Design Study

Learn About Feasibility and Regulatory Requirements

- Cohort Finding & Feasibility
- Evaluate Study Feasibility
- IRB Approval
- Regulatory Support and Resources
- Protocol Development
- Consent Development
- FDA Device (IDE) Submission
- FDA Drug or Biologics (IND) Submission
- Scientific Review
- Explore Funding Opportunities & Budget Preparation
- Identify Funding Opportunities
- Proposal Preparation and Submission
- Data Management Planning
- Set Up Research Budgets
- Overview of Coverage Analysis
- Billing Codes and Research Pricing Information
- Seek Study Design Assistance & Resources
- Request a CTSI Ethics Consultation
- Research Data Management Best Practices (Consultation)
- Review Project for HIPAA, Privacy & IT Security Requirements
- Biostatistics - Information and Consultation
- Study Design – Information and Consultation
- Clinical Data Related to Research - Electronic Health Record
- Clinical Research Services (CRS)
- Research Data Management Tools

Can't find what you need?
Contact ResearchGo
310-206-0360
Find a Mentor

Pre-Study Management

Last updated: 16 Nov 2016

- 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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