POLICY

Research activities that meet all applicable criteria set forth by the federal regulations (45 CFR 46.101 (b)) and involve no greater than “minimal risk” may qualify for exemption from IRB review. An exemption must be determined by the IRB, not an individual investigator. The exemption categories are posted on the IRB web site. The IRB may rule a proposal is exempt if it fits into one of the exemption categories and is no greater than minimal risk. However, the IRB may determine that there are studies that may pose ethical concerns and are reviewed as with any other new study application.

The IRB administrator is an IRB member designated by the IRB Chair to review all requests for exemption and make a final determination. Such claims of exemption are recorded and stored in the ERICA online system.

PROCEDURES

1. Exempt Research Activities

1.1. The investigator submits a new study application in ERICA.

1.2. The IRB administrator conducts a review of the research proposal using the IRB internal checklist (Determining if Human Research is Exempt from IRB review) within the ERICA online system. When one or more of the exemption categories are applicable to the research, the IRB administrator documents the applicable category(ies) using the internal checklist.

1.3. VA research determined exempt from IRB review must be reviewed by the R&D Committee prior to initiation and must be included in annual R&D Committee Review of research projects.

1.4. All exemption determinations are communicated to the investigator via the ERICA online system, and includes the applicable category(ies) justifying the exemption determination.