

**Trinity Valley Community College
Institutional Review Board
Expedited Review of Research Form**

Human subject research activities involving no more than minimal risk to the subjects may be eligible for expedited review by Trinity Valley Community College's Institutional Review Board Chair. The principal investigator/project director is authorized to make the first determination of eligibility for expedited review; however, the College bears the responsibility for concurring in that determination based on information provided by the principal investigator.

Research activities eligible for expedited review:

1. Collection of: hair and nail clippings, in a nondisfiguring manner; deciduous teeth; and permanent teeth if patient care indicated a need for extraction.
2. Collection of excreta and external secretions including sweat, uncannulated saliva, placenta removed at delivery, and amniotic fluid at the time of rupture of the membrane prior to or during labor.
3. Recording of data from subjects 18 years of age or older using noninvasive procedures routinely employed in clinical practice. This includes the use of physical sensors that are applied either to the surface of the body or at a distance and do not involve input of matter or significant amounts of energy into the subject or an invasion of the subject's privacy. It also includes such procedures as weighing, testing sensory acuity, electrocardiography, electroencephalography, thermography, detection of naturally occurring radioactivity, diagnostic echography, and electroretinography. It does not include exposure to electromagnetic radiation outside the visible range (for example, X-rays, microwaves).
4. Collection of blood samples by venipuncture; in amounts not exceeding 450 milliliters in an eight-week period and no more often than two times per week, from subjects 18 years of age or older and who are in good health and not pregnant.
5. Collection of both supra- and sub gingival dental plaque and calculus, provided the procedure is not more invasive than routine prophylactic scaling of the teeth and the process is accomplished in accordance with accepted prophylactic techniques.
6. Voice recordings made for research purposes such as investigations of speech defects.
7. Moderate exercise by healthy volunteers.
8. The study of existing data, documents, records, pathological specimens, or diagnostic specimens.
9. Research on individuals, such as studies of perception, cognition, game theory, or test development where the investigator does not manipulate subjects' behavior and the research will not involve stress to subjects.
10. Research on drugs or devices for which an investigational new drug exemption or an investigational device exemption is not required.

Expedited review may also be used to review minor changes in previously approved research. Questions about whether a research activity may be appropriate for expedited review can be directed to the Chair of the Institutional Review Board.

_____/_____/_____
Date Submitted

Trinity Valley Community
College
Institutional Review Board

File Number

Expedited Review of Research Form

Title of Research Project

Principal Investigator/Project Director

Department Phone Email

Co-investigator/Student Investigator

Department Phone Email

Co-investigator/Student Investigator

Department Phone Email

Anticipated Funding Source:

Projected Duration of Research: Months Projected Starting Date:

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

Expedited Review Category (see categories on page 1—check one) 1 2 3 4 5 6 7
8 9 10

SUMMARY ABSTRACT: Please supply the following information below: BRIEF description of the participants, the location(s) of the project, the procedures to be used for data collection, 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 and the measures (questionnaires) to be used in the project.

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.

_____/_____/_____
Investigator/Project Director Signature Co-Investigator/Student Signature (if appropriate)

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|-------------------------|-----------------------------------|---|---|
| Signature of IRB Chair: | | | Date: ____/____/____ |
| IRB Chair: Check 1 box: | <input type="checkbox"/> Approved | <input type="checkbox"/> Approved with Conditions | <input type="checkbox"/> Refer to Full Committee Review |