**Pilot Grant Application**

Institute for Clinical and Translational Science

Project Title:

Principal Investigator:

1. Are Human Subjects Involved? Y / N
	1. Please see IRB Guidelines statement below
2. Is this study exempt from Federal Regulations? Y / N
	1. If yes, what is the exemption number?
	2. If no, is the IRB review pending? Y / N
		1. IRB Approval Date (please attach IRB approval):
3. Clinical Trial Questionnaire
	1. Does the study involve human participants? Y / N
	2. Are the participants prospectively assigned to an intervention? Y / N
	3. Is the study designed to evaluate the effect of the intervention on the participants? Y / N
	4. Is the effect that will be evaluated a health-related biomedical or behavioral outcome? Y / N
	5. Provide the ClinicalTrials.gov identifier (e.g. NCT87654321) for this trial, if applicable:
4. Are Vertebrate Animals Used? Y / N
	1. If YES to Vertebrate Animals Is the IACUC review pending? Y / N
		1. IACUC Approval Date (please attach IACUC approval):
5. Is this a multi-site study? Y / N
6. Team Science:
	1. This proposal has potential for creating or expanding research that benefits from synergies between different groups: Y / N
	2. We agree to participate in a Seminar provided by the ICTS on Team Science should our Letter of Intent (LOI) be selected: Y / N

**IRB Guidelines**All projects involving human subjects research are required to receive IRB approval before funding is released. Some projects may be required to use a single IRB (sIRB). Some projects may also be required to undergo the NCATS human subject prior approval process. Once established, each applicant/team is required to meet all established IRB deadlines and comply with all information requests. Failure to meet deadlines or respond to information requests in a timely manner may result in the administrative removal of the non-compliant institution from the project team.