



CANCER PREVENTION & RESEARCH
INSTITUTE OF TEXAS

REQUEST FOR APPLICATIONS

RFA R-26.1-IIRAP

**Individual Investigator Research Awards
for Prevention and Early Detection**

**Please also refer to the Instructions for Applicants document, which will be posted on
January 14, 2025.**

**Due to the highly competitive nature of this RFA mechanism and available funds for FY26,
CPRIT projects that only 10% of submitted applications will be funded.**

Application Receipt Opening Date: February 18, 2025

Application Receipt Closing Date: May 6, 2025

FY 2026

Fiscal Year Award Period

September 1, 2025-August 31, 2026

TABLE OF CONTENTS

1. BRIEF DESCRIPTION OF RFA	4
2. KEY POINTS	4
3. ABOUT CPRIT	5
3.1. ACADEMIC RESEARCH PROGRAM PRIORITIES.....	6
4. RATIONALE	7
5. RESEARCH OBJECTIVES	7
6. FUNDING INFORMATION	9
7. ELIGIBILITY	10
8. RESUBMISSION POLICY	12
9. RENEWAL POLICY	12
10. RESPONDING TO THIS RFA	12
10.1. APPLICATION SUBMISSION GUIDELINES.....	12
10.1.1. <i>Submission Deadline Extension</i>	13
10.2. APPLICATION COMPONENTS.....	13
10.2.1. <i>Abstract and Significance (5,000 characters)</i>	13
10.2.2. <i>Layperson’s Summary (2,000 characters)</i>	14
10.2.3. <i>Aims and Subaims</i>	14
10.2.4. <i>Timeline (1 page)</i>	14
10.2.5. <i>Specific Aims (1 page)</i>	15
10.2.6. <i>Resubmission Summary (2 pages)</i>	15
10.2.7. <i>Renewal Summary (2 pages)</i>	15
10.2.8. <i>Research Plan (10 pages)</i>	15
10.2.9. <i>Vertebrate Animals (1 page)</i>	16
10.2.10. <i>Human Subjects (2 pages)</i>	16
10.2.11. <i>Publications/References</i>	16
10.2.12. <i>Budget and Justification</i>	16
10.2.13. <i>Biographical Sketches (5 pages each)</i>	18
10.2.14. <i>Current and Pending Support</i>	18
10.2.15. <i>Institutional/Collaborator Support and/or Other Certification (4 pages)</i>	18
10.2.16. <i>Previous Summary Statement</i>	18
10.3. FORMATTING INSTRUCTIONS.....	19
11. APPLICATION REVIEW	20
11.1. PRELIMINARY EVALUATION.....	20
11.2. FULL PEER REVIEW.....	20
11.3. CONFIDENTIALITY OF REVIEW.....	21
11.4. REVIEW CRITERIA.....	22
11.4.1. <i>Primary Criteria</i>	22
11.4.2. <i>Secondary Criteria</i>	23
12. KEY DATES	23
13. AWARD ADMINISTRATION	24
14. REQUIREMENT TO DEMONSTRATE AVAILABLE FUNDS	25
15. CONTACT INFORMATION	26
15.1. HELPDESK.....	26
15.2. SCIENTIFIC AND PROGRAMMATIC QUESTIONS.....	26

RFA VERSION HISTORY

- 1/14/25 RFA released
- 3/14/25 Clarified language on page 11 (Section 7: Eligibility) and 17 (Section 10.2.12: Budget and Justification) regarding clinical trials
- 3/26/25 Updated links to Texas Administrative Code

1. BRIEF DESCRIPTION OF RFA

- Supports applications that propose clinical and population-based projects designed to develop effective prevention and early detection interventions to reduce cancer risk, mortality, and morbidity among Texans.
- Applicants may request up to a maximum of \$300,000 per year for a period of up to 4 years.
- **Multi-Principal Investigators (MIs)** are allowed under this Request for Applications (RFA). Refer to the Instructions for Applicants (IFA) document for definition and eligibility of MIs.
- See application limitations for **Principal Investigators (PIs)/MIs**, regardless of whether the MI is from the primary institution or a subcontracted institution.
- **Note that Cancer Prevention and Research Institute of Texas (CPRIT) does not allow the use of the term Co-PI.**
- Minimum level of effort for the PI and/or MIs throughout the project period is required.
- A PI and/or MI may only submit **one application** for this RFA (defined as either a new application, resubmission application, or competitive *renewal application).
- *Renewal applications are limited to one competitive renewal under this RFA, regardless of the year of the funded parent IIRAP award.
- The FY26 salary cap is \$225,000 per year.

2. KEY POINTS

This version of the Individual Investigator Research Award for Prevention and Early Detection (IIRAP) invites clinical and population-based projects designed to develop effective prevention, early detection, or behavioral interventions to reduce cancer risk, mortality, and morbidity among Texans. Examples of projects responsive to this RFA include those planning clinical or population-based projects that will do the following:

- (1) Identify risk factors for cancers affecting Texans including social, environmental, and behavioral factors, genetic markers, and health disparities.
- (2) Validate biomarkers, genomic technologies, or imaging methods in Texas populations at high risk for cancer that will lead to detection of cancers at an early stage.

