

Subrecipient

will/

will not comply with this requirement.

FDP Subrecipient Pilot - Supplemental Project Information

Subrecipients **MUST** submit form prior to agreement execution with the University of California, Davis (UC Davis). It provides a checklist of documents and certifications required. Subrecipients complete this form on an annual basis for multi-year projects.

SECTION A General Information	······································
Subrecipient Name:	Subrecipient PI:
Project Title:	
Prime Sponsor:	RFP or Prime Sponsor #:
Budget Period: to	UEI Number:
SECTION B Program Specific Cor	npliance
1. Human Subjects	
a) Project involves Human Subjects Res	search: Yes No
b) Institutional Review Board Approval:	Pending Approved Exempt Not Applicable
NOTE: IRB Determination and Appro	
Date of IRB Determination/Approval:	Subrecipient Protocol Number:
c) Key Personnel have completed NIH I	Human Subjects Training requirements: Yes No
2. Animal Subjects	
a) Project involves Animal Subjects Res	search: Yes No
b) Institutional Animal Care and Use Co	mmittee Approval: Pending Approved Not Applicable
NOTE: IACUC Determination and Ap	proval MUST be submitted to UC Davis.
Date of IACUC Approval:	Subrecipient Protocol Number:
Subrecipient hereby certifies it will comport organizational, and conflicts of interest t	ndividual COI or any Organizational COI Requirements ONLY) bly with the additional standards for financial disclosure, both individual or hat are required by the Prime Sponsor and are not covered by the FDP FCOI es that, to the best of the Subrecipient's knowledge:
resulting agreement, and that are req 2) All identified conflicts of interest hat accordance with Subrecipient's conflict funds under any resulting agreement;	Sponsor's guidelines for conflict of interest reporting will be made to UC Davis
4. Cost-Sharing	
Cost Share or Matching is being provide	ed by the Subrecipient: O Yes O No Amount:
NOTE: Cost sharing MUST be detailed	and justified in the Subrecipient BUDGET.
5. NIH International Subrecipient Reporting	g Requirements (For non-U.S. Subrecipients ONLY)
the provisions of <u>NIH GPS 15.2.1</u> requir permissible) to copies of all lab noteboo	ponsor is the U.S. National Institutes of Health (NIH), Subrecipient is aware of ing that international subrecipients provide access (electronic access ks, all data, and all documentation associated with the research as described in a recipient and in alignment with progress report submission requirements, but

programmatic and administrative personnel involved in this applicatio	read, signed, and made by an authorized official of the Subrecipient. The appropriate n are aware of agency policy in regard to subawards and are prepared to establish the . Any work begun and/or experiences incurred prior to execution of an
Signature of Subrecipient Authorized Official	Date
Printed Name of Authorized Official	Title of Authorized Official
E-mail Address	Phone Number

SECTION C Subrecipient Signature