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The Good Clinical Practice (GCP) course is designed to prepare research staff in the conduct of clinical trials with human participants. The 12 modules included in the course are based on ICH GCP Principles and the Code of Federal Regulations (CFR) for clinical research trials in the U.S. The course is self-paced and takes approximately six hours to complete.

Good Clinical Practice (GCP) is an international ethical and scientific quality standard for designing, conducting, recording and reporting clinical trials that involve the participation of human subjects. Clinical Trials of Investigational Medicinal Products (CTIMPs) and GCP

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In addition, these Regulations integrate the principles of Good Clinical Practices (GCP) as described by the International Conference on Harmonization (ICH). The inspection of clinical trials will be initiated in close collaboration with the Therapeutic Products Directorate (TPD) and the Biologics and Genetic Therapies Directorate (BGTD).

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~~Good Clinical Practice (GCP) | NIHR~~

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~~HANDBOOK FOR GOOD CLINICAL RESEARCH PRACTICE (GCP)~~

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