- (3) Develop and validate innovative interventions for the prevention or early detection of cancers with particular emphasis on Texans at highest risk for cancer.
- (4) Increase implementation and/or dissemination of evidence-based interventions for the prevention and/or early detection of cancers among all Texans as well as Texas populations at high risk for cancer.
- (5) Conduct studies within the broad continuum of cancer survivorship research aimed at improving the health and quality of life of cancer survivors, including survivors of childhood or adolescent and adult cancers, and caretakers. For example, research to improve cancer prevention, screening, early detection, treatment and late effects, as well as to develop approaches to optimal [supportive care](#) are responsive to this RFA.
- (6) Conduct health services research in Texas populations at highest risk for cancer to identify the most effective ways to address the disparities (eg, through systems change, outreach, access) and delivery of early detection and preventive care.

Basic research using laboratory and/or animal models designed to identify underlying mechanisms causing cancer ***is not responsive*** to this RFA and should be directed to other CPRIT mechanisms supporting basic discovery research. Please contact research@cprit.texas.gov before initiating an application should you have questions regarding this statement of no responsiveness.

Basic and clinical research to discover or to evaluate chemoprevention strategies ***is not responsive*** to this RFA and should be directed to other CPRIT mechanisms supporting basic discovery research or clinical translational research. Please contact research@cprit.texas.gov before initiating an application should you have questions regarding this statement of no responsiveness.

Applicants may request up to \$300,000 per year in CPRIT funds over 4 years.

3. ABOUT CPRIT

The State of Texas has established CPRIT, which may issue up to \$6 billion in general obligation bonds to fund grants for cancer research and prevention.

CPRIT is charged by the Texas Legislature to do the following:

- Create and expedite innovation in the area of cancer research and in enhancing the potential for a medical or scientific breakthrough in the prevention of, or cures for, cancer;
- Attract, create, or expand research capabilities of public or private institutions of higher education and other public or private entities that will promote a substantial increase in cancer research and in the creation of high-quality new jobs in the State of Texas; and
- Develop and implement the Texas Cancer Plan.

3.1. Academic Research Program Priorities

The Texas Legislature has charged the CPRIT Oversight Committee with establishing program priorities on an annual basis. These priorities are intended to provide transparency with regard to how the Oversight Committee directs the orientation of the agency’s funding portfolio.

Established Principles:

- Scientific excellence and impact on cancer
- Increasing the life sciences infrastructure in all regions of the state
- Reducing cancer disparities

The program priorities for academic research adopted by the Oversight Committee include funding projects that address or utilize the following:

- Recruitment of outstanding cancer researchers to Texas
- Investment in core facilities
- A broad range of innovative, investigator-initiated research projects
- Implementation research to accelerate the adoption and deployment of evidence-based prevention, early detection or risk assessment approaches, and interventions
- Computational oncology and analytic methods
- Childhood and adolescent cancers
- Hepatocellular cancer
- Expanding access to innovative clinical trials
- Cancer survivorship research to enhance the health and well-being of cancer survivors and caregivers.

4. RATIONALE

Nowhere is there greater potential to reduce the burden of cancer than by reducing its incidence. It is estimated that more than half of all cancer deaths could be prevented. It is also well established that early detection of cancer improves cancer mortality. The goal of this RFA is to promote research to develop effective cancer prevention and early detection approaches for Texans.

Many opportunities exist to make greater progress in reducing cancer mortality in Texas through cancer prevention and early detection. Large numbers of Texans at average risk for cancers for which there are proven prevention and early detection strategies fail to take advantage of these strategies. This is particularly true among the medically underserved and hard-to-reach populations where numerous barriers inhibit the adoption and implementation of proven strategies for prevention or early detection. Early detection and prevention can improve mortality in individuals with an inherited predisposition to cancer, yet testing for inherited cancer syndromes in appropriate populations is vastly underused. Novel technologies and noninvasive approaches (eg, liquid biopsies, novel imaging strategies) to detect cancer biomarkers are undergoing rapid advances that offer opportunities to improve surveillance of individuals at average or high risk for cancer. Behavioral sciences research informs the development and uptake of cancer-reducing strategies, eg, tobacco cessation strategies and obesity control. Greater attention to implementation of evidence-based identification of at-risk individuals may empower them to make and act upon informed, cancer-preventing health decisions.

5. RESEARCH OBJECTIVES

This RFA solicits clinical and population-based projects designed to improve current methods and develop new strategies for the prevention and early detection of cancer among Texans. Examples of projects responsive to this RFA include those planning clinical or population-based novel projects to do the following:

- (1) To identify social, environmental, and behavioral factors, genetic markers, and health disparities that underlie cancer risk and predisposition among Texans at high risk for cancer.

