DOW and VA

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areas of research interest or potential opportunities for collaboration can be found in [Appendix 10](#) of the GAI.

3.4. Funding Instrument

The funding instrument for awards made under the program announcement will be grants (31 USC 6304).

3.5. Funding Details

Period of Performance: The maximum period of performance is **3** years.

3.5.1. Application Submissions With a Single PI

Cost Cap: The application's total costs budgeted for the entire period of performance should not exceed **\$0.8M**.

3.5.2. Application Submissions With the Early-Career Partnership Option

Cost Cap: The combined total costs budgeted for the entire period of performance in the applications of the Initiating PI and the Partnering PI should not exceed **\$0.96M**.

A separate award will be made to each PI's organization.

The PIs are expected to be partners in the research, and direct cost funding should be divided accordingly unless otherwise warranted and clearly justified.

Any application that requests the higher level of funding and that does not an eligible [early career](#) PI will have its budget reduced as appropriate.

3.5.3. For Both Options Within This Award Mechanism

If indirect cost rates have been negotiated, indirect costs are to be budgeted in accordance with the organization's negotiated rate. Collaborating organizations should budget associated indirect costs in accordance with each organization's negotiated rate.

All direct and indirect costs of any subaward or contract must be included in the direct costs of the primary award.

The applicant may request the entire maximum funding amount for a project that may have a period of performance less than the maximum **3** years.

The appropriateness of the budget for the proposed research will be assessed during peer review.

Direct Cost Restrictions: For this award mechanism, direct costs:

May be requested for (not all-inclusive):

- Data and research resource sharing costs.
- Travel in support of multi-institutional collaborations.
- Costs for one investigator, per award, to travel to one scientific/technical meeting per year. The intent of travel to scientific/technical meetings should be to present project information or disseminate project results from the FY26 SCIRP IIRA.

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Must not be requested for:

- Clinical trial costs.
- Costs for travel to scientific/technical meeting(s) beyond the limits stated above.

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4. Application Contents and Format

4.1. Application Overview

Application submission is a two-step process requiring both a **pre-application** submitted via the Electronic Biomedical Research Application Portal ([eBRAP](#)) and a **full application** submitted through eBRAP or Grants.gov. Depending on the submission portal, certain aspects of the application will differ.

Intramural DOW organizations submitting a full application should follow instructions for submission through eBRAP.



Extramural organizations submitting a full application must follow instructions for submission through Grants.gov.



4.2. Pre-Application Components

The PI, or Initiating PI if selecting the **Early-Career Partnership Option**, must submit the following pre-application components.

Upload documents as individual PDF files unless otherwise noted. Files must comply with the [formatting guidelines](#) listed in the GAI.

- **Preproposal Narrative (one-page limit):** The Preproposal Narrative page limit applies to text and non-text elements (e.g., figures, tables, graphs, photographs, diagrams, chemical structures, drawings) used to describe the project. Inclusion of URLs that provide additional information to expand the Preproposal Narrative and could confer an unfair competitive advantage is prohibited and may result in administrative withdrawal of the pre-application.

The Preproposal Narrative should include the following:

- **Background/Rationale:** State the scientific rationale on which the proposed research project is based. Clearly demonstrate that there is sufficient rationale, background data, and readiness to support the study. ***Indicate how the research question or study design is informed by the lived-experience community and/or patient perspectives research.***
- **Specific Aims and Study Design:** Concisely state the project's specific aims and describe the scientific approach to address them. Include the animal model proposed, if applicable, along with its relevance to human SCI.
- **Impact:** Describe the potential impact of the proposed research on at least one of the FY26 priority areas. Indicate how the proposed research will help to achieve a near-term programmatic goal identified in [Section 3.1](#).
- **Relevance to Military Health:** Describe whether/how the proposed research project is applicable to spinal cord-injured military Service Members, Veterans, and/or their Family members and care partners. Applicants are encouraged to review the [Relevance to Military Health](#) section of this program announcement prior to preparing this section.

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- **Pre-Application Supporting Documentation:** The items to be included as supporting documentation for the pre-application *must be uploaded as individual files* and are limited to the following:
 - **References Cited (one-page limit):** List the references cited (including URLs if available) in the Preproposal Narrative using a standard reference format that includes the full citation (i.e., author[s], year published, reference title, and reference source, including volume, chapter, page numbers, and publisher, as appropriate).
 - **List of Abbreviations, Acronyms and Symbols:** Provide a list of abbreviations, acronyms, and symbols used in the Preproposal Narrative.

4.3. Full Application Components

Applicants must receive an invitation to submit a full application. Uninvited full application submissions will be rejected.

Early-Career Partnership Option: The CDMRP requires separate full application package submissions for the Initiating PI and Partnering PI, even if the PIs are located within the same organization. The application submission process for the Partnering PI uses an [abbreviated full application package](#).

4.3.1. Full Application Components for the PI or Initiating PI

Each application submission must include the completed full application package for this program announcement. See [Appendix 1](#) for a checklist of the full application components.

(a) SF424 Research & Related Application for Federal Assistance Form (*Grants.gov submissions only*)



IMPORTANT: When completing the SF424 R&R, enter the **eBRAP log number** assigned during pre-application submission into **Block 4a – Federal Identifier**.

(b) Attachments:

Each attachment of the full application components must be uploaded as an individual file in the format specified and in accordance with the [formatting guidelines](#) in the GAI.

- **Attachment 1: Project Narrative (10-page limit): Upload as “ProjectNarrative.pdf”.**



Describe the proposed project in detail using the outline below.

- **Background:** Present the ideas and scientific reasoning behind the proposed research project. Provide a review and analysis of relevant literature and completed/ongoing studies, including community/patient perspectives research as appropriate. If applicable, discuss how the proposed project builds off previous research efforts. This section should establish the relevance of the study and explain the applicability of the proposed findings to the SCI community and at least one of the [FY26 SCIRP Program Priorities](#) and associated near-term goal.
- **Objectives/Specific Aims/Hypotheses:** Describe the purpose of the proposed study with detailed objectives. State the hypothesis/research question to be tested and detail the specific aims that will address the hypothesis/research question. Only describe aims that the SCIRP award would fund.

