

Department of Medicine – Clinical Trial Program

## New/Amended Study Notice from Study Team to Fund Manager

### Study Details (after site selection)

Principal Investigator:

Sub-Investigator(s):

Type of Study:  PI-Initiated       Sponsor-InitiatedFunding Type:  For-Profit       Non-Profit (due date/deadline, if applicable: \_\_\_\_\_)

Protocol Number:

IND Number:  
(for regulatory use)

Protocol Title:

**Attachments:**

- |             |  |  |
|-------------|--|--|
| 1. Protocol | <input type="checkbox"/> Final Version | <input type="checkbox"/> Draft (PI initiated ONLY) |
| 2. ICF      | <input type="checkbox"/> Draft         | <input type="checkbox"/> Pending                   |
| 3. Contract | <input type="checkbox"/> Draft         | <input type="checkbox"/> Pending                   |
| 4. Budget   | <input type="checkbox"/> Draft         | <input type="checkbox"/> Pending                   |

Anticipated # of Patients at UCLA site:

Study Duration (# of years):

IRB Number: \_\_\_\_\_ or  PendingNCT Number: \_\_\_\_\_ or  Pending

## Sponsor Information

Name:

Address:

Contact Name:

Contact Role/Title (if known):

Responsible for:     Contract     Budget     Contract & Budget

Contact Email:

Contact Phone:

## CRO Information (if applicable)

Name:

Address:

Contact Name:

Contact Role/Title (if known):

Responsible for:     Contract     Budget     Contract & Budget

Contact Email:

Contact Phone:

## Notes/Comments:

### Additional resources:

- [CT Application Checklist](#) (minimum documents/forms needed)
- [DOM CTP - Service Menu and Application](#)