Date __________________

HUMAN SUBJECT RESEARCH REVIEW FORM
NEW JERSEY INSTITUTE OF TECHNOLOGY
INSTITUTIONAL REVIEW BOARD (IRB) APPLICATION

Name of Principal Investigator(s) _________________________________

Principal Investigators must be staff or faculty members. Students can serve as co-principal investigators under faculty/staff supervision for expedited projects.

NJIT Address: _________________________________________________

Department: _________________________________________________

E-mail Address: _______________________________________________

NJIT Affiliation of Principal Investigators (Check all that apply):
☐ Faculty    ☐ Staff    ☐ Student    ☐ Other - Describe:

Students and doctoral candidates applying for IRB approval must submit written documentation from their faculty advisors (via e-mail) stating that research is being conducted under their supervision.

Project Title: _______________________________________________

This project will be conducted:
☐ On Campus    ☐ Off Campus (Location): _________________________________    ☐ Both

Is this research funded by outside source(s)?    ☐ Yes    ☐ No

If yes, indicate name(s) and type of funding source(s).

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<th>Name(s):</th>
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| Type: | ☐ Government (County, State or Federal) |
|       | ☐ Foundation |
|       | ☐ Corporation |
|       | ☐ Other |

Anticipated Starting Date:

Anticipated End Date:

Number of Subjects:

NOTE: All principal investigators, faculty, and students who will be interfacing with human subjects in this study must complete an online training course in the protection of human subjects. This course can be accessed by going to the US Department of Health and Human Services Office of Extramural Research. Certificates indicating course completion must be submitted with this application.

- For investigator training, please see the following URL: http://phrp.nihtraining.com/users/login.php.
- In addition, all NJIT investigators must complete the Responsible Conduct in Research training (RCR Basic Course) of the Citi Program (http://citiprogram.org).

To Principal Investigator: In addition to the questions below, please furnish copies of any questionnaires interview formats, testing instruments or other documents necessary to carry out the research. Any advertising materials used to recruit subjects must also be submitted.
The completed forms should be sent electronically to: irb@njit.edu

1. Project Title:

2. List the names and status (faculty, student, etc.) of the persons conducting the research:
   a. Principal Investigator(s):
   
   b. Other Members of Research Team:
   
   c. NJIT Faculty Advisor(s), if Student Project:

3. Describe the objectives, methods and procedures of the research project. This summary will be used to describe your project to the IRB. Use up to 2 pages, if necessary. You may also attach a copy of an abstract or full research proposal describing this work.

4. List names and institutional affiliations of research assistants, workers and students working on this project.

5. If research assistants, workers or students will be working on the project, describe their qualifications, special training and how they will be supervised.

6. What is the age of the subjects?

7. How will they be recruited?
8. Indicate any physical, psychological, social or privacy risk or pain, which may be incurred by human subjects, or any drugs medical procedures that will be used. (This includes any request for the subjects to reveal any embarrassing, sensitive, or confidential information about themselves or others.)

9. Indicate if any deception will be used, and if so, describe it in detail. Include your plans for debriefing.

10. Evaluate the risks presented in 7.
   a. Is it more than would normally be encountered in daily life?

   b. Do your procedures follow established and accepted methods in your field?

11. How will the risk be kept at a minimum? (E.g., describe how the procedures reflect respect for privacy, feeling, and dignity of subject and avoid unwarranted invasion of privacy or disregard anonymity in any way.) Also, if subjects will be asked to reveal any embarrassing, sensitive, or confidential information, how will confidentiality of the data be insured? Also include your plans for debriefing. If subjects will be placed under any physical risk, describe the appropriate medical support procedures.

12. Describe the benefits to be derived from this research, both by the subject and by the scientific community (this is especially important if research involves children).
13. Describe the means through which human subjects will be informed of their right to participate, not to participate, or withdraw at any time. Clearly describe how the subjects will be adequately informed about the procedures of the experiment so that they can make an informed decision on whether or not to participate.

14. Complete the attached Consent Form and the IRB will determine if your subjects will be at risk. This Consent Form must include the following five points: (1) The purpose of the research, (2) the procedures involved in the work, (3) the potential risks of participating, (4) the benefits of the research including the benefits for both society and the subjects, and (5) that the subjects are free to withdraw from the research at any time with no adverse consequences.

15. Provide copies of recruitment flyers or emails, questionnaires, interview formats, testing instruments or other documents to carry out the research. If questionnaires are not complete, please submit an outline of the questions to be used. You will have to submit the completed questionnaire to the IRB before the research can begin.

16. If the subjects include minor children, complete the Consent Form as prescribed in paragraph 12 for signature by parent or guardian. If the project is approved (regardless of the IRB determination concerning risk), it will be necessary that a Consent Form be secured for every minor child.

17. Attach a copy of the permission for the facility to conduct the proposed research (if other that NJIT).
CONSENT TO PARTICIPATE IN A RESEARCH STUDY

TITLE OF STUDY:

I, __________________________________________, have been asked to participate in a research study under the direction of Dr(s). _____________ (Insert name(s) of faculty or staff.)

Other professional persons who work with them as study staff may assist to act for them.

PURPOSE: (INSERT DESCRIPTION OF THE PURPOSE OF YOUR STUDY HERE.)

DURATION:

My participation in this study will last for ______________.

I have been told that my participation in this research study is important for the success of the research and that the results of this research study are expected to produce the following benefits to society and for me as a subject.

BENEFITS FOR SOCIETY AND THE SUBJECT:

I have been told that the benefits are: (INSERT DESCRIPTION OF BENEFITS TO BE DERIVED FROM THIS RESEARCH)

PROCEDURES:

I have been told that, during the course of this study, the following will occur: (INSERT EXPECTATIONS WITH REGARD TO PARTICIPATION HERE.)

PARTICIPANTS:

I will be one of about __________ participants in this study.

EXCLUSIONS:

I will inform the researcher if any of the following apply to me: (INSERT EXCLUSIONS HERE.)
RISKS/DISCOMFORTS:
I have been told that the study described above may involve the following risks and/or discomforts: (INSERT ANY RISKS HERE.)

There also may be risks and discomforts that are not yet known.

I fully recognize that there are risks that I may be exposed to by volunteering in this study which are inherent in participating in any study; I understand that I am not covered by NJIT’s insurance policy for any injury or loss I might sustain in the course of participating in the study.

CONFIDENTIALITY:
I understand confidential is not the same as anonymous. Confidential means that my name will not be disclosed if there exists a documented linkage between my identity and my responses as recorded in the research records. Every effort will be made to maintain the confidentiality of my study records. If the findings from the study are published, I will not be identified by name. My identity will remain confidential unless disclosure is required by law.

VIDEOTAPING/AUDIOTAPING: (INCLUDE ONLY IF APPLICABLE, OTHERWISE, REMOVE THIS SECTION.)
I understand that I will be video and audio taped during the course of this study. Video and audio tapes will be stored for (insert time frame; minimum 3 years) after the end of this project (enter date in parentheses). After that time, the recordings will be erased. Recordings will be stored digitally on the NJIT servers and will not be made available to anyone except investigators who are involved in this research.

PAYMENT FOR PARTICIPATION:
I have been told that I will receive $_________ compensation for my participation in this study.

RIGHT TO REFUSE OR WITHDRAW:
I understand that my participation is voluntary and I may refuse to participate, or may discontinue my participation at any time with no adverse consequence. I also understand that the investigator has the right to withdraw me from the study at any time.

