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 Study Team section in KRP. 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)		
Performing research procedures that involve greater than minimal risk		List the LR, Co-Researcher(s) and Research Personnel in the Study Team section in KRP.
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		