

Form FDA 1572 Revision Notification

I. STUDY INFORMATION	
Principal Investigator:	
Protocol Number:	
Sponsor name and/or Funding Agency:	

II. CHANGES
1. Detail Changes to Form FDA 1572:
2. Does Informed Consent Form require updating with new information? <input type="checkbox"/> Yes* <input type="checkbox"/> No *If Yes, indicate what should be updated:
3. Were new sub-investigators added to the study? <input type="checkbox"/> Yes* <input type="checkbox"/> No *If Yes, CVs and licenses are required for new sub-investigators. Submit along with this form.

A copy of the updated Form FDA 1572 is attached submitted with this notification form.

III. PERSON COMPLETING FORM

By submitting this form I certify that the information contained is complete and accurate and that no facts have been suppressed or misstated. The Principal Investigator (PI) is aware of all information in this form and I am authorized to submit to the IRB on the PI's behalf.

Form Completed By

Date

Title, Company

Phone:

Email: