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Clinical Research Handbook. This Handbook was originally developed by the University of Washington's School of Medicine in order to ensure a quick and efficient start-up process for industry-sponsored clinical trials. Over time, this Handbook has evolved to present practical information not only about the start-up process of clinical trials, but also about other information relating to clinical research.

Clinical Research Handbook - ITHS
Purpose of Handbook The purpose of the Chesapeake Institutional Review Board (Chesapeake IRB) Handbook is to orient Principal Investigators/research staff, sponsors, contract research organizations (CROs), and site management organizations (SMOs), to Chesapeake IRB's policies, procedures, and guidelines.

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The Investigator Handbook published by NCI is a helpful reference for investigators conducting NCI-funded trials. 16 The Clinical Trials Support Unit is also a helpful resource for individuals conducting NCI-sponsored phase III clinical trials. 17

Clinical Investigator Responsibilities
This handbook is issued as an adjunct to WHO's "Guidelines for good clinical practice (GCP) for trials on pharmaceutical products" (1995), and is intended to assist national regulatory authorities, sponsors, investigators and ethics committees in implementing GCP for industry-sponsored, government-sponsored, institution-sponsored, or inves-

HANDBOOK FOR GOOD CLINICAL RESEARCH PRACTICE (GCP)
compliance program, BIMO, clinical investigator, IRB, sponsor, monitor, GLP, bioequivalence, inspection

Bioresearch Monitoring Program (BIMO) | FDA
This handbook outlines the responsibilities of the Principal Investigator and should be read by the key personnel on the research team. We look forward to working with you to ensure the safeguarding of the rights, privacy and welfare of those who volunteer to participate in research studies.