Administrative & Limited (Exempt) Review Research Projects

Before proceeding to the application on the next page, please confirm that your research activities satisfy one of the following criteria. Otherwise, your project may require Expedited or Standard Review. ***If your project involves the use of identifiable secondary or existing data, please complete the IRB Application for Existing & Secondary Data.***

You should only fill this form out if your project involves any of the following:

* **Normal educational practices in an educational setting. The research activities cannot adversely affect students’ opportunity to learn required educational content OR adversely affect the instructors who provide instruction.**

**Example: Comparing multi-media lesson plans in Fall Semester to paper-based lesson plans in Spring semester. The same educational material is covered, only the teaching technique varies.**

* **Surveys, interviews, observation, educational test or behavioral interventions where the data collection is verbal, in writing or recorded (audio or video) and *MUST* satisfy one of the following:**
  + **Anonymous data collection (no links to identifiers such as names)**
  + **Data collected does not place subjects at risk or harm**
  + **Data that are identifiable, but protections are provided for confidentiality**

***Data collection involving children (under 18) or other vulnerable populations requires Expedited Review and submission via the IRB Application for Expedited and Standard Review.***

**Example 1:** Qualtrics survey to collect data on CSULB students’ study habits during finals for the purpose of a senior thesis; whether data are collected anonymously or confidentially.

**Example 2:** Interviews with transgender women to understand the harassment they endure in the workplace. No names, employers or company names are used. Pseudonyms used instead. Interviews will be audio recorded with permission in the consent form prior to the interview.

* **Benign behavioral interventions involving authorized deception (subjects must consent to deception prior to the research procedures). *The use of concealed deception or manipulation requires Expedited Review and submission via the IRB Application for Expedited and Standard Review*.**

**Example:** Group activity involving a white woman in scenario 1 and a black man in scenario 2 who both demonstrate an anger problem and subjects are surveyed to determine underlying gender and racial bias. Subjects were consented to participate in a puzzle competition that involved an element of deception that the PI agrees to reveal at the conclusion of the experiment.

* **Public benefit and service programs that are federally funded or conducted by the federal government.**

**Example:** An assessment of emergency room wait times in a Veterans Affairs hospital.

* **Taste and food quality tests involving ingredients the FDA determines safe for consumption.**

**Example:** A marketing taste test evaluation of a chocolate chip cookie compared to a nutrition bar flavored like a chocolate chip cookie. Participants will submit surveys to determine whether the healthy nutrition bar is a tasty substitute for real chocolate chip cookies

PLEASE DELETE THIS PAGE BEFORE SUBMISSION

IRB Application for Administrative & Limited (Exempt) Review

Projects involving less than minimal risk

**Instructions:** The Principal Investigator (PI) is responsible for all information in this form. The PI should fill out the form completely; incomplete forms will be returned. Use as much space as necessary to answer each question adequately. **Please upload the completed application and all relevant appendices to IRBNet (www.irbnet.org).**

**INITIAL SUBMISSION DOCUMENTS:** (1) IRB Application, (2) Informed Consent form(s), and (3) Appendices (relevant documents such as faculty advisor letter, permission letter(s), informed consent forms, recruitment material(s), etc.).

**IMPORTANT:** NO ACTIVITY MAY BEGIN ON THIS PROJECT UNTIL THE PRINCIPAL INVESTIGATOR HAS RECEIVED FORMAL NOTIFICATION FROM THE CSULB IRB THAT THE PROJECT HAS BEEN APPROVED BY THE IRB. In addition, the IRB will confirm that all individuals listed on this project have successfully completed the CITI online training for the Social & Behavioral Basic/Refresher course before issuing final approval ([www.citiprogram.org](http://www.citiprogram.org)).

1. **BASIC INFORMATION**

|  |  |
| --- | --- |
| **PI’s Name** (Last, First, Degree) |  |
| **Telephone Number** |  |
| **Email** |  |
| **CITI Member ID #** |  |
| **Completion of CITI Social & Behavioral Basic/Refresher Course (Check one)** | [ ] Yes [ ] No [ ] Not Sure |
| **Department** |  |
| **Affiliation** | [ ] Student\* [ ] Staff [ ] Faculty [ ] Other: |
| **\*If you are a student, please complete the information below for your Faculty Advisor:** | |
| **Faculty Advisor Name** |  |
| **Email** |  |
| **CITI Member ID #** |  |
| [ ] I have attached a signed and completed letter from my faculty advisor as an appendix to this project in IRBNet. | |

