**Please note: You are to submit one Research Equipment form for each equipment system.** Researchers planning to use physical stimulation or physiological data acquisition equipment with human subjects must complete this form and *append it to their original or amended New Protocol Application, as applicable.*

**IMPORTANT: DO NOT CHANGE THE FORMAT OF THE FORM. ANY CHANGE OR MODIFICATION OF THE FORM WILL RESULT IN REJECTION OF THE APPLICATION BY THE CANDIDATE.**

**1. RESPONSIBLE PROJECT INVESTIGATOR (PI)/SUPERVISOR**

|  |
| --- |
| Full Name: |

**2. PROJECT TITLE**

|  |  |
| --- | --- |
| |  | | --- | |  | |

**3. DESCRIPTION OF EQUIPMENT** Describe the system sufficiently so that the IERB can evaluate the risks associated with its use.

Include any information relevant in a product brochure, equipment specification sheet, or other printed material from the manufacturer, please check here  and provide the material *in addition to* the following information:

Describe the equipment to be used on the human subjects or system (or attach manufacturer’s printed material, as appropriate).

Describe specifications, industry ratings, and industry standards for all stimulation, amplification, transduction, and data acquisition equipment, as applicable. Describe alterations made to commercially available equipment.

If additional information is attached, check here: