**INSTRUCTIONS** **All forms must be typewritten and submitted via email to j.bain@sonoran.edu. 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.

**1. RESPONSIBLE PROJECT INVESTIGATOR (RPI)**

|  |  |
| --- | --- |
| Last Name: | First Name: |

**2. PROJECT TITLE**

|  |
| --- |
|  |

**3. DESCRIPTION OF EQUIPMENT:**

Describe the system sufficiently so that the IRB can evaluate the risks associated with its use. If this information is included in a product brochure, specification sheet, or other printed material from the manufacturer, check here  and provide the material *in addition to* the following information:

Describe the equipment or system, including how pieces interface with each other and with the subject (or attach manufacturer’s printed material, as appropriate). Include 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: