**PURPOSE:** To ensure proper calibration of equipment utilized to collect and process clinical trial data.

**SCOPE:** Applies to all site personnel involved in the implementation and coordination of clinical research.

**PERSONNEL RESPONSIBLE:**  Principal Investigator, Sub-Investigators, Study Coordinator and/or other pertinent staff who will conduct research.

**PROCEDURES:**

* The equipment must be calibrated per manufacturer or site standards.

**RESOURCES:**

* 21 CFR 312.60- General responsibilities of Investigator

**TOOLS:**

* Manufacture equipment guidelines