INDIVIDUAL TO CONTACT:
If I have any questions about my treatment or research procedures, I understand that I should contact the principal investigator at: (INSERT CONTACT INFORMATION (I.E., MAILING ADDRESS, TELEPHONE NUMBER AND E-MAIL OF) FACULTY OR STAFF HERE.)
If I have any additional questions about my rights as a research subject, I may contact:

Office of Research  
New Jersey Institute of Technology  
323 Martin Luther King Boulevard  
Newark, NJ 07102  
(973) 642-4877  
irb@njit.edu (email is preferred)

SIGNATURE OF PARTICIPANT

I have read this entire form, or it has been read to me, and I understand it completely. All of my questions regarding this form or this study have been answered to my complete satisfaction. I agree to participate in this research study.

Participant Name

Signature

Date

SIGNATURE OF READER/TRANSLATOR IF THE PARTICIPANT DOES NOT READ ENGLISH WELL  
(Only needed if English fluency is not an exclusion criteria)

The person who has signed above, _____________________________________, does not read English well. I understand English and am fluent in (name of the language) ____________________________________________, a language the subject understands well. I have translated for the subject the entire content of this form. To the best of my knowledge, the participant understands the content of this form and has had an opportunity to ask questions regarding the consent form and the study, and these questions have been answered to the complete satisfaction of the participant (or his/her parent/legal guardian).

Reader/Translator Name

Signature

Date

SIGNATURE OF INVESTIGATOR OR RESPONSIBLE INDIVIDUAL

To the best of my knowledge, the participant named in this form has understood the entire content of the above consent form, and comprehends the study. The questions of the participant and those of his/her parent/legal guardian have been accurately answered to his/her/their complete satisfaction.
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<th><strong>Investigator’s Name</strong></th>
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DISCLOSURE OF FINANCIAL RELATIONSHIP FOR SPONSORED PROJECTS

The following form must be completed by all Principal Investigators and members of the research team, including faculty advisors if student projects. Please use a separate form for each person.

Date: ___________________________

Name (Print and SIGN): ____________________________________________________________

(ORIGINAL SCANNED, FAXED, OR HARDCOPY SIGNATURE REQUIRED)

This form shall be completed by all members of the research team.

1. Funding Source. Does the research involve financial relationships that could create potential or actual conflicts of interest?

   ☐ Yes ☐ No

   How is the research supported or financed?

   __________________________________________________________

2. Payment or Compensation for Services. Are you receiving any salary and other payment for services (e.g., compensation in the form of equipment, consulting fees; honoraria, study design; management position, independent contractor, service on advisory committees or review panels for for-profit entities, board membership of for-profit entities; seminars, lectures or teaching engagements for for-profit entities) for this research? [Do not include salaries received from federal grant funding agencies.]

   ☐ Yes ☐ No

   If Yes, note amounts with explanation of source and activities:

   __________________________________________________________

   If Yes, is this payment or compensation affected by the study outcome?

   ☐ Yes ☐ No

   If Yes, explain:

   __________________________________________________________

   Do you receive payment per participant or incentive payments?

   ☐ Yes ☐ No
If Yes, note amounts with explanation of terms.

3. **Equity (Ownership) Interests.** Do you have any or all equity interests or ownership interests (e.g., stock, stock options, and partner) in entities related to the research activity?
   - [ ] Yes
   - [ ] No

   If Yes, note amount with explanation of source:

4. **Other Financial Interests or Relationships.** Do you have any financial interests in the product, including patents, trademarks, copyrights, or licensing agreements?
   - [ ] Yes
   - [ ] No

   If Yes, note amount with explanation of source:

5. **Incentives.** Will you receive any money, gift or anything of monetary value above and beyond the actual costs of enrollment, conducting of the research, and reporting on the results, including, but not limited to, finders fees, referral fees, recruitment bonuses, enrollment bonus for reaching an accrual goal or similar types of payments?
   - [ ] Yes
   - [ ] No

   If Yes, note amount with explanation of source:

6. **Other.** Are there any other interests or relationships (including volunteer services) that might constitute a conflict of interest or an appearance of conflict of interest in connection with the research project?
   - [ ] Yes
   - [ ] No

   If Yes, note amount with explanation of source:

Please address questions or concerns to the Office of Research (irb@njit.edu)