- (2) To validate biomarkers, genomic technologies, or imaging methods in populations at risk for cancer that will lead to detection of cancers at an early stage.
- (3) To develop and validate innovative interventions for the prevention or early detection of cancers with particular emphasis on Texans at highest risk for cancer.
- (4) To conduct behavioral science studies that inform the development of interventions to reduce cancer risk, eg, conducting obesity research, increasing uptake of screening or cancer vaccinations, or reducing use of combustible tobacco and vaping products, particularly in populations with cancer disparities.
- (5) To increase implementation and/or dissemination of evidence-based interventions for the prevention and/or early detection of cancers among all Texans as well as those populations at high risk.
- (6) To conduct health services research in populations at highest risk to identify the most effective ways to address the disparities (eg, through systems change, outreach, access) and delivery of early detection and preventive care.
- (7) To conduct survivorship research to predict outcomes, to identify children and adolescents at risk of developing late effects of treatment, or to improve quality of life for survivors of childhood and adolescent cancers or caretakers of individuals with cancer.
- (8) To conduct survivorship research on adults following a diagnosis of cancer, eg, the effects of cancer and its treatment on patients with other health conditions, health care utilization and access, and health equity, or the effects on caretakers.

Basic research using laboratory and/or animal models designed to identify underlying mechanisms causing cancer ***is not responsive*** to this RFA and should be directed to other CPRIT mechanisms supporting basic discovery research.

Basic and clinical research to discover or to evaluate chemoprevention strategies ***is not responsive*** to this RFA and should be directed to other CPRIT mechanisms supporting basic discovery research or clinical translational research.

Basic and clinical research to predict outcomes, to identify children and adolescents at risk of developing late effects of treatment or to improve quality of life for survivors of childhood and

adolescent cancers ***is not responsive*** to this RFA and should be directed to other CPRIT mechanisms supporting research on cancer in children and adolescents.

The *degree of relevance* to reducing the burden of cancer in Texas is a critical criterion for evaluation of projects for funding by CPRIT ([section 11.4.1](#)). Although CPRIT is required to spend 10% of its budget on cancer prevention, CPRIT's Cancer Prevention Program focuses exclusively on the delivery of evidence-based interventions to underserved populations and does not fund prevention research. **Thus, there is a unique opportunity for applicants to this RFA in CPRIT's Academic Research Program to develop collaborations with CPRIT Prevention Program grantees who have developed community-based partnerships and can provide access to Texas populations for research designed to improve current practices. Projects that propose such research collaborations with existing CPRIT Prevention Program awardees including the CPRIT-funded *Texas Collaborative Center for Hepatocellular Cancer* (<https://www.bcm.edu/research/labs-and-centers/research-centers/texas-collaborative-center-for-hepatocellular-cancer>) are strongly encouraged, as well as research that focuses on cancer survivorship to enhance the health and well-being of all cancer survivors and caregivers.**

6. FUNDING INFORMATION

Applicants may request a maximum up to \$300,000 in total costs per year for up to 4 years for clinical or population-based research, including implementation or dissemination research designed to accelerate the adoption and deployment of sustainable, evidence-based cancer prevention and screening interventions at multiple levels and in different clinical and community settings. Funds may be used for salary and fringe benefits, research supplies, equipment, subject participation costs, and travel to scientific/technical meetings or collaborating institutions. Requests for funds to support construction and/or renovation will not be approved under this funding mechanism. State law limits the amount of indirect costs to no more than 5% of the total award amount.

Please see section [10.2.12](#) and the IFA for additional information.

7. ELIGIBILITY

- The applicant must be a Texas-based entity. Any not-for-profit institution or organization that conducts research is eligible to apply for funding under this award mechanism. A public or private company is not eligible for funding under this award mechanism; these entities must use the appropriate award mechanism(s) under CPRIT's Product Development Research Program.
- The PI must have a doctoral degree, including MD, PhD, DDS, DMD, DrPH, DO, DVM, or equivalent, and be a full-time resident of Texas during the time the research that is the subject of the grant is conducted.
- The PI must hold an appointment at the rank of assistant, associate or professor, tenure track or tenured (or equivalent), at an accredited academic institution, research institution, industry, government agency, or private foundation. The PI must also have appropriate designated laboratory or clinical space, or other facilities in which to conduct the proposed studies.
- This award mechanism allows MIs for projects that require a team science approach (see the IFA for guidelines on CPRIT rules for MI awards).
- For applications that include 1 PI, the PI is required to maintain a minimum of 15% level of effort through the duration of the entire award period. For applications that include MIs, each PI is required to maintain a minimum of 10% effort through the entire duration of the award period.
- A PI may not submit applications to this RFA and to RFA R-26.1-IIRA, RFA R-26.1-IIRACCA, RFA R-26.1-IIRACSBC, RFA R-26.1-IIRACT, or RFA R-26.1-IIRAEOC.
- A PI/MI may not submit more than 1 application, either a new, resubmission, **or** renewal application under this RFA during this funding cycle
- A PI may not be a part of more than one application, whether as a single applicant or as part of an MI on an application, under this RFA and RFA R-26.1-IIRA, RFA R-26.1-IIRACCA, RFA R-26.1-IIRACSBC, RFA R-26.1-IIRACT, or RFA R-26.1-IIRAEOC.
- An individual may serve as a PI on no more than 3 active CPRIT Academic Research grants. Recruitment grants and Research Training Awards do not count toward the 3-grant maximum; however, CPRIT considers MIRA Project MIs equivalent to a PI. For the purpose of calculating the number of active grants, CPRIT will consider the number

of active grants at the time of the award contract effective date (for this cycle expected to be December 1, 2025).

- Applications that address basic or animal-based projects, cancers in children and adolescents, computational systems biology of cancer, or therapeutic clinical trials should be submitted under the appropriate targeted RFA. Note that a smoking cessation clinical trial using drugs, can fit under this RFA.
- Collaborating organizations may include public, not-for-profit, and for-profit entities. Such entities may be located outside of the State of Texas, but non-Texas-based organizations are not eligible to receive CPRIT funds.
- An applicant is eligible to receive a grant award only if the applicant certifies that the applicant institution or organization, including the PI, any senior member or key personnel listed on the grant application, or any officer or director of the grant applicant's institution or organization (or any person related to 1 or more of these individuals within the second degree of consanguinity or affinity), has not made and will not make a contribution to CPRIT or to any foundation specifically created to benefit CPRIT.
- An applicant is not eligible to receive a CPRIT grant award if the applicant PI, any senior member, or key personnel listed on the grant application, or any officer or director of the grant applicant's organization or institution is related to a CPRIT Oversight Committee member.
- The applicant must report whether the applicant institution or organization, the PI, or other individuals who contribute to the execution of the proposed project in a substantive, measurable way, regardless of whether those individuals are slated to receive salary or compensation under the grant award, are currently ineligible to receive federal grant funds or have had a grant terminated for cause within 5 years prior to the submission date of the grant application.
- CPRIT grants will be awarded by contract to successful applicants. Certain contractual requirements are mandated by Texas law or by administrative rules. Although applicants need not demonstrate the ability to comply with these contractual requirements at the time the application is submitted, applicants should make themselves aware of these standards before submitting a grant application. Significant issues addressed by the

CPRIT contract are listed in [section 13](#) and [section 14](#). All statutory provisions and relevant administrative rules can be found at www.cprit.texas.gov.

8. RESUBMISSION POLICY

An application previously submitted to CPRIT but not funded may be resubmitted **once** and must follow all resubmission guidelines and be eligible for this mechanism under the criteria in Section 5. If a previously submitted application does not meet the new criteria for this mechanism outlined in [section 5](#), it must be resubmitted under the IIRA mechanism. **More than 1 resubmission is not permitted.** Applications resubmitted more than once will be administratively withdrawn. An application is considered a resubmission if the proposed project is the same project presented in the original submission. A change in the identity of the PI for a project or a change in the project title that was previously submitted to CPRIT, or omission or modification of an aim do not constitute a new application; the application would be considered a resubmission. This policy is in effect for all applications submitted to date. See [section 10.2.6](#).

9. RENEWAL POLICY

An application funded by CPRIT under this mechanism may be submitted for **one** competitive renewal. An application originally funded by CPRIT as an IIRA that is now appropriate for the IIRAP mechanism may be submitted under this RFA for a competitive renewal. See [section 10.2.7](#). An application originally funded by CPRIT under this mechanism that does not meet the new criteria outlined in [section 5](#) must be submitted under the IIRA mechanism. Competitive renewals are not subject to preliminary evaluation. Renewal applications move directly to the full peer review phase. See [section 11.2](#).

10. RESPONDING TO THIS RFA

10.1. Application Submission Guidelines

Applications must be submitted via the CPRIT Application Receipt System (CARS) (<https://CPRITGrants.org>). **Only applications submitted through this portal will be considered eligible for evaluation.** The applicant is eligible solely for the grant mechanism specified by the RFA under which the grant application was submitted. The PI must create a user account in the system to start and submit an application. Furthermore, the Application Signing

Official (a person authorized to sign and submit the application for the organization) and the Grants Contract/Office of Sponsored Projects Official (the individual who will manage the grant contract if an award is made) also must create a user account in CARS. Applications will be accepted beginning at 7 AM central time on February 18, 2025, and must be submitted by 4 PM central time on May 6, 2025. **Submission of an application is considered an acceptance of the terms and conditions of the RFA.**

10.1.1. Submission Deadline Extension

The submission deadline may be extended upon a showing of extenuating circumstances. A request for a deadline extension based on the need to complete multiple CPRIT or other grants applications will be denied. All requests for extension of the submission deadline must be submitted via email to the CPRIT [Helpdesk](#) within 24 hours of the submission deadline. Submission deadline extensions, including the reason for the extension, will be documented as part of the grant review process records. Please note that deadline extension requests are very rarely approved.

10.2. Application Components

Applicants are advised to follow all instructions to ensure accurate and complete submission of all components of the application. Please refer to the IFA for details. The IFA will be available when the application receipt system opens. Submissions that are missing 1 or more components or do not meet the eligibility requirements listed in [section 7](#) will be administratively withdrawn without review.

10.2.1. Abstract and Significance (5,000 characters)

It is the responsibility of the applicant to capture CPRIT's attention primarily with the Abstract and Significance of the research alone. Therefore, applicants are advised to prepare this section wisely. Based on the Abstract and Significance statement (and the Specific Aims), applications that are judged to offer only modest contributions to the field of cancer research or that do not sufficiently capture the reviewers' interest may be excluded from peer review (see [section 11.1](#)). Applicants should not waste this valuable space by stating obvious facts (eg, that cancer is a significant problem; that better diagnostic and therapeutic approaches are needed urgently; or that the type of cancer of interest to the PI is important, vexing, or deadly).

