**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**