Regulatory Training

Training takes place over two half-days on consecutive weeks. This training utilizes International Conference on Harmonization Good Clinical Practice (ICH-GCP) as the gold standard for the conduct of clinical research and will cover key topics and concepts to ensure human/clinical research professionals at BMC and BU Medical Campus are able to successfully conduct research study activities. The completion of both sessions of this in-person training fulfills the GCP training requirement for 3 years.

This ICH E6 GCP Investigator Site Training meets the Minimum Criteria for ICH GCP Investigator Site Personnel Training identified by TransCelerate BioPharma as necessary to enable mutual recognition of GCP training among trial sponsors.

Target Audience:

- Clinical Research Professionals (clinical research coordinators, research nurses, research project managers, and research assistants, etc.) with less than one year of experience;
- Recent hires to BMC and BU Medical Campus (regardless of duration of experience);
- Experienced clinical research support staff interested in a review of fundamental concepts related to the conduct of human research studies.

Learn More here