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: 7 Mar 2019

• Group 1
  o Clinical Research Information Systems
  o Clinical Research Business Partners

• Group 2
  o Office of Research Administration
  o Jonsson Comprehensive Cancer Center

• Group 3
  o Office of Human Subjects Protection
  o CareConnect Website

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