University of Nevada, Reno
Research & Innovation

UNIVERSITY OF NEVADA, RENO

SUBRECIPIENT INFORMATION FORM - FDP MEMBERS ONLY (SP-Sub-1.1)

Section A: Prime Award Information

Project Title: _____

Prime Sponsor: _____ InfoEd # or Prime Sponsor Award ID: _____

UNR PI: _____

UNR Period of Performance: _____ Total Proposal/Award Amount to UNR: _____

Section B: Subrecipient Information

Subrecipient Legal Name: _____

Performance Site Address (if different from organizational address):

Subrecipient PI Name: _____ E-mail Address: _____

Period of Performance: From: _____ To: _____

Subrecipient Total Funding: _____ Amount cost shared: _____ ☐ N/A

First Increment Budget Period: _____ First Increment of Funding: _____

Section C: Project Specific Information

1. ☐ Yes ☐ No Will Human Subjects be involved in Subrecipient's portion of the project?

If human subjects are involved in this project, subrecipient shall conduct the activities in accordance with the DHHS regulations codified at 45 CFR Part 46 - Protection of Human Subjects and obtain IRB approval of the planned involvement of human subjects in the project. Upon UNR's request, subrecipient will provide certification of the review and date of approval by the subrecipient's IRB. As required, subrecipient will ensure that all personnel participating in the project complete the National Institutes of Health education requirement on the protection of human subjects, addressed in NIH Notice OD-00-039.

2. ☐ Yes ☐ No Will Animal Subjects be involved win Subrecipient's portion of the project?

If animal subjects are involved in this project, subrecipient shall conduct the activities in accordance with NIH "Principles for Use of Animals", the Animal Welfare Act (7 U.S.C. 2131 et. seq.) and all other applicable Federal laws, and policies. Practices for the procurement, housing, and care of laboratory animals shall conform to NIH Guide for the Care and Use of Laboratory Animals in Research and all USDA requirements. Upon UNR's request, subrecipient will provide certification of the review and date of approval by the subrecipient's IACUC.

3. ☐ Yes ☐ No Will Recombinant DNA, Human, Plant, or Animal Pathogens or Biological Toxins be involved in subrecipient's portion of the project?

If Recombinant or Synthetic Nucleic Acid Molecules are involved in this project, subrecipient shall conduct the activities in accordance with the NIH Guidelines for Research Involving Recombinant or Synthetic Nucleic Acid Molecules. Upon UNR's request, subrecipient will provide certification of the review and date of approval by the subrecipient's IBC.

4. ☐ Yes ☐ No Under Subrecipient's planned project, does Subrecipient expect to provide or receive any information, technology, or materials that may be subject to U.S. export control regulations? If 'Yes' attach an explanation.

Section D: Certifications

1. Responsible and Ethical Conduct of Research (RECR): (applicable to projects funded by NSF or any other programs requiring Ethics in Research training; see instructions for applicability):
☐ Yes ☐ No ☐ NA My organization certifies that it has a training program in place and will train all personnel in the responsible and ethical conduct of research, in accordance with the Sponsor's program-specific requirements.

2. CHIPS and Science Act of 2022 Public Law 117 - 167: (only for U.S. Federal Projects):
☐ Yes ☐ No My organization certifies that, per Section 10634, each Covered Individual listed in the Subaward Proposal has completed research security training that meets the guidelines developed under subsection (b) of Section 10634, as required by the Federal Awarding Agency. (As of May 1, 2025, applies only to DOE awards.)
☐ Yes ☐ No My organization certifies that, per Section 10632, each Covered Individual listed in in the Subaward Proposal has certified that they are not a party to a Malign Foreign Talent Recruitment Program, as required by the Federal Awarding Agency.

3. Dual Use Research of Concern (DURC) and Pathogens with Enhanced Pandemic Potential (PEPP): (only for U.S. Federal Projects)
☐ Yes ☐ No Will your organization's portion of this project involve DURC/PEPP? (As of May 6, 2025, applies only to NIH awards) If "Yes", my organization is aware of and will comply with the U.S. Government Policy for Oversight of Dual Use Research of Concern (DURC) and Pathogens with Enhanced Pandemic Potential (PEPP), as required by the Federal Awarding Agency.

Section H: Authorized Representative Approval

The information and representations above have been read, signed, and made by the authorized official of the subrecipient named below. The appropriate programmatic and administrative personnel involved in this project are aware of prime sponsor policy in regard to subrecipient agreements and are prepared to establish the necessary inter-organizational agreements consistent with those policies.

_____ _____
Signature of Subrecipient's Authorized Official Date

_____ _____
Printed Name and Title of Authorized Official Email and Phone