Clearly explain the question or problem to be addressed and the approach to its answer or solution. Provide the hypothesis to be tested, the specific aims, and the overall experimental approaches.

Clearly address how the proposed project, if successful, will have a major impact on cancer. Indicate whether this research addresses the identification, or mitigation, of cancer disparities. Summarize how the proposed research creates new paradigms or challenges existing ones. Indicate whether this research plan represents a new direction for the PI.

10.2.2. Layperson's Summary (2,000 characters)

Provide a layperson's summary of the proposed work. Describe, in simple, nontechnical terms, the overall goals of the proposed work, the type(s) of cancer addressed, the potential significance of the results, and the impact of the work on advancing the field of cancer research, early detection, prevention, or treatment. The information provided in this summary will be made publicly available by CPRIT, particularly if the application is recommended for funding. **Do not include any proprietary information in the layperson's summary.** The layperson's summary will also be used by advocate reviewers ([section 11.2](#)) to evaluate the significance and impact of the proposed work during peer review.

10.2.3. Aims and Subaims

Please provide a description of the aims and subaims and milestones to be achieved for each year of the project. At least 1 specific aim and 1 subaim are required. Provide 2 to 3 sentences describing activities to be performed and anticipated milestones. These aims will also be used during the submission and evaluation of progress reports and assessment of project success. Up to 5 specific aims and 5 subaims per specific aim may be entered.

10.2.4. Timeline (1 page)

Provide an outline of anticipated major milestones to be tracked. Timelines will be reviewed for reasonableness, and adherence to timelines will be a criterion for continued support of successful applications.

If the application is approved for funding, this section will be included in the award contract. Applicants are advised not to include information that they consider confidential or proprietary when preparing this section.

10.2.5. Specific Aims (1 page)

Please provide a 1-page summary of the aims of the proposal. The Specific Aims page should identify the problem or gap in our current knowledge. It should present a hypothesis and briefly describe the aims, subaims, and approaches. Address the proposal's innovation, novel approaches and significance and impact on the field and cancer research. Please also refer to the template located in [Current Funding Opportunities](#) for Academic Research in CARS.

10.2.6. Resubmission Summary (2 pages)

Applicants preparing a resubmission must describe the approach to the resubmission. If a summary statement was prepared for the original application review, applicants are advised to address all noted concerns.

Note: An application previously submitted to CPRIT but not funded may be resubmitted **once** after careful consideration of the reasons for lack of prior success. Applications that received overall numerical scores of 5 or higher are likely to need considerable attention. Applicants may prepare a new research plan or modify the original research plan and mark the changes.

However, **all resubmitted applications should be carefully reconstructed**; a simple revision of the prior application with only editorial or technical changes is not sufficient, and applicants are advised not to direct reviewers to such modest changes.

10.2.7. Renewal Summary (2 pages)

Applicants preparing a renewal must describe and demonstrate that appropriate/adequate progress has been made on the current funded award to warrant further funding. Publications and manuscripts in press that have resulted from work performed during the initial funded period should be listed in the renewal summary.

10.2.8. Research Plan (10 pages)

Background: Provide the rationale behind the proposed project, emphasizing the pressing problem in cancer research that will be addressed.

Hypothesis and Specific Aims: Concisely state the hypothesis to be tested, and specific aims to be addressed by the research described in the application. Describe the significance, innovation, and potential impact of the research.

Research Strategy: Describe the experimental design, including methods, anticipated results, potential problems or pitfalls, and alternative approaches. Preliminary data that support the proposed hypothesis are strongly encouraged.

10.2.9. Vertebrate Animals (1 page)

If vertebrate animals will be used, provide a detailed plan of the protocols that will be followed and justification for the number of animals used with reference to biostatistical input for sample selection and evaluation. Certification of approval by the Institutional Animal Care and Use Committee (IACUC) of the proposed animal use will be required before funding can occur.

10.2.10. Human Subjects (2 pages)

If human subjects or human biological samples will be used, provide a detailed plan for recruitment of subjects and/or the acquisition of samples that will meet the time constraints of this award mechanism. Certification of approval of these plans by the institutional review board (IRB) will be required before funding can occur.

10.2.11. Publications/References

Provide a concise and relevant list of publications/references cited for the application.

10.2.12. Budget and Justification

Provide a detailed justification of the budget for the entire proposed period of support, including salaries and benefits, supplies, equipment, patient care costs, animal care costs, and other expenses, subject participation costs, and travel to scientific/technical meetings or collaborating institutions. The PI and related research project staff are expected to attend CPRIT's conference. CPRIT funds may be used to send up to 2 people to the conference. Meals are not reimbursable for trips that do not include an overnight stay. Requests for funds to support construction and/or renovation will not be approved under this funding mechanism. Do not exceed \$300,000 per year for studies, including implementation research designed to accelerate the adoption and deployment of sustainable, evidence-based cancer prevention and screening interventions at

multiple levels and in different clinical and community settings. Applicants are advised not to interpret the maximum allowable request under this award as a suggestion that they should expand their anticipated budget to this level. Reasonable budgets clearly work in favor of the applicant.

