Human Participants Form (4)

Required for all research involving human participants not at a Regulated Research Institution. If at a Regulated Research Institution, use institutional approval forms for documentation of prior review and approval. (IRB approval required before recruitment or data collection.)

Student's Name(s)	Title of Project
Adult Sponsor	Phone/Email
MUST BE COMPLETED BY STUDENT RESEARCHER(S) IN COLLABORATION WITH THE ADULT SPONSOR/DIRECT SUPERVISOR/QUALIFIED SCIENTIST:	
1. 🔲 I have submitted my Research Plan/Project Summary which addresses ALL areas indicated in the Human Participants Section of the	
Research Plan/Project Summary Instructions. 2. I have attached any surveys or questionnaires I will be using in my project or other documents provided to human participants.	
Any published instrument(s) used was /were legally obtained.	
 I have attached an informed consent that I would use if required by the IRB. Yes No Are you working with a Qualified Scientist? If yes, attach the Qualified Scientist Form 2. 	
BELOW – IRB USE ONLY	
MUST be completed by Institutional Review Board (IRB) after review of the research plan. All questions must be answered for the approval to be valid. (If not approved, return paperwork to the student with instructions for modifications.)	
	lired) and the following conditions: (All 6 must be answered)
	imal Risk 🔲 More than Minimal Risk
2. Qualified Scientist (QS) Required (Form 2): 🔲 Yes	(a risk assessment form 3 is required).
3. Risk Assessment Required (Form 3):	
 4. Written Minor Assent required for minor participants: Yes No Not applicable (No minors in this study) 	
5. Written Parental Permission required for minor participants:	
Yes No Not applicable (No minors in this study)	
6. Written Informed Consent required for participants 18 years or older: Yes No Not applicable (No participants 18 yrs or older in this study) 	
IRB SIGNATURES (All 3 signatures required) None of these individuals may be the adult sponsor, direct supervisor, qualified	
scientist or related to (e.g., mother, father of) the student (conflict of interest).	
I attest that I have reviewed the student's project, that the checkboxes above have been completed to indicate the IRB determination and that I agree with the decisions above.	
Medical or Mental Health Professional (a psychologist, medical doctor, licensed social worker, licensed clinical professional counselor,	
physician's assistant, doctor of pharmacy, or registered nurse) with expertise related to this project.	
Printed Name	Degree/Professional License
Signature/Date (prior to experimentation)	Email
Educator	
Printed Name	Degree/Professional License
Signature/Date (prior to experimentation)	Email
School Administrator	
School Administrator	
Printed Name	Degree/Professional License
Signature/Date (prior to experimentation)	Email