Stockton University

Institutional Review Board
Documentation of Full Review

Title of Project: _____

Project Director: _____

IRB Reviewer: _____ Date of Review: _____

Requested Length of Approval: _____ Granted Length of Approval: _____

Does the study include biospecimens or identifiable private information? ☐ yes ☐ no
If yes, approval can be only for 1 year and then every 4 years thereafter. If no, approval can be granted for the length of time requested.

Does this project involve a clinical trial or behavioral health intervention? ☐ yes ☐ no

If yes, has external posting been included? ☐ yes ☐ no

Reviewer decision	Approved ☐	Request Modifications ☐	Not Approved ☐

7.1 Risk Has risk to the subject been minimized?

(i) By using procedures which are consistent with sound research design and which do not unnecessarily expose subjects to risks; and (ii) whenever appropriate, by using procedures already being performed on the subjects.

Notes:

7.2 Risks to subjects are reasonable in relation to anticipated benefits, if any, to subjects, and the importance of the knowledge that may reasonably be expected to result.

7.3 Selection of subjects is equitable (particularly regarding vulnerable populations: children, prisoners, individuals with impaired decision-making, or economically or educationally disadvantaged persons).

7.4 and 7.5 Informed Consent Does the ICF:

kton/other letterhead?..... Yes