

Institutional Review Board (IRB)

Human Subjects Research Proposal Form

Project Title:	
Anticipated: Start Date:	End Date:
Principle Investigator: Student	□ Faculty
Print Name:	Department:
Campus email:	Campus phone:
I, the Principle Investigator, have read the Gui Subjects Research Proposals for IRB Review, http://www.sjcme.edu/resources/institutional-r	available at:
Signature of Principle Investigator:	
Faculty Supervisor (Required for Student Re supervisor attests that (s)he has read this research appropriate education and supervision of the stud	h proposal and agrees to provide
Print Name:	Department:
Campus email:	Campus phone:
Signature of Faculty Supervisor:	

Part I: Training

Please complete the online training module for investigators provided by the Collaborative Institutional Training Initiative (CITI Program). CITI is "dedicated to promoting the public's trust in the research enterprise by providing high quality, peer reviewed, web-based educational courses in research, ethics, regulatory oversight, responsible conduct of research, research administration, and other topics pertinent to the interests of member organizations and individual learners."

https://about.citiprogram.org/en/homepage/

- At the website, click on Register and then under Select Your Organization Affiliation, enter: St. Joseph's College of Maine (this will auto-populate)
- Check the two boxes below (Agreement of Terms of Service and that you are an affiliate of the college)
- Create username and password
- Print the Certificate of Completion that is provided when you complete the training. Save a copy for your records and attach one copy to this proposal form. The IRB will keep the Certificate on file. If you have an existing NIH training certificate, please submit a copy of that.

Part II: Narrative

Please attach a typed narrative that provides the following information. Your narrative should clearly follow the structure and numbering provided. Each numbered section should include the title of that section in the heading (for example, the first section should begin with "1. <u>Description of Research</u>"). You can answer briefly, and include details as necessary. Remember that some reviewers are not specialists in your field; therefore, write your narrative so that an educated but naïve person will understand it.

- 1. <u>Description of Research</u>: Describe the following:
 - a. Purpose of research
 - b. Nature of the data to be collected
 - c. Data collection procedures
 - d. Data collection instruments (if surveys, etc., please attach copies)
 - e. Methods for selection/recruitment of participants
 - f. Information about participants (age, number, sex, etc.)
 - g. Incentives or compensation for participation.
- 2. <u>Risks</u>: Describe in detail any psychological, social, legal, economic, or physical risks to which participants might be exposed. Describe how subjects will be protected from these risks.
- 3. <u>Benefits</u>: Federal guidelines require that risks of participation be outweighed by potential benefits to participants and/or to humankind in general.
 - a. Describe any benefits to participants resulting from this research
 - b. Describe any benefits to humankind in general resulting from this research
- 4. <u>Informed Consent</u>: Describe the consent process to be followed in the research. If your research requires written documentation of informed consent, attach a copy of the informed consent form. If deception is to be used, explain the justification for the use of deception.
- 5. <u>Vulnerable Populations</u>: If minors or other vulnerable populations will be included as research participants, describe the procedures to be used in obtaining their agreement (assent) to participate, in addition to the consent of their authorized representative.
- 6. <u>Confidentiality</u>: Describe how the confidentiality of participant information will be maintained and how participant information will be kept secure. Note whether or not data collection will be anonymous.
- 7. <u>Dissemination</u>: Describe how results will be disseminated to the research community (e.g., publication or presentation at professional meetings) and/or to other interested groups (e.g., an on-campus poster session).

Part III: Summary Checklist

Please complete the following brief checklist. Some of these items address information that is also covered in the Narrative (Part II of this form).

- 1. Who will be participating in the study? (check all that apply):
 - □ Adults
 - □ Children / minors (individuals younger than 18 years old)
 - \Box Students
 - □ Pregnant women
 - □ Institutionalized people (e.g., prisoners)
 - □ Other; please list:
- 2. How will participants be recruited? (check all that apply):
 - □ Class announcements
 - □ Advertisement
 - □ Telephone
 - Email
 - □ Letter
 - □ Other; please list:

Please attach copies of any documents or materials that will be used for recruitment.

- 3. Will anyone under the age of 18 be participating in the study?
 - □ Yes
 - □ No
- 4. Will participant data be collected in a way that allows the Investigator to identify individual participants and link individual participants to their data?
 - □ Yes
 - \Box No, data will be anonymous (this is <u>not</u> the same thing as confidential)
- 5. Will participants be offered compensation for participation?
 - □ Yes
 - □ No
- 6. Will participants be exposed to *more than minimal risk*, as defined in the Guidelines?
 - □ Yes
 - 🛛 No
- 7. Will any participant data be collected from institutional files or archives?
 - □ Yes
 - □ No
- 8. Will any institutions besides SJC be involved in this research project?
 - \Box Yes If yes, list the institutions in the space below.
 - 🛛 No

Part IV: Signatures

1. Name and signature of Principal Investigator:

I understand that, as the Principal Investigator, it is my responsibility to ensure that all researchers involved in this project understand and comply with the applicable ethical guidelines for protecting human subjects.

Name:	
Signature:	Date:
2. Names and signatures of all	the researchers involved in this project:
Name:	
Signature:	Date:
Name:	
Signature:	Date:
Name:	
Signature:	Date:
Name:	
Signature:	Date:
Name:	
Signature:	Date: