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.)

CONTRACTOR SECTION AND SECTION AND SECTION AND SECTION ASSESSMENT OF CONTRACTOR SECTION AND SECTION AN	
Nellie + Bailey Fisher	arkinsons
	Title of Project
Adult Sponsor Must be completed by Student Researcher(s) in collaboration with the Adult Sponsor/Designated Supervisor/Qualified Scientist: 1.	
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.) Approved with Full Committee Review (3 signatures required) and the following conditions: (All 6 must be answered) 1. Risk Level (check one):	
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	PSV. D. Licensed Drychologist Degree/Professional License
Signature	Daté of Approval (Must be prior to experimentation.) (mm/dd/yy)
Christopher J. Tyler Ph.D.	
Printer Mame Poler	Degree/Professional License 9.18.19 Date of Approval (Must be prior to experimentation.) (mm/dd/yy)
School Administrator	
BRIAN LADEWIG	
Printed Name	Degree/Professional License
Blu July	Ed.D.
Signature	Date of Approval (Must be prior to experimentation.) (mm/dd/yy)