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381-2476 FOR CURRENT PRICING**

ST. LUKE'S REGIONAL MEDICAL CENTER
MOUNTAIN STATES TUMOR INSTITUTE
DEPARTMENT OF PHARMACY

RESEARCH STUDY FEE SCHEDULE

Protocol Number _____ Principle Investigator _____

The following fees that are checked, will be assessed for Pharmacy Department involvement in drug research studies. Fees are based on the complexity of the study and anticipated time involvement of the Pharmacy Department management and staff.

PROTOCOL START-UP FEE

\$500 Familiarizing pharmacy with protocol and working out logistics of dispensing, RX staff education, taking pre-printed orders to the P& T Committee

PATIENT ENROLLMENT FEE

Minimal Involvement - \$100.00/patient

- Drug storage
- Record maintenance
- Drug labeling and dispensing
- Single dose dispensing

Moderate Involvement - \$150.00/patient

- Drug storage
- Record maintenance
- Sterile drug preparation
- Drug labeling and dispensing
- Multiple dose dispensing

Complex Involvement - \$200.00/patient

- Drug storage
- Record maintenance
- Sterile drug preparation
- Drug labeling and dispensing
- Multiple dose dispensing
- Drug randomization/record keeping – additional \$50/patient
- In-patient & out-patient dispensing

St. Luke's Department of Pharmacy responsibilities include, but are not limited to the following:

1. Review the study protocol to determine the feasibility of pharmacy department participation in the study.
2. Receive and account for each shipment of study drug.
3. Provide in-service education to the pharmacy staff in consultation and cooperation with the sponsor.
4. Meet with study sponsors and investigators.
5. Provide for storage of the drug.
6. Perform an inventory of drug stock and reconcile all discrepancies.
7. Arrange for return of the drug at the completion of the study.
8. Provide copies of the drug accountability records to investigators and sponsors.
9. Provide necessary copies of all records at study close as required by the protocols and sponsor contacts.
10. Store the records in accordance with the study protocol.

11. Serve as a Drug Information resource for protocol medications for other St. Luke's staff.

Responsibilities of the physician's research coordinator include, but are not limited to the following:

1. Provide a copy of study protocol to St. Luke's Pharmacy study coordinator(s).
2. Meet with St. Luke's Pharmacy study coordinator(s) prior to implementation of the study and as necessary thereafter.
3. Develop pre-printed drug study order form in accordance with St. Luke's Pharmacy and Nursing guidelines.
4. Assist in acquisition of study drug as necessary.
5. As appropriate, transmit completed study drug pre-printed orders to the appropriate St. Luke's departments in a timely manner.
6. Keep St. Luke's study coordinator(s) informed of any protocol changes.
7. Provide a list of approved sub-investigators and any changes to St. Luke's study coordinator(s).
8. Notify St. Luke's at the time patient enrollment is concluded.

Jim Francis, RPh, MS Clinical Director of Pharmacy

Date

Principal Investigator

Date