Set Up Study

Pre-Study Setup

**Oncore - CRMS**

**Proposal Preparation and Submission**

**Set Up Research Budgets**

**Overview of Coverage Analysis**

**Billing Codes and Research Pricing Information**

**Clinical Trials & Industry Research Contracting**

**CareConnect/ResearchConnect**

**Clinical Labs at UCLA**

**Anatomic Pathology**

**Biostatistics**

**Study Management**

Initiate Regulatory Tasks

**WebIRB Submission**

Submit Application to Embryonic Stem Cell Research Oversight (ESCRO) Committee

Submit Application for Animal Subjects Research

Submit Applications for Studies Requiring Biosafety & Radiation Approvals

Investigational New Drug (IND) Submission to FDA

Investigational Device Exemption (IDE) Submission to FDA

Prepare Regulatory Binder

**Radiation Safety**

**CTSI Partner Information**

Complete These Tasks Before Enrolling Study Participants

Register Study on ClinicalTrials.Gov

Radiology Set up
Pathology Research Portal

Investigational Pharmacy Application

Investigational Pharmacy Set Up

Investigational New Drug (IND) Approval from FDA

Investigational Device Exemption (IDE) Approval from FDA

Last updated: 21 Oct 2020

- Group 1
  - Clinical Research Information Systems
  - Clinical Research Business Partners
- Group 2
  - Office of Research Administration
  - Jonsson Comprehensive Cancer Center
- Group 3
  - Office of Human Subjects Protection
  - CareConnect Website

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