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- **Study Design and Feasibility:** Describe the research strategy, methods, and analyses, including controls, in sufficient detail for evaluation of their appropriateness and feasibility. Describe how the study is designed to achieve the project aims. Consult appropriate [guidelines](#) to ensure relevant aspects of rigorous and reproducible research are adequately planned for and, ultimately, reported. Address potential problem areas, and present alternative methods and approaches where appropriate. Indicate whether and how the study design is informed by the lived-experience community and/or patient perspectives research.

As appropriate, describe the statistical model and data analysis plan with respect to the study objectives. If applicable, include power analysis calculations. Further information describing the strategy for how sex will be considered as a biological variable will be requested in [Attachment 2](#).

If applicable, briefly describe the use of animal model(s) within the study. Include a rationale for the choice of animal model and the model's relevance to human SCI. Full details will be required in the Animal Research Plan ([Attachment 8](#)).

If human subjects, human anatomical samples or data will be used, include a detailed plan for the recruitment of subjects or the acquisition of samples or data, including evidence that the research team has access to subjects, samples and/or data. Describe the strategy for the inclusion of women and minorities appropriate to the objectives of the study, including a description of the composition of the proposed study population in terms of sex, racial, and ethnic group, and an accompanying rationale for the selection of subjects. Anticipated enrollment table(s) with the proposed enrollment distributed on the basis of sex, race, and ethnicity should be provided as part of the application's Supporting Documentation ([Attachment 2](#)). ***This award may not be used to conduct [clinical trials](#).***

If applicable, describe the SCI CDEs to be collected.

- **Research Team Composition:** Describe the composition of the research team in enough detail to demonstrate that the study team has appropriate expertise to accomplish the goals of the proposed research.
- **Attachment 2: Supporting Documentation: Combine and upload as a single file named "Support.pdf".** 

There are no page limits for these components unless otherwise noted. Include only components described below; inclusion of items not requested or viewed as an extension of the Project Narrative will result in the removal of those items or may result in administrative withdrawal of the application.

- **References Cited:** List the references cited in the Project Narrative using a standard reference format (include URLs, if available).
- **List of Abbreviations, Acronyms and Symbols:** Provide a list of abbreviations, acronyms and symbols.
- **Facilities, Existing Equipment and Other Resources:** Describe the facilities and equipment available for performance of the proposed project; include any additional facilities or equipment proposed for acquisition at no cost to the award. Indicate whether government-furnished facilities or equipment are proposed for use. If so, reference the original or present government award under which the facilities or equipment items are now accountable. There is not a standardized form for this information.

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- **Publications and/or Patents:** Include a list of relevant publication URLs and/or patent abstracts. If articles are not publicly available, then copies of up to five published manuscripts may be included in Attachment 2. Extra items will not be reviewed.
- **Letters of Support (two-page limit per letter is recommended):** Provide individual letters signed by collaborating individuals and/or organizational officials demonstrating that the PI has the support and resources necessary for the proposed work. Letters from the PI's Department Chair, or appropriate organization official, should also confirm that the PI(s) meet [eligibility criteria](#). If applicable, provide a letter of support, signed by the lowest-ranking person with approval authority, confirming participation of intramural DOW collaborator(s) and/or access to military populations, databases or DOW resources. If applicable, provide a letter of support signed by the VA Facility Director(s), or an individual designated by the VA Facility Director(s), confirming access to VA patients, resources and/or VA research space.
- **Sex as a Biological Variable Strategy (two-page limit is recommended):** Describe the strategy for how sex will be considered as a biological variable. This strategy should include a brief discussion of what is currently known regarding sex differences in the applicable research area. Clearly articulate how sex as a biological variable will be factored into the data analysis plan and how data will be collected and disaggregated by sex. If needed, provide a strong rationale for proposing a single-sex study, based on justification from scientific literature, preliminary data or other relevant considerations. Refer to the [CDMRP Directive on Sex as a Biological Variable in Research](#) for additional information.
- **Research Sharing Plan:** Describe the type of data or research resources (e.g., bio-specimen, analysis tool/software, training material) to be made publicly available as a result of the proposed work. Describe the mechanism (e.g., direct sharing, repository, mixed mode) by which data and resources generated during the period of performance will be shared with the research community and other affected communities, including clinical research participants. Include the name of the repository(ies) where scientific data and resources arising from the proposed study will be archived, if applicable. Identify and provide the rationale for any data or resources that will not be shared (e.g. for intellectual property, feasibility, cost, or other considerations). The plan should also protect participant privacy, confidential and proprietary data, and performer/third-party intellectual property. Provide a milestone plan for disseminating data/results including when data and resources will be made available to other users. In cases where the study participant could potentially derive medical or other benefit from the information, explain whether the results of screening and/or study participation will be shared with the participant or their primary care provider, including results from any screening or diagnostic tests performed as part of the study.

Do not submit a copy of the National Institutes of Health Data Management and Sharing Plan or duplicate the Data Management Plan, which will be requested only after a recommendation for funding is made.

Refer to the [CDMRP Directive on Sharing Data and Research Resources](#) for more information about the CDMRP's expectations for making data and research resources publicly available.

- **Inclusion Enrollment Plan (only required if [clinical research](#) is proposed):** Provide an anticipated enrollment table(s) for the inclusion of women and minorities

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using the [Public Health Service \(PHS\) Inclusion Enrollment Report](#), a three-page fillable PDF form that can be downloaded from eBRAP. The enrollment table(s) should be appropriate to the objectives of the study with the proposed enrollment distributed on the basis of sex, race and ethnicity.

- **Attachment 3: Technical Abstract (one-page limit): Upload as “TechAbs.pdf”.** 

Write the technical abstract using the outline below. Clarity and completeness within the space limits are highly important.

- **Background:** Present the scientific rationale behind the proposed research project.
- **Hypothesis/Objective(s):** State the hypothesis to be tested and/or objective(s) to be reached.
- **Specific Aims:** State the specific aims of the study.
- **Study Design:** Briefly describe the study design, including appropriate controls.
- **Impact:** Briefly describe the impact of this study on one or more FY26 SCIRP priority areas. Indicate how the proposed research will help to achieve a near-term programmatic goal identified in [Section 3.1](#).
- **Military Relevance:** Describe how the study is relevant to military health.

- **Attachment 4: Lay Abstract (one-page limit): Upload as “LayAbs.pdf”.** 

The lay abstract should address the points outlined below *in a manner that is readily understood by readers without a background in science or medicine*. Avoid overuse of scientific jargon, acronyms and abbreviations. *Do not duplicate the technical abstract.*

- Summarize the objectives and rationale for the proposed research.
- Describe the intervention(s), if applicable.
- What population will the research help, and how will it help them?
- What are the potential applications, benefits, and risks of the anticipated outcomes?
- If the research is too basic for immediate clinical applicability, describe the interim outcomes.
- What is the projected time it may take to achieve a person-related outcome?
- Describe how the proposed study will directly address an FY26 SCIRP priority area and contribute to the associated near-term programmatic goal identified in [Section 3.1](#).
- What is the potential benefit of the proposed study and the anticipated outcome to Service Members, Veterans and/or their Families?

- **Attachment 5: Statement of Work (three-page limit): Upload as “SOW.pdf”.** 
Refer to eBRAP for the [Suggested SOW Format](#).

For guidance on preparing the SOW, refer to the [Example: Assembling a Generic Statement of Work](#). Include milestones for data or research resource(s) sharing.

Include the name(s) of the key personnel and contact information for each study site/subaward site. The contributions of the key personnel, including the PI or Initiating

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PI, Partnering PI (if applicable), and SCI Lived-Experience Consultants or community partners, should be noted for each task.

Early-Career Partnership Option: Each PI must submit an identical copy of a jointly created SOW.

- **Attachment 6: Impact Statement (one-page limit): Upload as “Impact.pdf”.** The impact statement summarizes the potential short- and long-term impact of the proposed research for the SCI lived-experience community and for the broader SCI research field. The statement should address the points outlined below written ***in a manner that will be readily understood by readers without a background in science or medicine.***
 - Detail the anticipated research outcome(s) that will be directly attributed to the results of the proposed study and describe the anticipated benefits of these outcomes for individuals with SCI and the broader SCI research field. Explain how the proposed project will advance at least one of the FY26 SCIRP priority areas and help achieve its associated near-term programmatic goal, as identified in [Section 3.1](#).
 - To demonstrate how the study is grounded in the needs of the SCI community, describe how the lived-experience community and/or patient perspectives research informed the research question or study design.
 - Assess the likelihood that a successful outcome of the proposed research project will lead to practical application in individuals living with SCI. If applicable, state whether/how the proposed project builds off previous research efforts to advance a product, outcome or finding closer to clinical utility.
 - If applicable, describe how the anticipated outcomes of the proposed study will make an impact in understanding health differences between sexes.
 - Describe whether/how the proposed research project is applicable to the unique health care needs and quality of life concerns of spinal cord-injured military Service Members, Veterans, and/or their Family members and care partners. Applicants are encouraged to review the [Relevance to Military Health](#) section of this program announcement prior to preparing this section.
- **Attachment 7: Long-Term Goal (three-page limit): Upload as “Longterm.pdf”.** The purpose of this attachment is to create a high-level roadmap detailing the steps needed to eventually translate outcomes of SCIRP-funded research into the standard of care, consistent with the [long-term programmatic goals](#) of the FY26 SCIRP. The long-term goal attachment should include the points listed below.
 - **Near-Term Action:** Outline the immediate next steps to be taken by the research team after successful completion of the proposed research project, including plans for dissemination of results, further outcome validation if applicable and engagement with end-users (e.g., patients, clinicians).
 - **Intermediate Objectives:** Discuss the unresolved questions that must be answered for the outcome to be adopted within the standard of care, considering the following as a starting place:
 - What additional research is necessary?
 - What is the likely regulatory pathway?
 - What is the commercial potential of the outcome?

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- What is the intellectual property strategy? If the intellectual property rights are not owned by the study team, how will the final product become accessible to the SCI community?
- **Long-Term Strategy:** Describe the long-term implementation and adoption strategy for the research outcome, outlining key milestones on the path to adoption.

Identify the key collaborations (e.g., clinical partners, commercial entities, patient groups, professional societies) required to execute the strategy. Be specific about the types of partners needed, and name them if possible.

Concisely explain how successfully executing this strategy will lead to the integration of the research outcome into the standard of care and directly fulfill the specific [long-term programmatic goal](#) for the relevant FY26 SCIRP priority area.
- **Attachment 8: Animal Research Plan (three-page limit): Upload as “AnimalResPlan.pdf”. (Applications proposing animal studies only.)** The purpose of this attachment is to provide additional details regarding the proposed animal research beyond what was included within the project narrative. Consult the [ARRIVE guidelines 2.0](#) (Animal Research: Reporting *In Vivo* Experiments) to ensure relevant aspects of rigorous animal research are adequately planned for. The Animal Research Plan may not be an exact replica of the protocol(s) submitted to the Institutional Animal Care and Use Committee (IACUC). The Animal Research Plan should address the following points to achieve reproducible and rigorous results for each proposed animal study:
 - Describe the research objective(s) of the animal study. Explain how and why the animal species, strain, and model(s) being used can address the scientific objectives and, where appropriate, the study’s relevance to human biology. **Be specific as to why an animal model is necessary to address the study aims, why the specific animal SCI model was chosen over other models, including any relevance of the model specific to military health, and how it is optimal for addressing the study aims. The model’s relevance to human SCI should also be detailed.**
 - Summarize the procedures to be conducted and the outcome measures to be collected. Describe how the study will be controlled.
 - Describe the randomization and blinding procedures for the study, and any other measures to be taken to minimize the effects of subjective bias during animal treatment and assessment of results. If randomization and/or blinding will not be utilized, provide justification.
 - Provide a sample size estimate for each study arm and the method by which it was derived, including power analysis calculations.
 - Describe how data will be handled, including rules for stopping data collection, criteria for inclusion and exclusion of data, how outliers will be defined and handled, statistical methods for data analysis, and identification of the primary endpoint(s).
- **Attachment 9: Partnership Statement (one-page limit): Upload as “Partnership.pdf”. (Early-Career Partnership Option only.)** Describe the experience of the Initiating and Partnering PI and indicate how the award will help to enhance research capacity within the SCI field. Describe the contribution and time commitment of each PI toward the proposed research project. Describe how the partners’ combined experience will better address the research question and explain why the work should be done together rather than through separate efforts.

