**PURPOSE:** To define the procedures for management of Investigational Product(s) [IP] according to the contractual requirements of the clinical trial.

**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:**

* Please refer to the individual pharmacy’s and entity’s SOPs regarding IP management
* Please refer to OVCR website policies <http://research.wustl.edu/Pages/default.aspx>

**RESOURCES:**

* ICH GCP 4.6- Investigational Product(s)

**TOOLS:**

* Investigational Product accountability logs