Conducting Clinical Research Course
Wednesday, October 9, 2013
8:30 AM - 2:30 PM
3.104A – Pestaña Lecture Hall/Medical Bldg.
Videoconferencing to: RAHC-Room 2.120 and
Laredo Campus-Room1.106

Agenda:

- Investigator Responsibility & Safety Data Monitoring
- The Informed Consent Process
- Study Documentation-it’s All About the Data
- Investigational Product Accountability
- Training, Qualifications, Delegation of Authority
- Unanticipated Problems Involving Risk to Subjects or Others (UPIRSO)
- Working with Affiliate Hospital Partners

Purpose

Conducting Clinical Research is intended to be a practical course useful to faculty, research nurses and coordinators who interact with human subjects in the many ongoing research projects at the Health Science Center and associated institutions. The course is designed to address the responsibilities of the Principal Investigator and the research team members in the conduct of human subject research. Focus is given to meeting regulatory and HSC requirements to promote best practices in human subject research while protecting the rights, safety and welfare of our research participants.

The structure of the course is intended to fully convey the regulated environment of human subject research while outlining operational procedures which maintain compliance.

This Course is designed to meet the educational requirement of HOP 7.2.3 Research Scope of Practice for Study Personnel. Although not required, Faculty, ACGME Residents/Fellows are encouraged to attend.

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Sponsored by:
The Office of Clinical Research

PROPOSED VISION STATEMENT:
To be the “One”...
that is, the #1 clinical research institution in South Texas and beyond.
To be “Collaborative” ...
with the investigative teams, sponsors, departments and affiliates.
To be a “Resource” ...
for investigators, coordinators, regulatory specialists, compliance officers, and industry sponsors.

MISSION STATEMENT:
The OCR is committed to supporting the research community in the conduct of ethical clinical research and human subject protection in compliance with the required federal, state, and local regulations. The office provides consultation on questions related to regulations and standards of practice governing human research. The OCR provides education and training; develops policy and processes; and leads quality improvement efforts in areas of clinical trial management. It endeavors to create an interdisciplinary collaborative environment among the research teams, fostering best practices.

Lunch will be provided!

Training materials will be available for printing via the Knowledge Center approximately one week prior to the course date.

Please access instructions for registration at the OCR website http://research.uthscsa.edu/ocr/training.shtml