Human Subjects Research Change Form

Any change to an approved protocol **must** be reviewed and approved by the IRB **before** the change is implemented. Such amendments could include changes to the study design, procedures, enrollment, methods of recruitment, personnel, funding source or the consent form/information sheet. This includes changes that appear to reduce risks to subjects.

Protocol Number

Study Title

Principle Investigator (PI)

Phone

E-mail

Mailing Address

Please describe *each* proposed amendment(s) and explain *why* it is being made. This includes changes to project personnel.

Signature of Principal Investigator

Date