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| **Names of all personnel involved in this protocol’s design, conduct, or reporting** | | | | | |
| **IRBNet ID #:** |  | | | | |
| **Project Title:** |  | | | | |
|  | **Name and Title** | **Institution and Department** | | | **Involved in consent?** |
| Principal Investigator: |  |  | | |  |
| Co-Principal Investigator:  *Please note a Co-PI is required for clinical trials* |  |  | | |  |
| Individuals included on the research personnel list must be UC Davis faculty, employees, students, or volunteers, or outside collaborators working under the oversight of the UC Davis Principal Investigator and covered by an [Individual Investigator Agreement](http://research.ucdavis.edu/policiescompliance/irb-admin/researchers/coll-research/). Contact the IRB if you have questions regarding research personnel. All research personnel are required to complete human subject research protection training prior to engaging in research activities. Required training is described in [HRP – 103 Investigator Manual.](http://research.ucdavis.edu/wp-content/uploads/HRP-103-INVESTIGATOR-MANUAL.pdf) | | | | | |
| **Name and Title** | | | | **Involved in consent?** | |
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| **Principal Investigator Acknowledgement** | | | | | |
| I attest that all personnel assigned to this study are qualified to perform the procedures assigned to them, have reported any and all conflicts of interest, and will complete all required training prior to engaging in research activities. | | | | | |
| Principal Investigator signature | | | Date | | |
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