

[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
 - [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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