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- **Attachment 10: Representations (*Grants.gov submissions only*): Upload as “RequiredReps.pdf”.** All extramural applicants must complete and submit the [Required Representations](#) document available on eBRAP. 
- **Attachment 11: Suggested Intragovernmental/Intramural Budget Form (*if applicable*): Upload as “IGBudget.pdf”.** If an [intramural DOW organization](#) will be a collaborator in the performance of the project, complete a separate budget for that organization using the [Suggested Intragovernmental/Intramural Budget](#) form available on eBRAP. 

(c) Additional Application Materials:

The following are additional forms for application submission. Follow the instructions specific to the submission portal, as found within the GAI.



Grants.gov



eBRAP.org

i. Research & Related Senior/Key Person Profile (Expanded)

- **Biographical Sketch**
- **Current/Pending Support**

Intragovernmental applicants must include their internally supported research and development programs.

ii. Research & Related Budget

Initiating and Partnering PIs must have a separate budget and justification specific to their distinct portions of the effort that the applicant organization will submit as separate Grants.gov or eBRAP application packages. The Initiating PI should not include budget information for the Partnering PI even if they are located within the same organization. Refer to [Section 3.5, Funding Details](#), for detailed budget information.

iii. Project/Performance Site Location(s)

iv. Research & Related Subaward Budget Attachment(s) (*if applicable, Grants.gov submissions only*)

4.3.2. Full Application Components for the Partnering PI

Refer to the equivalent attachment [above](#) for details specific to each of the following application components. See [Appendix 1](#) for a checklist of the full application components required for the Partnering PI.

(a) [SF424 Research & Related Application for Federal Assistance Form](#) (*Grants.gov Submissions Only*): 

(b) Attachments:

- **Attachment 5: Statement of Work (three-page limit): Upload as “SOW.pdf”.** Each PI must submit an identical copy of a jointly created SOW.
- **Attachment 10: Representations (*Grants.gov submissions only*): Upload as “RequiredReps.pdf”.**

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- [Attachment 11](#): Suggested Intragovernmental/Intramural Budget Form: Upload as “IGBudget.pdf”.

(c) [Additional Application Materials](#):

The following are additional application materials for application submission. Follow the instructions specific to the submission portal found within the GAI.



Grants.gov



eBRAP.org

i. Research & Related Senior/Key Person Profile (Expanded)

- **Biographical Sketch**
- **Current/Pending Support**

Intragovernmental applicants must include their internally supported research and development programs.

ii. Research & Related Budget

Initiating and Partnering PIs must have a separate budget and justification specific to their distinct portions of the effort that the applicant organization will submit as separate Grants.gov or eBRAP application packages. The Partnering PI should not include budget information for the Initiating PI even if they are located within the same organization. Refer to [Section 3.5, Funding Details](#), for detailed budget information.

iii. Project/Performance Site Location(s) Form

iv. Research & Related Subaward Budget Attachment(s) Form *(if applicable, Grants.gov submissions only)*

4.4. Other Application Elements

If recommended for funding, a data management plan compliant with Section 3.c, Enclosure 3, [DoD Instructions 3200.12](#) will be requested.



The government reserves the right to request a revised budget, budget justification and/or additional information for applications recommended for funding.

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5. Submission Requirements

5.1. Location of Application Package

Download the application package components for HT942526SCIRPIIRA from [Grants.gov](#) or [eBRAP](#), depending on which submission portal will be used.

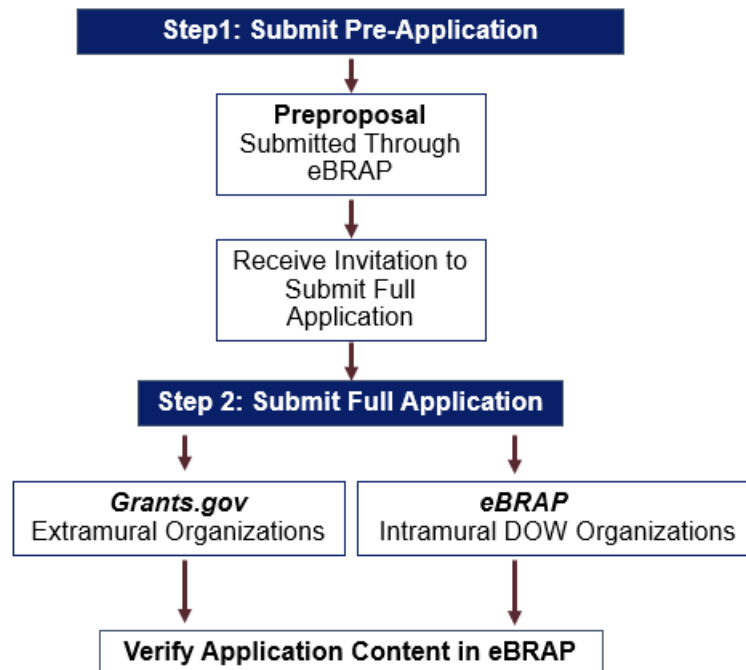
5.2. Unique Entity Identifier and System for Award Management

The applicant organization must be registered as an entity in the System for Award Management (SAM), [SAM.gov](#), and receive confirmation of an “Active” status before submitting an application through Grants.gov. Organizations must include the unique entity identifier (UEI) generated by the SAM in applications to this funding opportunity and maintain an active registration in the SAM at all times during which it has an active Federal award or an application under consideration. i

5.3. Submission Instructions

The CDMRP uses two portal systems to accept pre- and full application submissions. The workflow below shows which portal system to use for pre- and full application submissions, respectively.

