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
Clinical Research Services (CRS)

Research Data Management Tools

Find a Mentor

Pre-Study Management

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