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| **Names of all personnel involved in this IRB application’s design, conduct, or reporting** |
| **Title of the research study:**  |  |
|  | **Name and Title**  | **Department/School** | **Involved in the consent process?** |
| Researcher\*: |       |       | [ ]  |
| (If applicable) UCR Faculty Advisor/ Supervisor\*: |       |       | [ ]  |
| Co - Investigator (Co-I):*Please note a Co-I is required for clinical trials* |       |       | [ ]  |
| ***All investigators and staff conducting research must complete CITI*** [***training***](https://research.ucr.edu/ori/guidance/citi-instruction.aspx)***. Investigators and staff conducting clinical trialsare required to take GCP training, either through CITI or by providing a copy of their ACRP or SoCRA certification. Required training must be renewed every 5 years.*** ***\*IRB approval cannot be issued until the researcher and advisor/supervisor (if applicable) have completed the required ethics training.***  |
| **Name (Last, First) /** **Role in study (design, conduct, reporting, other)**  | [**CITI training**](https://research.ucr.edu/ori/guidance/citi-instruction.aspx) | **UCR Tutorial or****NIH Protection Human Research Participants Course (PHRP)** | **GCP Training** | **ACRP/ SoCRA****Certified** | **Involved in consent?** |
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| **Researcher-Faculty Advisor/Supervisor Acknowledgement** |
| I attest that the personnel delegated are qualified to perform the study-related procedures assigned to them and that all conflicts of interest these individuals have with this research have been reported to the UC Riverside [Promoting Research Objectivity (PRO) Committee](https://research.ucr.edu/ori/committees/pro.aspx). I also attest that these individuals have received the training or have the qualifications indicated above. |
| UCR Faculty Researcher or Faculty Advisor signature: | Date |
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