Arapahoe Community College

Institutional Review Board

## ACTIVITIES EXEMPT FROM COMMITTEE REVIEW

Research activities involving human subjects in the following categories may be exempt from review by ACC’s Institutional Review Board. The principal investigator/project director is authorized to make the first determination of eligibility for exemption; however, the College bears the responsibility for concurring in that determination based on notice provided by the principal investigator to the Institutional Review Board.

The following exemptions do NOT apply when (a) deception of subjects may be an element of the research; (b) subjects are under the age of eighteen; (c) the activity may expose the subject to discomfort or harassment beyond levels encountered in daily life; or (d) fetuses, pregnant women, human in vitro fertilization, children, or individuals involuntarily confined or detained in penal institutions are subjects of the activity.

EXCEPT FOR THE ABOVE EXCLUSIONS, the federally approved Categories of Exemption are:

1. Research conducted in established or commonly accepted educational settings involving normal educational practices, such as: (a) research on regular and special education instructional strategies; (b) research on the effectiveness of or the comparison among instructional techniques curricula, or classroom management methods.
2. Research involving the use of educational tests (cognitive, diagnostic, aptitude, achievement), survey procedures, interview procedures or observation of public behavior, unless: (a) information obtained is recorded in such a manner that human subjects can be identified, directly or through identifiers linked to the subjects; **and** (b) any disclosure of the human subjects’ responses outside the research reasonably place the subjects at risk of criminal or civil liability or be damaging to the subjects’ financial standing, employability, or reputation.
3. Research involving the use of educational tests (cognitive, diagnostic, aptitude, or achievement), survey procedures, interview procedures, or observation of public behavior that is not exempt under Category 2 if: (a) the human subjects are elected or appointed public officials, or candidates for public office, **or** (b) federal statute(s) require(s) without exception that the confidentiality of the personally identifiable information will be maintained throughout the research and thereafter.
4. Research involving the collection or study of existing data, documents, records, pathological specimens, or diagnostic specimens, if these sources are publicly available or if the information is recorded by the investigator in such a manner that subjects cannot be identified directly or through identifiers linked to the subjects.
5. Research and demonstration projects which are conducted by or subject to the approval of department or agency heads, and which are designed to study, evaluate, or otherwise examine: (a) public benefit or service programs; (b) procedures for obtaining benefits or services under those programs; (c) possible changes in or alternatives to those programs or procedures; or (d) possible changes in methods or levels of payment for benefits or services under those programs.
6. Taste and food quality evaluation and consumer acceptance studies: (a) if wholesome foods without additives are consumed, or (b) if a food is consumed that contains a food ingredient or at or below the level and for a use found to be safe, or agricultural chemical or environmental contaminant at or below the level found to be safe by the U.S. Food and Drug Administration or approved by the U.S. Environmental Protection Agency or the Food Safety and Inspection Service of the U.S. Department of Agriculture.

NOTE: Decision charts available at <https://www.hhs.gov/ohrp/regulations-and-policy/decision-charts-2018/index.html> can be useful in determining whether IRB review is needed, and what level of review. Questions about whether a research activity may be exempt from human subject’s review can be directed to IR@arapahoe.edu.

Exempting an activity from review does not absolve the investigator(s) of the activity from ensuring that the welfare of subjects in the activity is protected and that methods used, and information provided to gain subject consent are appropriate to the activity.

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| \_\_\_/\_\_\_\_/\_\_\_\_ | **Arapahoe Community College** | \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ |
| **Date Submitted** | **Institutional Review Board** | **File Number** |

**Exempt Protocol Summary Form** R-6-2025

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**Title of Research Project**

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**Principal Investigator/Project Director Institution & Department Phone Extension Email**

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**Co-investigator/Student Investigator Institution & Department Phone Extension Email**

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| **Anticipated Funding Source:** |  |

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| --- | --- | --- | --- | --- |
| **Projected Duration of Research:** |  | **months** | **Projected Starting Date:** |  |

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| --- | --- |
| **Other organizations and/or agencies, if any, involved in the study:** |  |

**Exempt under code (see definitions on page one – check one) 1**  2  3  4  5  6

**SUMMARY ABSTRACT: Please supply below a brief description of: (1) Summary of the project, involved researcher(s) and the intended benefit to ACC students or institution; (2) the participants, (3) the location(s) of the activities, (4) the procedures to be used for data collection, and (5) whether data will be confidential or anonymous, disposition of the data, who will have access to the data.**

**Attach copy of the Informed Consent Form, the instruments (i.e., questionnaires) to be used in the project.**

**Note: if no investigator(s) are affiliated with ACC, the investigator(s) must pursue a collaborating department or investigator at ACC prior to submitting this IRB request.**

**RESPONSIBILITIES OF THE PRINCIPAL INVESTIGATOR:**

* Any additions or changes in procedures in the protocol will be submitted to the IRB for written approval prior to these changes being implemented.
* Any problems connected with the use of human subjects once the project has begun must be communicated to the IRB Chair.
* The principal investigator is responsible for retaining informed consent documents for a period of three years after the project.

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| Principal Investigator Signature | |  | | Co-Investigator/Student Signature (if appropriate) | | |  |
|  | |  | |  | | |  |
| **Signature of IRB Committee Chair:** | | | | | | **Date:** \_\_/\_\_/\_\_ | | |
| **IRB Chair: Check 1 box:** | **Approved** | | **Approved with Conditions** | | **Refer to Full Committee Review** | | | |