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