1 PURPOSE

1. This policy establishes the definitions followed by the human research protection program.

2 REVISIONS FROM PREVIOUS VERSION

1. Revision of compliance definitions. Removal of determinations of Human Research and exempt from Non-Committee Review definition.

2. Added definition of clinical investigation.

3 POLICY

1. Allegation of Non-Compliance: An unproved assertion of Non-Compliance.

2. Children: persons who have not attained the legal age for consent to treatments or procedures involved in the research, under the applicable law of the jurisdiction in which the research will be conducted. For purposes of Department of Education sponsored research, Children are persons enrolled in research not above the elementary or secondary education level, who have not reached the age of majority as determined under state law.

3. Clinical investigation: any experiment in which a drug is administered or dispensed to, or used involving, one or more human subjects. For the purposes of this part, an experiment is any use of a drug except for the use of a marketed drug in the course of medical practice.

4. Clinical Trial: A research study in which one or more human subjects are prospectively assigned to one or more interventions (which may include placebo or other control) to evaluate the effects of those interventions on health-related biomedical or behavioral outcomes.

5. Continuing Non-Compliance: A pattern of noncompliance that indicates an inability or unwillingness to comply with the requirements of an applicable law, regulation, or institutional policy pertaining to the protection of human subjects and/or with the requirements or determinations of an IRB.

6. Designated Faculty: A faculty member trained and authorized by the IRB to approve exempt Human Research conducted by investigators or students in their department, division or school.

7. Designated Reviewer: The IRB chair or an Experienced IRB Member designated by the IRB chair to conduct Non-Committee Reviews.

8. Experienced IRB Member: An IRB member is considered experienced if the IRB chair considers the IRB member to have sufficient experience in and knowledge of conducting IRB reviews.

9. Experimental Subject: An activity, for research purposes, where there is an intervention or interaction with a living individual for the primary purpose of obtaining data regarding the effect of the intervention or interaction. Research involving experimental subjects as defined in DODI 3216.02 is a subset of research involving human participants.

10. Expiration Date: The first date that the protocol is no longer approved. The date after the end date of the approval period.


12. Human Research: Any activity that either:

   3.12.1 Is Research as Defined by DHHS and involves Human Subjects as Defined by DHHS; or

   3.12.2 Is Research as Defined by FDA and involves Human Subjects as Defined by FDA.

13. Human Subject as Defined by DHHS: A living individual about whom an investigator (whether professional or student) conducting research obtains (1) data through Intervention or Interaction with the individual, or (2) information that is both Private Information and Identifiable Information. For the purpose of this definition:

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1 The terms “Human Subject Research,” “Research Involving Human Subjects,” “Human Subject Research,” “Research Involving Human Subjects,” “Clinical Research,” “Clinical Investigation,” “Clinical Study” and similar phrases are considered to be synonyms for the term Human Research.
3.13.1 Intervention: Physical procedures by which data are gathered (for example, venipuncture) and manipulations of the subject or the subject’s environment that are performed for research purposes.

3.13.2 Interaction: Communication or interpersonal contact between investigator and subject.

3.13.3 Private Information: Information about behavior that occurs in a context in which an individual can reasonably expect that no observation or recording is taking place, and information which has been provided for specific purposes by an individual and which the individual can reasonably expect will not be made public (for example, a medical record).

3.13.4 Identifiable Information: Information that is individually identifiable (i.e., the identity of the subject is or may readily be ascertained by the investigator or associated with the information).

14. **Human Subject as Defined by FDA:** An individual who is or becomes a subject in research, either as a recipient of the test article or as a control. A subject may be either a healthy human or a patient. A human subject includes an individual on whose specimen a medical device is used.

15. **Immediate Family:** Spouse, domestic partner; and dependent children.

16. **Minimal Risk:** The probability and magnitude of harm or discomfort anticipated in the research that are not greater in and of themselves than those ordinarily encountered in daily life or during the performance of routine physical or psychological examinations or tests.

17. **Non-Committee Review:** Any of the following:
   - Reviews of non-exempt research using the expedited procedure.
   - Determinations of which subjects can continue in expired research.

18. **Non-Compliance:** Failure to comply with the requirements of an applicable law, regulation, or institutional policy pertaining to the protection of human subjects, and/or with the requirements or determinations of an IRB.

In the case of research funded or conducted by the Department of Defense (DOD), Non-Compliance includes failure of a person, group, or institution to act in accordance with Department of Defense (DOD) instruction 3216.02, its references, or applicable requirements.

19. **Institutional Official:** Vice Chancellor for Research.

20. **Prisoner:** Any individual involuntarily confined or detained in a penal institution. The term is intended to encompass individuals sentenced to such an institution under a criminal or civil statute, individuals detained in other facilities by virtue of statutes or commitment procedures which provide alternatives to criminal prosecution or incarceration in a penal institution, and individuals detained pending arraignment, trial, or sentencing.

21. **Related Financial Interest:** An individual involved in research review is automatically considered to have a related financial interest when the individual or the individual’s spouse, domestic partner, children, and dependents have any of the following interests in the sponsor, product or service being tested, or competitor of the sponsor held by the individual or the individual’s immediate family:
   - Involvement in the design, conduct, or reporting of the research.
   - Ownership interest, stock options, or other ownership interest of any value exclusive of interests in publicly-traded, diversified mutual funds.
   - Compensation of any amount in the past year or of any amount expected in the next year, excluding compensation for costs directly related to conducting research.
   - Proprietary interest including, but not limited to, a patent, trademark, copyright or licensing agreement.
   - Board or executive relationship, regardless of compensation.
   - Reimbursed or sponsored travel by an entity other than a federal, state, or local government agency, higher education institution or affiliated research institute, academic teaching hospital, or medical center.
   - Any other reason for which the individual believes that he or she cannot be independent.
22. **Related to the Research:** A financial interest is **Related to the Research** when the interest is in:
   3.22.1 A sponsor of the research;
   3.22.2 A competitor of the sponsor of the research;
   3.22.3 A product or service being tested; or
   3.22.4 A competitor of the product or service being tested.

23. **Research as Defined by DHHS:** A systematic investigation, including research development, testing and evaluation, designed to develop or contribute to generalizable knowledge.

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2 The phrase “ordinarily encountered in daily life or during the performance of routine physical or physiological examinations or tests” should not be interpreted to include the inherent risks certain categories of subjects face in their everyday life. For example, the risks imposed in research involving human subjects focused on a special population should not be evaluated against the inherent risks encountered in their environment (e.g., emergency responder, pilot, soldier in a combat zone) or having a medical condition (e.g., frequent medical tests or constant pain).
24. **Research as Defined by FDA:** Any experiment that involves a test article and one or more Human Subjects, and that meets any one of the following:
   
   3.24.1 Must meet the requirements for prior submission to the Food and Drug Administration under section 505(i) of the Federal Food, Drug, and Cosmetic Act meaning any use of a drug other than the use of an approved drug in the course of medical practice;
   
   3.24.2 Must meet the requirements for prior submission to the Food and Drug Administration under section 520(g) of the Federal Food, Drug, and Cosmetic Act meaning any activity that evaluates the safety or effectiveness of a device; OR
   
   3.24.3 Any activity the results of which are intended to be later submitted to, or held for inspection by, the Food and Drug Administration as part of an application for a research or marketing permit.

25. **Research or experimentation program or project, as defined by Department of Education, means any program or project in any research that is designed to explore or develop new or unproven teaching methods or techniques.**

26. **Restricted:** Applies to investigators who are delinquent in meeting IRB requirements.

27. **Serious Non-Compliance:** Noncompliance that adversely affects the rights or welfare of participants. For Department of Defense (DOD) research **Serious Non-Compliance** includes failure of a person, group, or institution to act in accordance with Department of Defense (DOD) Instruction 3216.02 and its references such that the failure could adversely affect the rights, safety, or welfare of a human subject; place a human subject at increased risk of harm; cause harm to a human subject; affect a human subject's willingness to participate in research; or damage or compromise the scientific integrity of research data.

28. **Suspension of IRB Approval:** An action of the IRB, IRB designee, Institutional Official, or designee of the Institutional Official to temporarily or permanently withdraw IRB approval of some or all research procedures short of a **Termination of IRB Approval.** Suspended studies remain open and are subject to continuing review.

29. **Termination of IRB Approval:** An action of the IRB, IRB designee, Institutional Official, or designee of the Institutional Official to permanently withdraw IRB approval of all research procedures. Terminated studies are permanently closed and no longer require continuing review.

30. **Unanticipated Problem Involving Risks to Subjects or Others:** An incident, experience, or outcome that meets all of the following criteria:
   
   3.30.1 unexpected (in terms of nature, severity or frequency) given (a) the research procedures that are described in the protocol-related documents, such as the IRB-approved research protocol and informed consent document; and (b) the characteristics of the subject population being studied;
   
   3.30.2 related or possibly related to participation in the research (possibly related means there is a reasonable possibility that the incident, experience, or outcome may have been caused by the procedures involved in the research); and
   
   3.30.3 suggests that the research places subjects or others at a greater risk of harm (including physical, psychological, economic, or social harm) than was previously known or recognized, even if no harm has actually occurred.

4 **RESPONSIBILITIES**
   
   1. Individuals writing policies and procedures are to indicate terms defined in this policy with a double underline.
   
   2. Individuals using policies and procedures are to consult this policy for the definitions of double underlined terms.

5 **PROCEDURE**
   
   1. None

6 **MATERIALS**
   
   1. None

7 **REFERENCES**
   
   7.1 45 CFR §46.102.
   
   7.2 21 CFR §50.3, 21 CFR §56.102, 21 CFR §312.3, 21 CFR §812.2(a), 21 CFR §812.3(p)