In preparing the requested budget, applicants should be aware of the following:

- Equipment having a useful life of more than 1 year and an acquisition cost of \$10,000 or more per unit must be specifically approved by CPRIT. An applicant does not need to seek this approval prior to submitting the application. Per Texas Grant Management Standards, as of September 1, 2024, for all CPRIT grantees, the minimum threshold for equipment purchases is \$10,000. Generally, any purchases up to \$9,999 should now be categorized as supplies.
- Texas law limits the amount of grant funds that may be spent on indirect costs to no more than 5% of the total award amount (5.263% of the direct costs). Guidance regarding indirect cost recovery can be found in CPRIT's Administrative Rules, which are available at www.cprit.texas.gov. So-called grants management and facilities fees (eg, sponsored programs fees; grants and contracts fees; electricity, gas, and water; custodial fees; maintenance fees) may not be requested. Applications that include such budgetary items will be rejected administratively and returned without review.
- The maximum annual salary (also referred to as direct salary or institutional base salary) that an individual may request under a CPRIT award for FY 2026 is \$225,000; CPRIT FY 2026 is from September 1, 2025, through August 31, 2026. Salary does not include fringe benefits and/or facilities and administrative costs, also referred to as indirect costs. An individual's institutional base salary is the annual compensation that the applicant organization pays for an individual's appointment, whether that individual's time is spent on research, teaching, patient care, or other activities. Base salary excludes any income that an individual may be permitted to earn outside of his or her duties to the applicant organization.
- Funds can be used to pay for costs that a cancer clinical trial participant may have associated with their participation in a non-therapeutic clinical trial, including (1) transportation, including car mileage, parking, bus fare, taxi or ride hailing fare exclusive

of tips, and commercial economy class airfare within the borders of the State of Texas and (2) lodging.

10.2.13. Biographical Sketches (5 pages each)

Applicants should provide a biographical sketch that describes their education and training, professional experience, awards and honors, and publications relevant to cancer research. A biographical sketch must be provided for the PI and, if applicable, any additional MIs (for an MI application), as required by the online application receipt system. Up to **5 additional** biographical sketches for key personnel may be provided; these should be concatenated into a single PDF with a 25-page limit. **Each biographical sketch must not exceed 5 pages.** The NIH biosketch format is appropriate.

10.2.14. Current and Pending Support

Describe the funding source and duration of all current and pending support for all personnel who have included a biographical sketch with the application. For each award, provide the title, a 2-line summary of the goal of the project and, if relevant, a statement of overlap with the current application and a preliminary brief statement on plan to mitigate potential overlap. At a minimum, current and pending support of the PI and, if applicable, any additional MIs (for an MI application), must be provided. Refer to the sample current and pending support document located in [Current Funding Opportunities](#) for Academic Research in CARS.

10.2.15. Institutional/Collaborator Support and/or Other Certification (4 pages)

Applicants may provide letters of institutional support, collaborator support, and/or other certification documentation relevant to the proposed project. A maximum of 4 pages may be provided.

10.2.16. Previous Summary Statement

If the application is being resubmitted, the summary statement of the original application review, if previously prepared, will be automatically appended to the resubmission. The applicant is not responsible for providing this document.

Applications that are missing 1 or more of these components; exceed the specified page, word, or budget limits; or that do not meet the eligibility requirements listed above will be administratively rejected without review.

10.3. Formatting Instructions

Formatting guidelines for all submitted CPRIT applications are as follows:

- **Language:** English
- **Document Format:** PDF only
- **Font Type/Size:** Arial (11 point), Calibri (11 point), or Times New Roman (12 point)
- **Line Spacing:** Single
- **Page Size:** 8.5 x 11 inches
- **Margins:** 0.75 inch, all directions
- **Color and High-Resolution Images:** Images, graphs, figures, and other illustrations may be submitted as part of the submitted document. Applicants should include text to explain illustrations that may be difficult to interpret when printed in black and white.
- **Scanning Resolution:** Images and figures must be of lowest reasonable resolution that permits clarity and readability. Unnecessarily large files will NOT be accepted, especially those that include only text.
- **References:** Applicants should use a citation style that includes the full name of the article and that lists at least the first 3 authors. Official journal abbreviations may be used. An example is included below; however, other citation styles meeting these parameters are also acceptable if the journal information is stated. URLs of publications referenced in the application may be included.

Smith, P.T., Doe, J., White, J.M., et al (2006). Elaborating on a novel mechanism for cancer progression. *Journal of Cancer Research*, 135: 45-67.
- **Internet URLs:** Applicants are encouraged to provide the URLs of publications referenced in the application; however, applicants must not include URLs directing reviewers to websites containing additional information about the proposed research.
- **Headers and Footers:** These should not be used unless they are part of a provided template. Page numbers may be included in the footer (see **Page Numbering section** below).

