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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