



Cape Cod Community College

Institutional Review Board (IRB) Research Application

NOTE: All investigators need to complete the online Office for Human Research Protections Self-Assessment Training, available at [Human Research Protection Training | HHS.gov](https://www.hhs.gov/ohrt/). Completion certificates should be on file with IRB in the Office of Institutional Research. Please send completion certificates to the CCCC IRB at IRB@capecod.edu

Research Project Information

Title of Research Project:

Anticipated Funding Source:

Principal Investigator/Faculty Advisor

Department

Phone Ext.

Email Address

Co-Investigator/Student Researcher

Department

Phone Ext.

Email Address

Projected Duration of Research:

Projected Start Date:

Other Organizations/Agencies
Involved (if any):

Research Summary

Describe the purpose of the research:

Provide a detailed description of the research objectives, goals, and anticipated contributions. Clearly state the research question(s) and hypothesis, if applicable.

Participant Description:

Provide an overview of the participants involved, including demographic details (e.g. age range, gender, academic status) and any inclusion/exclusion criteria.

Recruitment Process:

Outline how participants will be recruited, specifying methods such as flyers, email invitations, classroom announcements, or social media outreach. Ensure compliance with ethical standards for voluntary participation.

Research Location:

Specify the setting where data collection will take place (e.g. classrooms, online surveys, or campus labs)

Data Collection Procedures/Methods:

Provide a step-by-step overview of the data collection methods (e.g. surveys, interviews, or experiments). Include the duration and frequency of interactions with participants.

Collection of Personal Identifiers:

State whether any personally identifiable information (e.g. names, emails, ID numbers) will be collected. If so, explain how confidentiality will be maintained.

Data Protection and Storage:

Specify the measures in place to secure participant data (e.g. encryption, password-protected files, locked storage)

Data Retention and Disposal:

Detail the duration for which the data will be retained and the methods for secure disposal.

Data Access:

Specify who will have access to the data and the conditions under which it may be shared.

Use of Data and Results Sharing:

Explain how the data will be used, including potential publications, presentations, or reports to external stakeholders.

Attachments:

1. **Informed Consent Form:**
Attach a detailed consent form outlining participant rights, the nature of the study, potential risks, and confidentiality protections.
2. **Additional Project Materials (if applicable)**
Include copies of all relevant materials (e.g. surveys, questionnaires, recruitment flyers, or brochures)

Signatures:

- **Principal Investigator/Faculty Advisor:**

Signature: _____ *Date:* _____

- **Co-Investigator/Student Researcher:**

Signature: _____ *Date:* _____