- **Page Numbering:** Pages should be numbered at the bottom right corner of each page.
- All attachments that require signatures must be , printed, signed, scanned, and then uploaded in PDF format.

11. APPLICATION REVIEW

11.1. Preliminary Evaluation

To ensure the timely and thorough review of only the most innovative and cutting-edge research with the greatest potential for advancement of cancer research, all eligible applications shall undergo a preliminary evaluation for scientific merit and potential impact by a CPRIT Scientific Peer Review Panel consisting of scientific experts.

The preliminary evaluation is based on a subset of material presented in the application—namely, the Abstract and Significance and the Specific Aims page. Applications that do not sufficiently capture the reviewers’ interest at this stage will not be considered for full peer review. Such applications will have been judged to offer only modest contributions to the field of cancer research and will be excluded from further peer review.

The applicant will be notified of the decision to disapprove the application after the preliminary evaluation has concluded. Due to the volume of applications to be reviewed, comments provided by reviewers at the preliminary evaluation stage may be limited.

11.2. Full Peer Review

Applications that pass preliminary evaluation will undergo further review using a 2-stage peer review process: (1) Full peer review and (2) Prioritization of grant applications by the CPRIT Scientific Review Council. In the first stage, applications will be evaluated by an independent peer review panel consisting of scientific experts as well as advocate reviewers using the criteria listed in [section 11.4](#). In the second stage, applications judged to be most meritorious by the peer review panels will be evaluated and recommended for funding by the CPRIT Scientific Review Council based on comparisons with applications from all the peer review panels and programmatic priorities. Applications approved by the Scientific Review Council will be forwarded to the CPRIT Program Integration Committee (PIC) for review. The PIC will consider factors including program priorities set by the Oversight Committee, portfolio balance across

programs, and available funding. The CPRIT Oversight Committee will vote to approve each grant award recommendation made by the PIC. The grant award recommendations will be presented at an open meeting of the Oversight Committee and must be approved by two-thirds of the Oversight Committee members present and eligible to vote. The review process is described more fully in CPRIT's Administrative Rules, [chapter 703, sections 703.6 to 703.8](#).

11.3. Confidentiality of Review

Each stage of the application review is conducted confidentially, and all CPRIT Scientific Peer Review Panel members, Scientific Review Council members, PIC members, CPRIT employees, and Oversight Committee members with access to grant application information are required to sign nondisclosure statements regarding the contents of the applications. All technological and scientific information included in the application is protected from public disclosure pursuant to Health and Safety Code §102.262(b).

Individuals directly involved with the review process operate under strict conflict-of-interest prohibitions. All CPRIT Scientific Peer Review Panel members and Scientific Review Council members are non-Texas residents.

An applicant will be notified regarding the peer review panel assigned to review the grant application. Peer review panel members are listed by panel on CPRIT's website.

By submitting a grant application, the applicant agrees and understands that the only basis for reconsideration of a grant application is limited to an undisclosed Conflict of Interest as set forth in CPRIT's Administrative Rules ([Texas Administrative Code RULE §703.9](#)).

Communication regarding the substance of a pending application is prohibited between the grant applicant (or someone on the grant applicant's behalf) and the following individuals: an Oversight Committee Member, a PIC Member, a Scientific Peer Review Panel member, or a Scientific Review Council member. Applicants should note that the CPRIT PIC comprises the CPRIT Chief Executive Officer, the Chief Scientific Officer, the Chief Prevention Officer, the Chief Product Development Research Officer, and the Commissioner of State Health Services. The prohibition on communication begins on the first day that grant applications for the grant mechanism are accepted by CPRIT and extends until the grant applicant receives notice regarding a final decision on the grant application. The prohibition on communication does not

apply to the time when preapplications or letters of interest are accepted. Intentional, serious, or frequent violations of this rule may result in the disqualification of the grant application from further consideration for a grant award.

11.4. Review Criteria

Full peer review of applications will be based on primary scored criteria and secondary unscored criteria, listed below. Peer review committees will evaluate and score each primary criterion and subsequently assign an overall score that reflects a complete assessment of the application. **The overall score will not be an average of the individual criteria scores; rather, it will reflect the reviewers' overall impression of the application. Evaluation of the scientific merit of each application is within the sole discretion of the peer reviewers.**

11.4.1. Primary Criteria

The primary criteria will evaluate the scientific merit and potential impact of the proposed work contained in the application. Concerns with any of these criteria potentially indicate a major flaw in the significance and/or design of the proposed study. Primary criteria include the following:

Significance and Impact: Will the results of this research, if successful, significantly change the research of others or the opportunities for better cancer prevention or early detection? Is the application innovative? Does the applicant propose new paradigms or challenge existing ones? Does the project develop state-of-the-art technologies, methods, tools, or resources for cancer research or address important underexplored or unexplored areas? If the research project is successful, will it lead to truly substantial advances in the field rather than add modest increments of insight? Projects that modestly extend current lines of research will not be considered for this award. Projects that represent straightforward extensions of ongoing work, especially work traditionally funded by other mechanisms, will not be competitive.

Research Plan: Is the proposed work presented as a self-contained research project? Does the proposed research have a clearly defined hypothesis or goal that is supported by sufficient preliminary data and/or scientific rationale? Are the methods appropriate, and are potential experimental obstacles and unexpected results discussed with potential alternative approaches?

