Study Staff

Pre-Study

Consent Development
Coordinator Training
Good Clinical Practice
Pre-Study Budgeting
Recruitment Strategy
ResearchConnect
WebIRB Submissions

Regulatory

21 CFR Part 11
Clinicaltrials.gov
Data & Safety Monitoring
IND Development
IDE Development
Regulatory Binder
Safety Reporting
SOPs

Study Management

Data Management
Research Billing
Specimen Management
Study Management
Subject Enrollment Logs
Accountability Logs
Quality Management
Regulatory Binder Logs

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- **Group 1**
  - Clinical Research Information Systems
  - Clinical Research Business Partners
- **Group 2**
  - Office of Research Administration
  - Jonsson Comprehensive Cancer Center
- **Group 3**
  - Office of Human Subjects Protection
  - CareConnect Website

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