Design Study

Learn About Feasiblity and Regulatory Requirements

Cohort Finding & Feasibility

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 (for MCC Studies)

Explore Funding Opportunites & Budget Preparation

Identify Funding Opportunities

Proposal Preparation and Submission

Data Management Planning

Set Up Research Budgets

Overview of Coverage Analysis

Billing & Coding Compliance

ACTRI Recharge Rates

Seek Study Design Assistance & Resources

UCSD Research Ethics Program

Research Data Management Best Practices

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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