



- If application is Funded/Sponsored, ask the research team to confirm that the application is still funded through their approved Sponsors/Funder.
- Review the application to make sure the application does not need a non-significant determination.
- Follow-up on conditional approvals.
- IRB of Record is AAHRPP Accredited
  - If IRB is not AAHRPP Accredited, request copies of SOP's.
- IAA is needed for projects that are funded where we don't have a standing agreement. See Deferral Cheat Sheet.
- Partial Waiver of PHI for recruitment purposes, have research team fill out the required Appendix for Alterations/Waivers of Consent or PHI.

**Review Consent Forms for:**

- If the research is being conducted at a BUMG site, the correct ICF template is being used.
- Sponsor required language for all Sponsor funded projects.
- Version dates are correct.

**Within KC**

- Check COI for individuals on the Personnel List (PI and CO-PI).
  - Update Special Review Tab to reflect COI.
  - Update Special Review Tab to Reflect any SRA/SRC/Biological/ Radiation approval.
- Make sure the *protocol type* reflects the review level associated with the application.
- Update the *Additional Information* section within the Protocol tab to add *Other Identifiers*.
- Funding source is filled out completely based on the application.
- Cover to cover copy of the grant/award is in file.
- Drugs/ Devices are listed in the Keywords, under *Additional Information*.
- Vulnerable Population matches what the current project is enrolling.
- Custom data, Location is complete.
- Update Questionnaire Tab to reflect the requested information (This information will pull into the Approval Letter.)
- Add Waiver of PHI for those projects approved with a partial waiver of PHI for recruitment/screening purposes.

Effective Date	Version #	Authors	Description
5/1/2018	001	Andi Encinas	