Clinical Lab Services for Research Studies

Overview:

This document describes the process for working with the Stanford Clinical Lab group when lab processing is required as part of a clinical research study conducted through Stanford Children’s Health (SCH) or Stanford Health Care (SHC). For questions or to suggest additional content, please contact the Clinical Research Support Office (CRSO) at crso@stanfordchildrens.org

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I. Request a Feasibility Assessment for Research Studies with Clinical Labs

WHY:
To determine if Clinical Labs can support your study.

As a general guidance, a feasibility assessment is required except when:

- All study labs are routine and/or available on the dropdown menu of the budgeting/billing workbook (Ex: CBC, MetC, Urinalysis)
If you are still not sure if a feasibility assessment is required, please contact Clinical Labs (Ester Bengil & Kristine Ubungen) via email. You can also contact the Clinical Research Support Office (CRSO) for assistance.

**WHEN:**
Prior to submitting the study workbook to Research Management Group (RMG) ([http://med.stanford.edu/spectrum/b1_researcher_resources/b1_2_forms_templates.html](http://med.stanford.edu/spectrum/b1_researcher_resources/b1_2_forms_templates.html)), you should request a feasibility assessment from Clinical Labs to ensure that they can meet the laboratory needs of the study. If you expect the study start-up process to take more than 3 months, please be sure to let Clinical Labs know.

**HOW:** Complete the following step:

1. **Initiate an e-mail communication with Clinical Labs** (Ester Bengil & Kristine Ubungen). The subject line of the e-mail should be: “eProtocol xxxxx: Study Feasibility”
   
   In your e-mail, **include:**
   
   - **Section of Lab Manual that contains lab collection, processing and shipping instructions (not the whole manual).**
     
     This should include tube types, temperature requirements for storage and transport, maximum time to process or ship, centrifuge instructions, etc.

   **Example:**

   **Specimen Collection Procedures**

<table>
<thead>
<tr>
<th>TESTS</th>
<th>VISITS</th>
<th>COLLECT</th>
<th>RETURN</th>
</tr>
</thead>
<tbody>
<tr>
<td>PLASMA BIOMARKER</td>
<td>Screening Day</td>
<td>1 x 1 mL lavender top EDTA tube</td>
<td>2 mL</td>
</tr>
<tr>
<td></td>
<td>Study Treatment Discontinuation</td>
<td>1 x cryovial</td>
<td>TO:</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>FROZEN -70°C OR LOWER WEEKLY</td>
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</table>

   If Clinical Labs has questions regarding the study labs and the CRC cannot answer these questions, the CRC will need to reach out to the Sponsor to provide clarification. Clinical Labs personnel do not contact Sponsors/monitors.
Whether or not your study may require after hours or weekend sample processing.

After hours processing requires notifications and additional documentation.

Clinical Labs will respond to the feasibility assessment by e-mail within 2 weeks and indicate if study is feasible and, if so, whether or not after hours/weekend processing will be available. Please note: Clinical Labs can only store research samples up to 3 months.

II. Submit the ‘Request for Clinical Lab Requisition Form’

WHY:
A completed ‘Request for Clinical Lab Requisition Form’ is required by Clinical Labs in order to create a Clinical Lab Requisition Form. The Clinical Lab Requisition Form allows Clinical Labs to collect and process research samples and charge them appropriately. This ensures that patients are not charged for study-related activities. DO NOT create your own requisition form!

WHEN:
The ‘Request for Clinical Lab Requisition Form’ should be submitted to Clinical Labs as soon as all of the following items are obtained and before the first patient is enrolled:

- IRB approval letter
- Finalized Budget/Billing workbook including active hospital account number (see HOW section below)
- Executed contract (where applicable)

HOW: Complete the following step:
1. Initiate an e-mail communication with Clinical Labs (Ester Bengil & Kristine Ubungen).
The subject line of the e-mail should be: “eProtocol xxxxx: New Study Submission”
In your e-mail, include:

- Complete the ‘Request for Clinical Lab Requisition Form’ - click here to obtain the form. The Request for Clinical Lab Requisition Form requires an active hospital account number or account name
  - To obtain an account name or account number, you must submit the final workbook to Patient Financial Services (PFS) for their approval. Once approved, PFS will provide the account name/account number. PFS will require a PTA#. Workbook templates and other instructions can be found by here (Under Study Team Resources: Budget & Billing)
  - All studies require a workbook be submitted to PFS, even “No bill” studies or those using discarded specimens from routine labs.
Example of the ‘Request for Clinical Lab Requisition Form’:

- **IRB approval Letter**
- **Section of Final Lab Manual that contains specimen collection, processing and shipping instructions (not the whole manual)**
  - If there have been any changes to the specimen collection, processing or shipping instructions since the feasibility assessment, please indicate what those changes are.
  - Please do not send the entire lab manual to Clinical Labs
- **Copy of all feasibility assessment e-mail communications with Clinical Labs.**

Clinical Labs may initiate a meeting with the CRC if required. Approval takes approximately 2 weeks.

III. Obtain the Lab Requisition Form

**WHY:**
The Lab Requisition Form will be used as an order for Clinical Labs to collect and/or process the samples. Clinical Labs cannot draw and/or collect, accept, or process research samples without the lab requisition form.
**WHEN:**
Before enrolling first patient.

