



INSTITUTIONAL REVIEW BOARD

APPLICATION FOR USE OF HUMAN SUBJECTS

Title of Proposed Study: (NOTE: Title must be consistent on all documents)

Principal Investigator (PI):

Position:

TSC Faculty

TSC Staff

Non-TSC

List all co-investigators and key personnel, their affiliation and their role in the research:

Source of funding:

Describe the purpose (research objective) of the study:

Describe the procedure (methodology) for selecting and recruiting subjects/participants:

Number of Participants (*refers to the maximum number from whom consent is obtained. Estimate an adequate number of subjects to allow for dropout and subjects who may be screened and not meet the inclusion criteria.*)

Age of Participants eligible for inclusion:

Describe the Subjects/participants:

Check any categories of vulnerable populations that may apply to proposed subjects:

Children	TSC Students	Prisoners
Pregnant Women	Decisionally Impaired*	Economically Disadvantaged
Non-English Speaking	Racial/Ethnic Minority	Educationally Disadvantaged
Students in subject pool	Students in PI's class	Other (please explain)

**Decisionally impaired refers to subjects who may not be able to provide informed consent for their own participation, due either to a permanent or temporary condition.*

If other was selected, explain:

Describe how informed consent will be obtained and by whom (or note if a waiver of consent is being requested, or any alterations in informed consent are being requested.)

Check type of procedure to obtain consent from subjects (Attach copies of any applicable consent forms):

Consent form	Parental consent form	Child assent form
Verbal (waiver of documentation of consent)*	Waiver of consent*	Other (provide explanation)

If other was selected, explain:

Will participants be deceived, misled or misinformed about any aspect of the project?

Yes

No

If yes, please describe:

Check type of proposed research:

Questionnaire/Survey/Data

Collection Instrument(s)

(attach copy)

Existing Records (attach
document explaining

records)

Telephone Survey (attach
copy of questions)

Interview (attach copy of
questions)

Public Observation

Controlled Observation

Treatment/Experimental

Other (attach document with explanation)

Describe the research procedures to which participants will be exposed:

How will the data be recorded?

Audio recorder

Video recorder

Other (explain)

If other was selected, explain

If recordings will be used, when will the recordings be destroyed?

How will the confidentiality of the data be maintained (be specific)?

Where and how will your records be stored?

Describe any incentives and compensation related to participation and how/when this will be dispersed.

Does the study involve an intervention from which the subject could potentially benefit?

(NOTE: Incentives are not considered benefits)

No

Yes

If yes, please describe:

Will project involve stress, discomfort, or physical, legal, financial, psychological, or social risk to the participants?

No

Yes

If yes, describe all potential risks related to the participant and likelihood of occurrence:

Describe any special safety procedures to be used or methods to minimize risk:

What plans have been made to deal with adverse psychological or physiological reactions or stress which might be caused or triggered by the study?

Please review all documentation before submission. An incomplete submission means a delay in review and approval.