**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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