# Revision History

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| --- | --- | --- |
| **Requestor** | **Change(s)** | **Date** |
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|  |  |  |

Start

1. What is the title of the project?

2. Where will the project take place?

a. University of Arkansas for Medical Sciences

b. ACH/ACHRI

c. Other

**If Other**, Please describe:

Basic Details

1. What is the lay summary of the project?

* *This section can often be taken from the study description in the informed consent document.*
* *Failure to make this section understandable to non-scientists will delay approval.*

1. What is the primary intent of the project (choose one):
2. To develop or contribute to generalizable knowledge (to extend outside the local context)
3. To evaluate an existing program or practice and to use the findings within the institution to improve the practice or program (note that this aim does not preclude publication).
4. Other (please describe
5. Would this project still be conducted as proposed if there were no possibility of any form of academic recognition for the project (for example, no possibility of publication in a journal or presentation at a conference)?

Data Collection

1. Does the project involve obtaining information about living individuals?
   * 1. Please explain why or why not.
2. Does the project involve intervention or interaction with individuals?
   * 1. Please explain why or why not

* *Intervention: Physical procedures by which data are gathered (for example, venipuncture) and manipulations of the individual’s environment that are performed for the purposes of the project.*
* *Interaction: Communication or interpersonal contact.*

1. Is the information that will be obtained individually identifiable?
   * 1. Please explain why or why not.

* *Individually identifiable: Information that allows, or could allow, the investigator to ascertain the identity of individuals, or information that is associated with the identities of individuals.*

1. Is the information about behaviors that occur in contexts in which individuals can reasonably expect that no observation or recording is taking place?
   * 1. Please explain why or why not.
2. Is the information provided by individuals for specific purposes, and can individuals reasonably expect the information will not be made public?
   * 1. Please explain why or why not.
3. Check all of the following that the project will involve.
   1. Human embryonic stem cells (hESC), adult human stem cells, pluripotent cells or

somatic nuclear transplantation

* 1. Fetal tissue
  2. Biohazardous/infectious agents
  3. VA sites, resources, or personnel
  4. N/A

Staff

Submission Screen

The Human Subject Research Determination Request will be submitted to IRB for review.