

Institutional Review Board
Office of Research and Sponsored Programs
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Statement describing how data will be stored. If collected identifiers or confidential information, provide a statement describing the extent confidential records identifying subjects will be maintained and how (will identifiers be collected? Will they be stored with the data or separately? Where? For how long?).

Statement about future use of data. Indicate ~~IF IR~~ that

A) information may be used for future research studies or shared with another investigator for future research studies without additional informed consent from the subject; indicate whether identifiable information will or will not be shared
OR

B) that the subject's information, even if identifiers are removed, will not be used or distributed for future research studies.

Explicit disclosure of contact person(s) to address concerns and pertinent questions about research, research subject rights, and who to contact if a ~~researched~~ injury is sustained by the subject.

Include IRB contact information (irb@stockton.edu).

Reminder statement that participation is voluntary, refusal to participate will involve no penalty or loss of benefits to which the subject is otherwise entitled, and the subject may discontinue participation at any time without penalty or loss of benefits, to which the subject is otherwise entitled.

This checklist are based ~~the~~ requirements set forth by the [Office for Human Research Protections §46.116](#).