**HOW:**
Once Step II (above) is completed, lab personnel will create the Lab Requisition Form and provide an electronic version to the CRC. Enrollment can then begin.

**IV. Conduct the Study with Clinical Labs**

**WHY:**
To ensure that all study visits run smoothly and Clinical Labs has everything they need to process and collect samples in a timely manner.

**WHEN:**
When study is open to enrollment.

**HOW:**
Complete the following steps for each patient visit:

1. Complete the Lab Requisition Form and notify Clinical Labs at least 24 hours prior to the study visit.
   a. This requisition should accompany the specimens. Clinical Labs requires the paper requisition form for each sample processing, even if orders have also been placed in EPIC.
2. Provide pre-labeled tubes and shipping materials for each patient visit.
   a. Please note the lab does not provide any study supplies.
3. Provide a copy of the specific processing instructions for each particular visit, not the entire laboratory manual.
4. If your visit is taking place outside the Clinical Lab usual business hours of 7:30am to 6:00pm Monday through Friday or on the weekends, provide items above to Clinical Labs in advance of the visit, and include information about the anticipated collection time(s).

**V. Study Renewals**

Lab Requisition Forms expire after one year and need to be renewed annually along with the IRB approval letter. Please note that some minimal risk studies may not have an IRB expiration date or may be approved by the IRB for a period longer than 3 years. These studies still require annual renewal with Clinical Labs even if IRB renewal is not required annually. Complete the following step **within one week after IRB approval of continuing review:**
1. Initiate an e-mail communication with Clinical Labs (Ester Bengil & Kristine Ubungen). The subject line of the e-mail should be: “eProtocol xxxxx: IRB Annual Renewal”

In your e-mail, include:

- IRB annual renewal approval letter
- Expired lab requisition form

Clinical Labs will then update the Lab Requisition Form within 2 weeks.

VI. Close the Study with Clinical Labs

**WHY:**
To ensure that Clinical Labs is aware they no longer need to support the study.

**WHEN:**
The study team should notify Clinical Labs that the study is closing when their services are no longer required.

**HOW:**
Complete the following steps:

1. Email Clinical Labs (Ester Bengil and Kristine Ubungen) to inform them no further lab services will be required for the study. The subject line of the e-mail should be: “eProtocol xxxxx: Study Closeout”
2. If you have frozen samples, please set up an appointment with Clinical Labs to coordinate shipping the samples. Clinical Labs needs at least 24 hours’ notice to ship any samples. Clinical Labs can store research samples up to 3 months.
Appendix

Clinical Labs Locations and Hours

- **Clinical Labs:** Monday – Friday: 7:30am – 5 pm

- **SCH (Pediatrics):**
  730 Welch – Mary Johnson Building (M-F 7:30 am – 5:30 pm – no holidays)
  725 Welch – LPCH Main (M-F 7:00 am – 5:30 pm; Saturday: 8:00 am – 4:30 pm, with 24 hours dispatch available)

- **SHC (Adults):**
  Blake Wilbur (M-F 7:30 am – 5:30 pm – no holidays)
  300 Pasteur Drive – Stanford Hospital – A101- Ambulatory clinic (M-F 7:30 am – 5:30 pm; Saturday: 6 am – 3:30 pm – (no holidays)

**Do not drop off samples with Clinical Labs.**

Contact Information

Clinical Labs:
- Ester Bengil: EBengil@stanfordhealthcare.org
- Kristine Ubungen: kubungen@stanfordhealthcare.org

When sending e-mails to Clinical Labs, please send the e-mail to both Ester Bengil and Kristine Ubungen

Send-Out Labs: (650) 725-5623

Clinical Research Support Office (CRSO):
- crso@stanfordchildrens.org
- Alyson Falwell
  Clinical Research Operations Manager
  Stanford Children’s Health
  Phone: (650) 304-7118
  AFalwell@stanfordchildrens.org

Patient Financial Services (PFS):
- SHC : Jacqueline Barajas @ JBarajas@stanfordhealthcare.org
- SCH: Sydney Piaia @ SPiaia@stanfordchildrens.org
  This team processes study financial workbooks to generate account numbers used in the Lab Requisition forms.
Leftover Samples

For access to discarded leftover samples from routine clinical labs, please contact Clinical Labs (Ester Bengil & Kristine Ubungen) via email. In order to access leftover clinical samples for research, the research study must be set up with Clinical Labs by following the standard set up process.

Other: Forms and Policy Updates

- Clinical Laboratory Documentation Request (Under Study Team Resources: Forms: Request & Requisitions / Lab Forms)
- New Clinical Laboratory Policy - Pediatric maximum blood draw volumes for clinical trials (5/24/19)

Acronyms and Definitions

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<thead>
<tr>
<th>Acronym</th>
<th>Definition</th>
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<tbody>
<tr>
<td>SCH</td>
<td>Stanford Children’s Health</td>
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<tr>
<td>LPCH</td>
<td>Lucile Packard Children’s Hospital</td>
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<tr>
<td>SHC</td>
<td>Stanford Health Care</td>
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<tr>
<td>CRC</td>
<td>Clinical Research Coordinator</td>
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<td>CRSO</td>
<td>Clinical Research Support Office</td>
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<td>PFS</td>
<td>Patient Financial Services</td>
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<tr>
<td>PTA</td>
<td>Project Task Award</td>
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<tr>
<td>RMG</td>
<td>Research Management Group</td>
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