**Design Study**

**Learn About Feasibility and Regulatory Requirements**

- **Cohort Finding**
- **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**
- **CTSI Integrating Special Populations Program (ISP)**
  - 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**

Can't find what you need? [Contact ResearchGo](https://www.researchgo.ucla.edu)
Clinical Research Services (CRS)
Research Data Management Tools
Find a Mentor
Pre-Study Management

Last updated: 25 Oct 2018

• 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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