Application Submission Workflow



5.3.1. Pre-Application Submission

All pre-application components must be submitted by the PI or Initiating PI through [eBRAP](#), including the submission of contact information for the Partnering PI if selecting the Early-Career Partnership Option. i

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During the pre-application process, eBRAP assigns each submission a unique log number. This unique log number is required during [the full application submission process](#). The eBRAP log number, application title and all information for the PI, Business Official(s), performing organization, and contracting organization must be consistent throughout the entire pre-application and full application submission process. Inconsistencies may delay application processing and limit or negate the ability to view, modify and verify the application in eBRAP. Contact the [eBRAP Help Desk](#) if any changes need to be made.

Early-Career Partnership Option: After the Initiating PI confirms submission of the pre-application, the Partnering PI will be notified of the pre-application submission via an email from eBRAP. ***The Partnering PI must follow the instructions provided in the email to associate the partnering pre-application with their eBRAP account.*** If not previously registered, the Partnering PI must register in eBRAP.


Partnering PIs should not initiate a new pre-application based on the same research project submitted by the Initiating PI. Partnering PIs are urged to associate the partnering pre-application with their eBRAP account as soon as possible. If this is not completed by the full application deadline:

- Any intramural Partnering PI will not be able to submit their full application package components to eBRAP.
- The Partnering PI will not be able to view and modify their full application during the verification period in eBRAP.

When starting the pre-application, PIs should select a Mechanism Option appropriate to their pre-application:


Application Includes:	Select Mechanism Option:
Single PI	No Option
Early-Career PI and other PI	IIRA With Early-Career Partnership Option

5.3.2. Full Application Submission

Grants.gov Submissions: Full applications from extramural organizations *must* be submitted through the Grants.gov Workspace. 

eBRAP Submissions: Only [intramural DOW organizations](#) may submit full applications through eBRAP. 

5.3.3. Applicant Verification of Full Application Submission in eBRAP

Independent of the submission portal, once the full application is submitted, it is transmitted to and processed in eBRAP; the transmission to eBRAP may take up to 48 hours. At this stage, the PI and organizational representatives will receive an email from eBRAP instructing them to log in to eBRAP to review, modify and verify the full application submission. ***The Project Narrative and Research & Related Budget Form cannot be changed after the application submission deadline.*** Other application components, including subaward budget(s) and subaward budget justification(s), may be changed until the [application verification period](#) ends. The full application cannot be modified once the application verification period ends. 

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5.4. Submission Dates and Times

The pre-application and full application submission process should be started early to avoid missing deadlines. Regardless of submission portal used, all pre- and full application components must be submitted by the deadlines stipulated in this program announcement. There are no grace periods for deadlines; failure to meet submission deadlines will result in application rejection. ***The DHACA cannot make allowances/exceptions for submission problems encountered by the applicant.***

Submission dates and times are specified in [Section 1, Basic Information](#).

5.5. Intergovernmental Review

Not applicable for this funding opportunity.

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6. Application Review Information

6.1. Application Compliance Review

Submitting applications that propose essentially the same research project to different funding opportunities within the same program and fiscal year is prohibited and will result in administrative withdrawal of the duplicative application(s).

While it is allowable to propose similar research projects to different programs within the CDMRP or to other organizations, duplication of funding or accepting funding from more than one source for the same research is prohibited. See the [CDMRP's Directive on Research Duplication](#).

Including classified research data within the application and/or proposing research that may produce classified outcomes or outcomes deemed sensitive to national security concerns, may result in application withdrawal.



Members of the FY26 SCIRP Programmatic Panel must not be involved in any pre-application or full application including, but not limited to, concept design, application development, budget preparation and the development of any supporting documentation, including personal letters of support/recommendation for the research and/or PI. Programmatic panel members **may** provide [letters](#) to confirm [PI eligibility](#) and access to laboratory space, equipment and other resources necessary for the project if that is part of their regular roles and responsibilities (e.g., as Department Chair). ***A list of the [FY26 SCIRP Programmatic Panel members](#) can be found on the CDMRP website.***

Additional restrictions and associated administrative responses are outlined in [Section 9.2, Administrative Actions](#).

6.2. Review Criteria

6.2.1. Pre-Application Screening Criteria

To determine the merits of the pre-application and the relevance to the mission the SCIRP, pre-applications will be screened based on the following criteria:

- **Funding Mechanism:** How well the study adheres to the intent of the funding mechanism.
- **Background/Rationale:** How well the background and scientific rationale demonstrate sufficient evidence to support the proposed research project. To what extent the research question or study design is informed by the lived-experience community and/or patient perspectives research.
- **Specific Aims and Study Design:** How well the specific aims are stated and addressed in the outlined research project.
- **Impact:** How well the proposed research project addresses one or more [FY26 SCIRP Priority Areas](#) and will make important contributions toward achieving a near-term programmatic goal.
- **Relevance to Military Health:** To what extent the proposed research project will directly or indirectly benefit spinal cord-injured military Service Members, Veterans, and/or their Family members and care partners.

