1. **Title of Study:**
2. **IRB Approval #:**
3. **Principal Investigator**

**Name:** **Title:**

**Phone:** **Email:**

**Campus Address:**

1. **Amendment Change Type:**

[ ]  Administrative Change [ ]  Protocol Revisions

**[ ]** Consent Form Revisions **[ ]** Other:

1. Briefly describe the modifications requested, please included a reasons for each change.

1. In your opinion, do the moditications requested increase, decrease or have no effect on the risk of harm to the subjects?

[ ]  No effect [ ]  Increase [ ]  Decrease

1. Do the modifications alter the approved consent from (If yes, please make sure to include a new version of the consent form with submission)

[ ]  Yes [ ]  No

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Signature of Principal Investigator Date