**SOCIAL & BEHAVIORAL TEMPLATE WITHOUT INSTRUCTIONS:**

Use this template to prepare a document for social and behavioral research with the information from the following sections.

If your study will ONLY involve secondary use of data and/or specimens. See **HRP-503b TEMPLATE** **SECONDARY USE PROTOCOL**.

Depending on the nature of what you are doing, some sections may not be applicable to your research. Mark N/A if it’s not applicable.

When you write a protocol, keep an electronic copy. You will need to modify this copy when making changes.

Use the Social & Behavioral protocol template WITH instructions as a guide.

Remove above instructions in red before submitting to the IRB

**PROTOCOL TITLE:**

*<Enter Information Here>*

Version Date:<Enter Information Here>

**PRINCIPAL INVESTIGATOR:**

Name <Enter Information Here>

Department <Enter Information Here>

Telephone Number <Enter Information Here>

Email Address *<*Enter Information Here>

**CO-INVESTIGATORS:**

Name <Enter Information Here>

Department <Enter Information Here>

Email Address <Enter Information Here>

**STUDENT INVESTIGATORS:**

Name <Enter Information Here>

Department <Enter Information Here>

Email Address <Enter Information Here>

Is this study is part of a dissertation or thesis:

Yes  No

Check any **applicable** boxes in the table below – you will be asked for further detail on these topics later in the protocol form:

|  |
| --- |
| International Research (check this box if you will collect data from individuals located outside the United States) List the locations: |
| Research involving external collaborators (Non-TAMU personnel).  List any external personnel and their organization: |
| This research has U.S. Federal government funding via one or more direct awards or a sub-award. Provide the source of federal support: |
| All other sources of funding: |

**1.0 Purpose of the Study:**

*<Enter Information Here>*

**2.0 Background / Literature Review / Rationale for the study:**

*<Enter Information Here>*

**3.0 Inclusion and exclusion criteria:**

*<Enter Information Here>*

**4.0 Procedures Involved:**

*<Enter Information Here>*

**5.0 Multiple sites:**

*<Enter Information Here>*

**6.0 Incomplete Disclosure or Deception:**

*<Enter Information Here>*

**7.0 Recruitment:**

*<Enter Information Here>*

**8.0 Consent Process**

*<Enter Information Here>*

**9.0 Process to Document Consent:**

*<Enter Information Here>*

**10.0 Risks to Participants:**

*<Enter Information Here>*

**11.0 Potential Benefits to Participants:**

*<Enter Information Here>*

**12.0 Financial Compensation:**

*<Enter Information Here>*

**13.0 Provisions to Protect the Privacy Interests of Participants:**

*<Enter Information Here>*

**14.0 Confidentiality and Data Management:**

*<Enter Information Here>*

**15.0 Data Monitoring Plan to Ensure the Safety of Participants:**

*<Enter Information Here>*

**16.0 Data and if applicable, Specimen Banking:**

*<Enter Information Here>*

**17.0 Qualifications to Conduct Research and Resources Available:**

*<Enter Information Here>*