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6.2.2. Peer Review Criteria

To determine technical merit, all applications will be evaluated individually according to the following **scored criteria**, listed in decreasing order of importance:

- **Study Design and Feasibility**

- How well the rationale for the proposed research is supported by the review and analysis of the relevant literature and completed/ongoing studies.
- To what degree the research question or study design is informed by the lived-experience community and/or patient perspectives research.
- How well the specific aims/hypotheses/research question, study design, experimental methods, data collection procedures, evaluations, and data analysis plan are designed to address the objective and purpose of the study.
- To what extent the proposed research project is feasible as described.
 - If applicable, whether the application includes sufficient evidence to support successful recruitment of and access to human subjects, data, and samples, and whether the strategy for the inclusion of women and minorities and the distribution of proposed enrollment are appropriate for the proposed research
- How well the application acknowledges potential problems and addresses alternative approaches.
- How well the study is designed to achieve reproducible and rigorous results, including the choice of model and the endpoints/outcomes.
 - If applicable, how well the chosen animal model of SCI simulates the human condition and is justified is over other models.
- Whether the strategy for considering sex as a biological variable is appropriate to the objectives of the study or whether the justification for a single-sex study is sufficiently strong.
- If applicable, whether the application describes SCI CDEs to be collected.

- **Scientific Impact**

- To what extent a successful outcome of the proposed research project will make important contributions toward advancing at least one of the FY26 SCIRP priority areas and help achieve its associated near-term programmatic goal, as identified in [Section 3.1](#).
- If applicable, to what extent the proposed project builds off of previous research efforts to advance a product, outcome, or finding closer to clinical utility.

- **Patient Impact**

- To what extent the proposed research addresses a critical need for the SCI community, specific to at least one [FY26 SCIRP Priority Area](#).
- How likely a successful outcome of the proposed research project will lead to a practical application in individuals living with SCI.
- If applicable, to what extent the anticipated outcomes of the proposed study will make an impact in understanding health differences between sexes.

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- To what degree the proposed research project is applicable to the unique health care needs and quality of life concerns of injured military Service Members, Veterans, and/or their Family members and care partners.
- **Long-Term Goal**
 - To what extent the application provides a credible and comprehensive strategy to eventually translate outcomes of the research into the standard of care, consistent with the [long-term programmatic goals](#) of the FY26 SCIRP.
 - Whether immediate next steps to be taken by the research team upon successful completion of the proposed research project are articulated and appropriate.
 - To what degree the discussion of unresolved questions that must be answered for the research outcome to be adopted within the standard of care demonstrates a realistic understanding of the primary barriers to adoption.
 - How well the implementation and adoption strategy details a clear, logical, and feasible roadmap with high potential to translate the research outcome into a product or clinical practice within the standard of care; and to what extent key collaborations are identified and appropriate to the plan.
- **Personnel**
 - To what extent the composition of the study team is appropriate to accomplish the proposed research project.
 - To what degree the levels of effort by the PI, Partnering PI (if applicable), and other key personnel are appropriate to ensure successful conduct of the proposed work.
 - **Early-Career Partnership Option:** How the partners' combined expertise will better address the research question and to what extent the award will help to enhance research capacity within the SCI field.

In addition, the following criteria will also contribute to the overall evaluation of the application, but will not be individually scored and are therefore termed **unscored criteria**:

- **Research Sharing Plan**
 - To what extent the plan for sharing of project data and research resources is appropriate and reasonable and includes dissemination to affected communities, study participants and/or the scientific community. If applicable, which specific repository(ies) were designated for storing project data and research resources?
- **Budget**
 - Whether the budget is appropriate for the proposed research.
- **Environment**
 - To what degree the scientific environment, clinical setting and the accessibility of institutional resources support the clinical trial at each participating center or institution (including collaborative arrangements).
 - Whether there is evidence for appropriate institutional commitment from each participating institution.
- **Application Presentation**
 - To what extent the writing, clarity and presentation of the application components influence the review.

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6.2.3. Programmatic Review

To make funding recommendations and select the application(s) that, individually or collectively, will best achieve the program objectives, the following criteria are used by programmatic reviewers:

- Ratings and evaluations of peer reviewers
- Relevance to the priorities of the FY26 SCIRP, as evidenced by the following:
 - Adherence to the intent of the funding opportunity
 - Relative impact
 - Program portfolio composition
 - Relevance to military health

6.3. Application Review and Selection Process

6.3.1. Pre-Application

Following the pre-application screening, PIs and Initiating PIs will be notified as to whether they are invited to submit full applications. The estimated date when PIs can expect to receive notification of an invitation to submit a full application is indicated in [Section 1, Basic Information About the Funding Opportunity](#). No feedback (e.g., a critique of the pre-application's strengths and weaknesses) is provided at this stage. Because the invitation to submit a full application is based on the contents of the pre-application, investigators should not change the title or research objectives after the pre-application is submitted.

6.3.2. Full Application

All applications are evaluated by scientists, clinicians and consumers in a two-tier review process. The first tier is **peer review**, the evaluation of applications against established criteria to determine technical merit, where each application is assessed for its own merit, independent of other applications. The second tier is **programmatic review**, a comparison-based process in which applications with high scientific and technical merit are further evaluated for programmatic relevance. Final recommendations for funding are subject to review and approval by a designated official. ***The highest-scoring applications from the first tier of review are not automatically recommended for funding. Funding recommendations depend on various factors as described in [Section 6.2.3, Programmatic Review](#).*** Additional information about the two-tier process used by the CDMRP can be found on the [CDMRP website](#).

Funding of applications received is contingent upon the availability of federal funds for this program, the number of applications received, the quality and merit of the applications as evaluated by peer and programmatic review, and the requirements of the government. Funds to be obligated on any award resulting from this funding opportunity will be available for use for a [limited time period](#) based on the fiscal year of the funds.

6.4. Risk, Integrity and Performance Information

Prior to making an assistance agreement award where the federal share is expected to exceed the simplified acquisition threshold, as defined in the Code of Federal Regulations, Title 2, Part 200.1 (2 CFR 200.1), over the period of performance, the federal awarding agency is required to review and consider any information about the applicant that is available in the SAM.

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An applicant organization may review the SAM and submit comments on any information currently available about the organization that a federal awarding agency previously entered. The federal awarding agency will consider any comments by the applicant, in addition to other information in the designated integrity and performance system, in making a judgment about the applicant's integrity, business ethics and record of performance under federal awards when determining a recipient's qualification prior to award, according to the qualification standards of the Department of Defense Grant and Agreement Regulations (DoDGARs), Section 22.415.

In accordance with National Security Presidential Memorandum-33 and all associated laws, all fundamental research funded by the DOW must be evaluated for affiliations with foreign entities. All applicant organizations must disclose foreign affiliations of all key personnel named on applications. Failure to disclose foreign affiliations of key personnel shall lead to withdrawal of recommendations to fund applications. Applicant organizations may be presented with an opportunity to mitigate identified risks, particularly those pertaining to influence from foreign entities specified in law. Implementation of mitigation discussions and utilization of the [DOD Component Decision Matrix](#) must decrease risk of foreign influence in accordance with the above-mentioned laws and guidance prior to award.

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7. Federal Award Notices

For each compliant full application received, the organizational representative(s) and PI will receive email notification when the funding recommendations are posted to eBRAP, typically within 6 weeks after programmatic review. At this time, each PI will receive a peer review summary statement on the strengths and weaknesses of the application and an information paper describing the application receipt and review process for the SCIRP award mechanisms. The information papers and a list of organizations and PIs recommended for funding are also posted on the program's page within the CDMRP website. After all awards are made, the CDMRP includes individual award information in a searchable [database](#).

If an application is recommended for funding, after the email notification is posted to eBRAP, a government representative will contact the person authorized to negotiate on behalf of the recipient organization.

Only an appointed DHACA Grants Officer may obligate the government to the expenditure of funds to an extramural organization. No commitment on the part of the government should be inferred from discussions with any other individual. ***The award document signed by the Grants Officer is the official authorizing document (i.e., assistance agreement).***

Intragovernmental obligations of funding will be made according to the terms of a negotiated Inter-Agency Agreement and managed by a CDMRP Science Officer.

Funding obligated to ***intragovernmental and intramural DOW organizations*** will be sent through the Military Interdepartmental Purchase Request (MIPR), Funding Authorization Document (FAD) or Direct Charge Work Breakdown Structure processes. Transfer of funds is contingent upon appropriate safety and administrative approvals. Intragovernmental and intramural DOW investigators and collaborators must coordinate receipt and commitment of funds through their respective Resource Manager/Task Area Manager/Comptroller or equivalent Business Official. 

An organization may, at its own risk and without the government's prior approval, incur obligations and expenditures to cover costs up to 90 days before the beginning date of the initial budget period of a new award.

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8. Post-Award Requirements


8.1. Administrative and National Policy Requirements

Applicable requirements in the DoDGARs found in 32 CFR, Chapter I, Subchapter C, and 2 CFR, Chapter XI, apply to grants and cooperative agreements resulting from this program announcement.

The GAI contain information regarding [administrative requirements](#) and [national policy requirements](#).

Refer to full text of the latest [DoD R&D Terms and Conditions](#) and the [DHACA Terms and Conditions](#) for further information.

If there are delinquencies in technical reporting requirements for any existing DHA or U.S. Army Medical Research and Development Command awards at the applicant organization, DHACA will not issue any new awards to the applicant organization until all delinquent reports have been submitted.

Applications recommended for funding that involve animals, human data, human specimens, human subjects or human cadavers must be reviewed for compliance with federal animal and/or human subjects protection requirements and must be approved by the DHA R&D Office of Research and Regulatory Compliance (ORRC), prior to implementation. This administrative review requirement is in addition to the local IACUC, Institutional Review Board (IRB) or Ethics Committee (EC) review. 

8.2. Reporting

Annual technical reports and quad charts, as well as a final technical report and quad chart, will be required. Annual and final technical progress reports must be prepared in accordance with the Research Performance Progress Report (RPPR).

Enrollment reporting on the basis of sex, race, and ethnicity will be required with each annual and final progress report. The [PHS Inclusion Enrollment Report](#) is available in eBRAP (**only required for [clinical research](#)**).

The Award Terms and Conditions will specify whether additional and/or more frequent reporting is required.

Awards resulting from this program announcement may entail additional reporting requirements related to recipient integrity and performance matters. Recipient organizations that have federal contract, grant and cooperative agreement awards with a cumulative total value greater than \$10M are required to provide information to the SAM about certain civil, criminal and administrative proceedings that reached final disposition within the most recent 5-year period and that were connected with their performance of a federal award. These recipients are required to disclose, semiannually, information about criminal, civil and administrative proceedings as specified in the applicable [Representations](#).

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8.3. Additional Requirements

PIs are expected to participate in at least one Interim Progress Review (IPR) for the funded project. For planning purposes, PIs can expect that the IPR will last no longer than one day and will be hosted virtually by the SCIRP. The invitation and format for the IPR will be provided by the Grants Officer's Representative at least 90 days prior to the scheduled IPR date.

Unless otherwise restricted, changes in the PI or organization will be allowed on a case-by-case basis, provided the intent of the award mechanism is met.



An organizational transfer of an award will not be allowed in the last year of the original period of performance or any extension thereof.

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9. Other Information

9.1. Program Announcement Version

Questions related to this program announcement should refer to the program name, the program announcement name and the program announcement version code CD26_01d.

9.2. Administrative Actions

After receipt of pre-applications or full applications, the following administrative actions may occur.

9.2.1. Rejection

The following will result in administrative rejection of the pre-application:

- Preproposal Narrative is missing.

The following will result in administrative rejection of the full application:

- Project Narrative is missing.
- Budget is missing.
- Submission of an application for which a letter of invitation was not issued.
- **Early-Career Partnership Option:** Partnership Statement ([Attachment 9](#)) is missing.

9.2.2. Modification

- Pages exceeding the specified limits will be removed prior to reviewing all documents.
- Documents not requested will be removed.

