Research Personnel Heat Map

Role of Research Personnel	Minimal Risk Protocol	Greater Than Minimal Risk Protocol
Access to subject identifiable data including Protected Health Information (PHI) for screening/ determining eligibility	**List only the LR and Co-Researcher(s) in the UCI IRB Application & Protocol Narrative. The LR is required to maintain a Study Team log or something similar to track Research Personnel independently.**	
Recruiting subjects directly via phone, email or in person		
Access to subject identifiable data which may include PHI for data collection purposes		
Involvement in the informed consent process (i.e., explaining the study to prospective subject)		
Interacting with subjects as part of study procedures; for greater than minimal risk research this may include more invasive procedures		**List the LR, Co-Researcher(s) and Research Personnel in the UCI IRB Application & Protocol Narrative.**
Involvement in the interpretation of study data		
Finalization of the informed consent process (i.e., able to sign off as the individual obtaining consent)		
Has a disclosable financial conflict of interest		