

Section: HRPP 3.8

Effective Date: 01/27/2012 Replaces Policy: 10/01/2010

I. POLICY

A. Introduction

Quality Improvement (QI) activities are done to improve quality of programs, improve services, or improve the provision of medical care, customer service, etc. QI projects are usually done for internal purposes only. However, some QI projects may fall under the federal definition of human subject's research, and therefore may require IRB review.

B. Requirements

To determine whether QI activities involving human participants or individually-identifiable data must be submitted to the IRB, consider the definition of research. This policy defines when a QI project involves research and is subject to IRB review.

II. DEFINITIONS

Definitions for the following terms may be found in the HRPP Program Guide Section 1.3 – Definitions of terms:

- A. Research DHHS and FDA Definitions
- B. Human Subject DHHS and FDA Definitions

III. PROCEDURES

A. Overview of the differences between QI and Research

	Research	QI
Purpose	To test a hypothesis OR to	To assess or improve a process,
	establish clinical practice	program, or system OR to improve
	standards where none are	performance as judged by
	already accepted	established/accepted standards
Benefits	Knowledge sought may or	Knowledge sought directly benefits a
	may not benefit current	process/ program/ system, and may or
	subjects, but may benefit	may not directly benefit patients
	future patients	
Risks/Burdens	May put subjects at risk	Does not increase risk to patients, with
		exception of possible
		privacy/confidentiality concerns
Methods	Systematic data collection	Systematic data collection
Analysis	Statistically prove or disprove	Compare a program/process/system to
	hypothesis	an established set of standards, or to
		establish internal benchmarks
Result	Answer a research question	Improves or creates a program/
		process/system that results in greater
		safety, efficiency or satisfaction

B. Issues to Consider

- 1. What often distinguishes QI activities from research is whether the activities are intended or designed to develop or contribute to generalizable knowledge. For purposes of this policy, "generalizable knowledge" is information (findings) that can be applied to populations or situations beyond those being immediately studied.
- 2. If there are no intentions to develop or contribute to generalizable knowledge, IRB review is not required.
- 3. If QI activities are a systematic investigation AND will develop or contribute to generalizable knowledge, IRB review is required. It is important to note that at the onset, many QI projects have only local (organizational) improvement intentions, but during the process of data collection or

- analysis, it becomes clear that findings could be generalizable or benefit others. IRB review should occur when there is an intention to make findings generalizable.
- 4. When an IRB Chair, designee or IRB staff member cannot in all fairness decide or agree on whether a submission is research or QI, that application may be referred to the full board for discussion and vote.

C. The QI project must be submitted to the IRB if any of the following are true:

- 1. there is an intent to use the data to contribute to generalizable knowledge,
- 2. there is a random assignment of participants to compare outcomes,
- 3. the activities are not normally done as part of standard operating procedures,
- 4. results will be used to apply knowledge to other programs outside the institution,
- 5. the project is subject to peer review (designed to be used outside of the institution),
- 6. anonymity of participants cannot be assured, or
- 7. the activities involve more than minimal risk to participants.
- D. If an investigator is unsure as to whether or not the project meets, or does not meet, the definitions above, please consult with the IRB.