1. **PROJECT SUMMARY**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Title of Project** | | | | |
|  | | | | |
| **Describe the purpose of the project. Provide context to the importance of the research and explain how the results of this study will contribute to the field of study or specific population.** | | | | |
|  | | | | |
| **Total number of subjects anticipated:** | |  | | |
| **Describe the subject population (Select all that apply):** | | | | |
| [ ] 18+  [ ] 65+  [ ] Other age range (Explain): | [ ] All genders  [ ] Females only  [ ] Males Only  [ ] Other (Explain): | | [ ] Non-English Speakers  [ ] College Students  [ ] General public  [ ] Public officials  [ ] Agency/Institutional officials | |
| **Provide any other subject characteristics for inclusion in your project:** | | | | |
|  | | | | |
| **List all methods of subject recruitment (Select all that apply):** | | | | |
| [ ] Flyers/letters  [ ] Emails  [ ] Telephone  [ ] In-class announcement | [ ] Subject pool  [ ] Tabling in public  [ ] Personal network/Snowball sampling | | | [ ] Passive recruitment  [ ] Social media post  [ ] Other (Explain): |
| **Describe the step-by-step recruitment methods for each subject group (do not include any other project procedures in this section):** | | | | |
|  | | | | |
| **List the setting(s) of the research activities (Select all that apply):** | | | | |
| [ ] CSULB (in class)  [ ] CSULB (out of class)  [ ] CSULB (online)  [ ] Other College Campus  [ ] K-12 School  [ ] Lab | [ ] Public setting  [ ] Nonprofit or Other Agency  [ ] Government Facility  [ ] Foreign Country  [ ] Subject’s work/home  [ ] Prison/Jail | | [ ] Telephone  [ ] Email  [ ] Social media  [ ] Amazon mTurk  [ ] Online (general public)  [ ] Other: | |
| **List the location of the research activities (Agency/school names, city, county or social media site):** | | | | |
|  | | | | |
| **Do you have permission from the facility, institution or director of the organization to conduct your project? *\* If yes, please attach signed permission letter or data sharing agreement to this package****.* | | | | |
| [ ] Yes\* [ ] No [ ] In Progress | | | | |
| **Is the project primarily for training purposes?** | | | | |
| [ ] Yes [ ] No [ ] Other (Explain): | | | | |
| **List all methods of data collection (Select all that apply):** | | | | |
| [ ] Surveys/Questionnaires  [ ] Focus Groups  [ ] Educational Tests/Assessments  [ ] Accessing Public Records  [ ] Observation  [ ] Audio recording  [ ] Video recording | | [ ] One-on-One interviews  [ ] Data Mining  [ ] Digital media (videos, pictures, simulation)  [ ] Access to medical/academic records  [ ] Biospecimen Analysis  [ ] Experimental Controls  [ ] Other (Explain): | | |
| **Is there an experimental group compared to a control group?** | | | | |
| [ ] Yes [ ] No | | | | |
| **Describe the step-by-step procedures of your project beginning with the informed consent process and continuing with all other project activities (all project activities must occur *AFTER* IRB approval):** | | | | |
|  | | | | |
| **Will subjects be compensated for their participation? If monetary compensation is to be provided, please specify the amount and provide a justification for that amount that avoids potential coercion.** | | | | |
| [ ] No  [ ] Yes (Complete below):  **Mode of compensation (Check one):** [ ] Cash [ ] Gift Card [ ] Course credit [ ] Other (Explain):  **Total Amount per Subject:**  **Justification (Explain):** | | | | |

1. **RISKS AND MITIGATIONS**

|  |
| --- |
| **Describe any reasonable risks subjects may experience participating in this project (Ex. Risk #1: Loss of confidentiality, Risk #2: Discomfort answering questions, Risk #3: etc…):** |
|  |
| **Provide a reasonable mitigation to reduce each potential risk listed above. Each mitigation should directly correspond to each risk (Mitigation for Risk #1, Mitigation for Risk #2, etc.):** |
|  |

1. **DATA ACCESS**

|  |  |
| --- | --- |
| **Who will have access to the data?** |  |
| **Where will the data be stored?** |  |
| **Will identifiable data be collected (names, student ID numbers, address, etc.)?** | [ ] Yes\* [ ] No [ ] Not Sure |
| ***\*If yes*, why is it necessary for identifiable data to be collected?** |  |
| **Will you have access to external data i.e. medical charts, public records, proprietary information?** | [ ] Yes\* [ ] No |
| **\*If yes, please describe the additional data you will have access to:** | |
|  | |
| **Is it possible for the information collected to be traced back to identify the individual with subject master list or by other means?** | [ ] Yes\* [ ] No |
| \*If yes, describe the method(s) to protect subject privacy and confidentiality: | |
|  | |

1. **FUNDING**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Is the project funded?** (*If no*, skip to Section 5) | | | [ ] Yes [ ] No | |
| **Name of Funding Agency:** | |  | | |
| **Grant Title or Number:** | |  | | |
| **Funding Dates:** | Start: | | | End: |

1. **RESULTS**

|  |  |
| --- | --- |
| **Will the results of this project be published?** | |
| [ ] Yes\* [ ] No | |
| ***\*If yes*, for what purpose will the results be published?** (Check all that apply) | |
| [ ] Thesis  [ ] Dissertation  [ ] Research Journal/Article  [ ] Employee Manual  [ ] Pilot Data  [ ] Research Conference | [ ] Course/Department Improvements  [ ] Course Assignments  [ ] Evidence Based Practice  [ ] Training  [ ] Other (Explain): |
| **What knowledge base will the results contribute to?** | |
| [ ] Personal knowledge [ ] Generalizable knowledge [ ] Program Evaluation | |

1. **ADDITIONAL PERSONNEL**

|  |
| --- |
| **List the name, CITI Member ID #, and email for any additional research staff or assistants that will have access to the research data:** |
|  |
| [ ] Not Applicable |

1. **Investigator Assurance**

**As the Principal Investigator (PI) for this project, I certify the following (please check and confirm all):**

[ ] All subjects in this project will be afforded the same rights and protections.

[ ] All provisions to maintain subject confidentiality and protect subject data will be upheld.

[ ] The CITI online training is complete for me and all personnel listed. Training will remain valid for the duration of research activity.

[ ] Research activities will be conducted as described in the approved version of this application.

[ ] Any modifications, including personnel changes to this project will be submitted, reviewed, and approved by the CSULB IRB prior to implementation.

[ ] I will comply with the CSULB IRB research policies (<https://www.csulb.edu/office-of-research-and-economic-development/research-related-policies>).

[ ] I will comply with the Department of Health and Human Services (HHS) research policy and regulation (<https://www.hhs.gov/ohrp/regulations-and-policy/regulations/45-cfr-46/index.html>).

Name: Date: MM/DD/YYYY