

# Download Conducting Gcp Compliant Clinical Research A Practical Guide

Whitehall Training specialises in providing clients with Good Clinical Practice Training. We offer affordable GCP, Pharmacovigilance and other courses around the world. The availability of high quality biological specimens is of utmost importance for cancer clinical trials and research. Standardizing methods for collection, long-term storage, retrieval and distribution of specimens across collection sites is essential to ensure the quality of the samples and enable consistent analysis of DNA, RNA and proteins. A step-by-step guide on how you can become a clinical research associate (CRA), clinical research coordinator (CRC) without two years of experience. The Clinical Program Manager will be responsible for developing, managing and implementing single and multi-site clinical trials. This position is located in Emeryville and reports to the Director, Clinical Operations or VP, Clinical Operations.