9.2.3. Withdrawal

The following may result in administrative withdrawal of the full application:

- A member of the FY26 SCIRP Programmatic Panel is named as being involved in the development or execution of the research proposed or is found to have assisted in the pre-application or application processes.
- The application includes the name(s) of personnel from either of the CDMRP peer or programmatic review companies for which conflicts cannot be adequately mitigated. For FY26, the identities of the peer review contractor and the programmatic review contractor may be found on the [CDMRP website](#).
- Personnel from applicant or collaborating organizations are found to have contacted persons involved in the review or approval process to gain protected evaluation information or to influence the evaluation process.
- The application from an extramural organization, including non-DOW federal agencies, is received through eBRAP.

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- The federal government recipient organization (including an intramural DOW organization):
(a) cannot accept and execute the entirety of the requested budget in FY26 funds; and/or (b) cannot coordinate the use of contractual, assistance or other appropriate agreements to provide funds to collaborators.
- The application fails to conform to this program announcement description.
- The application includes URLs, with the exception of links in the References Cited and Publication and/or Patent sections.
- The application includes research data that are classified and/or proposes research that may produce classified outcomes, or outcomes deemed sensitive to national security concerns.
- The invited application proposes a different research project than that described in the pre-application.
- The same research project is submitted to different funding opportunities within the same program and fiscal year.
- A clinical trial is proposed.
- Failure to submit all associated (Initiating and Partnering PI) applications by the deadline.
- The PI is named on more than one FY26 SCIRP Investigator-Initiated Research Award application.
- The PI, Initiating PI, or Partnering PI does not meet the [eligibility criteria](#).

9.2.4. Withhold

Applications that appear to involve research misconduct will be administratively withheld from further consideration pending organizational investigation. The organization will be required to provide the findings of the investigation to the DHACA Grants Officer for a determination of the final disposition of the application.

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Appendix 1. Full Application Submission Checklist

Full Application Components	Uploaded	
	PI/Initiating PI	Partnering PI
SF424 Research & Related Application for Federal Assistance <i>(Grants.gov submissions only)</i>	<input type="checkbox"/>	<input type="checkbox"/>
Summary (Tab 1) and Application Contacts (Tab 2) <i>(eBRAP submissions only)</i>	<input type="checkbox"/>	<input type="checkbox"/>
Attachments		
Project Narrative – Attachment 1, upload as “ProjectNarrative.pdf”	<input type="checkbox"/>	
Supporting Documentation – Attachment 2, upload as “Support.pdf”	<input type="checkbox"/>	
Technical Abstract – Attachment 3, upload as “TechAbs.pdf”	<input type="checkbox"/>	
Lay Abstract – Attachment 4, upload as “LayAbs.pdf”	<input type="checkbox"/>	
Statement of Work – Attachment 5, upload as “SOW.pdf”	<input type="checkbox"/>	<input type="checkbox"/>
Impact Statement – Attachment 6, upload as “Impact.pdf”	<input type="checkbox"/>	
Long-Term Goal – Attachment 7, upload as “Longterm.pdf”	<input type="checkbox"/>	
Animal Research Plan <i>(Animal studies only)</i> – Attachment 8, upload as “AnimalResPlan.pdf”	<input type="checkbox"/>	
Partnership Statement <i>(Early-Career Partnership Option only)</i> – Attachment 9, upload as “Partnership.pdf”	<input type="checkbox"/>	
Representations <i>(Grants.gov submissions only)</i> – Attachment 10, upload as “RequiredReps.pdf”	<input type="checkbox"/>	<input type="checkbox"/>
Suggested Intragovernmental/Intramural Budget Form <i>(if applicable)</i> – Attachment 11, upload as “IGBudget.pdf”	<input type="checkbox"/>	<input type="checkbox"/>
Additional Application Materials		
Research & Related Senior/Key Person Profile (Expanded)	<input type="checkbox"/>	<input type="checkbox"/>
Attach Biographical Sketch for Senior/Key Persons (Biosketch_LastName.pdf)	<input type="checkbox"/>	<input type="checkbox"/>
Attach Current/Pending Support for Senior/Key Persons (Support_LastName.pdf)	<input type="checkbox"/>	<input type="checkbox"/>
Research & Related Budget	<input type="checkbox"/>	<input type="checkbox"/>
Project/Performance Site Location(s)	<input type="checkbox"/>	<input type="checkbox"/>
Research & Related Subaward Budget Attachment(s) <i>(if applicable)</i>	<input type="checkbox"/>	<input type="checkbox"/>

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Appendix 2. Acronym List

ARRIVE	Animal Research: Reporting of In Vivo Experiments
CDMRP	Congressionally Directed Medical Research Programs
CFR	Code of Federal Regulations
DHA	Defense Health Agency
DHA R&D	Defense Health Agency Research and Development
DHACA	Defense Health Agency Contracting Activity
DOD	U.S. Department of Defense
DoDGARs	Department of Defense Grant and Agreement Regulations
DOW	U.S. Department of War
eBRAP	Electronic Biomedical Research Application Portal
EC	Ethics Committee
ET	Eastern Time
FAD	Funding Authorization Document
FY	Fiscal Year
GAI	General Application Instructions
IACUC	Institutional Animal Care and Use Committee
IIRA	Investigator-Initiated Research Award
IPR	Interim Progress Review
IRB	Institutional Review Board
M	Million
MIPR	Military Interdepartmental Purchase Request
ORRC	Office of Research and Regulatory Compliance
PDF	Portable Document Format
PHS	Public Health Service
PI	Principal Investigator
R&D	Research and Development
RPPR	Research Performance Progress Report
SAM	System for Award Management
SCI	Spinal Cord Injury
SCIRP	Spinal Cord Injury Research Program
SF424 R&R	Standard Form 424 (Application for Federal Assistance, Research & Related)
SOW	Statement of Work
STROBE	STrengthening the Reporting of OBServational studies in Epidemiology

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UEI	Unique Entity Identifier
URL	Uniform Resource Locator
USC	United States Code
VA	U.S. Department of Veterans Affairs