Applicant Investigator: Does the investigator(s) demonstrate the required creativity and expertise to make a significant contribution to the research? Applicants' credentials will be

evaluated by the biosketch provided and in a career stage-specific fashion. Have early-career-stage investigators received excellent training, and do their accomplishments to date offer great promise for a successful career? Has the applicant(s) devoted a sufficient amount of their time (percent effort) to this project?

Relevance: Does the proposed research have a high degree of relevance to cancer prevention research or early detection for Texans? This is a critical criterion for evaluation of applications for CPRIT support.

11.4.2. Secondary Criteria

Secondary criteria contribute to the overall score assigned to the application. Concerns with these criteria potentially question the feasibility of the proposed research.

Secondary criteria include the following:

Research Environment: Does the research team have the needed expertise, facilities, and resources to accomplish all aspects of the proposed research? Are the levels of effort of the key personnel appropriate? Is there evidence of institutional support of the research team and the project?

Vertebrate Animals and/or Human Subjects: Is the vertebrate animals and/or human subjects plan adequate and sufficiently detailed?

Budget: Is the budget appropriate for the proposed work?

Duration: Is the stated duration appropriate for the proposed work?

12. KEY DATES

RFA

RFA release January 14, 2025

Application

Online application opens February 18, 2025, 7 am central time

Application due May 6, 2025, 4 pm central time

Application review May 2025-November 2025

Award

Award notification	November 2025
Anticipated start date	December 1, 2025

13. AWARD ADMINISTRATION

Texas law requires that CPRIT grant awards be made by contract between the applicant and CPRIT. CPRIT grant awards are made to institutions or organizations, not to individuals. Award contract negotiation and execution will commence once the CPRIT Oversight Committee has approved an application for a grant award. CPRIT may require, as a condition of receiving a grant award, that the grant recipient use CPRIT's electronic Grant Management System to exchange, execute, and verify legally binding grant contract documents and grant award reports. Such use shall be in accordance with CPRIT's electronic signature policy as set forth in [chapter 701, section 701.25](#).

Texas law specifies several components that must be addressed by the award contract, including needed compliance and assurance documentation, budgetary review, progress and fiscal monitoring, and terms relating to revenue sharing and intellectual property rights. These contract provisions are specified in CPRIT's Administrative Rules, which are available at www.cprit.texas.gov. Applicants are advised to review CPRIT's administrative rules related to contractual requirements associated with CPRIT grant awards and limitations related to the use of CPRIT grant awards as set forth in [chapter 703, sections 703.10, 703.12](#).

Prior to disbursement of grant award funds, the grant recipient organization must demonstrate that it has adopted and enforces a tobacco-free workplace policy consistent with the requirements set forth in CPRIT's Administrative Rules, [chapter 703, section 703.20](#).

CPRIT requires award recipients to submit an annual progress report. These reports summarize the progress made toward the research goals and address plans for the upcoming year. In addition, fiscal reporting, human studies reporting, and vertebrate animal use reporting will be required as appropriate. Continuation of funding is contingent upon the timely receipt of these reports. Failure to provide timely and complete reports may waive reimbursement of grant award costs and may result in the termination of the award contract. Forms and instructions will be made available at www.cprit.texas.gov.

14. REQUIREMENT TO DEMONSTRATE AVAILABLE FUNDS

Texas law requires that prior to disbursement of CPRIT grant funds, the award recipient must demonstrate that it has an amount of funds equal to one-half of the CPRIT funding dedicated to the research that is the subject of the award. A grant recipient that is a public or private institution of higher education, as defined by §61.003, Texas Education Code, may credit toward the Grant Recipient's Matching Funds obligation the dollar amount equivalent to the difference between the indirect cost rate authorized by the federal government for research grants awarded to the Grant Recipient and the 5% indirect cost limit imposed by §102.203(c), Texas Health and Safety Code. Grant applicants are advised to consult CPRIT's Administrative Rules, [chapter 703, section 703.11](#), for specific requirements regarding demonstration of available funding. The demonstration of available matching funds must be made at the time the award contract is executed, and annually thereafter, not when the application is submitted.

15. CONTACT INFORMATION

15.1. Helpdesk

The Helpdesk will answer queries submitted via email within 1 business day. Helpdesk support is available for questions regarding user registration, online submission of applications as well as page limitations, formatting and correct folders to place content in CARS. Helpdesk staff cannot answer scientific questions and/or programmatic questions. Before contacting the Helpdesk, please refer to the *IFA* document, which provides a step-by-step guide on preparing an application and using CARS.

Hours of operation: Monday through Friday, 8 AM to 6 PM central time

Tel: 866-941-7146

Email: Help@CPRITGrants.org

15.2. Scientific and Programmatic Questions

Scientific and Programmatic Questions-should be directed to the CPRIT Director of Academic Research. **Before contacting CPRIT, please refer to the IFA document and contact the Helpdesk for any questions related to CARS, page limitations, formatting, etc.**

Email: research@cprit.texas.gov

Website: www.cprit.texas.gov