

Clinical Trials Manual – Edmonton Zone
Formal Quote Guidelines
Effective Date: August 31, 2014

Cross Cancer Institute Laboratory Guidelines
Document Number: RCTGNR00014CCI
Version: 1.2

Formal Quote Guidelines

NOTE: All Laboratory set-ups will be subject to a Lab Set Up Fee (\$350.00) upon submission.

To establish laboratory services for a clinical trial and/or research project at the **Cross Cancer Institute**:

Step	Action	Additional Information
1.	<p>Please complete the Formal Quote Form.</p> <p>The Formal Quote Form must be completed in full and all necessary documentation must be included before we will proceed with Operational Approval and Formal Quote.</p> <p>Forms with incomplete information will be returned to sender.</p>	<p>See NACTRC AHS Laboratory Services / Cross Cancer Institute / Formal Quotes for Budget Preparation and Laboratory Setup</p> <p>Example of completed form can be found at:</p> <ul style="list-style-type: none"> See NACTRC AHS Laboratory Services / Cross Cancer Institute / Formal Quotes for Budget Preparation and Laboratory Setup <p>Note: Refer to Regional Laboratory Services Guide To Laboratory Services for specific test names and test code(s). Website: http://www4.albertahealthservices.ca/labservices/mmenu.asp</p> <p>If the test is not listed in the Guide To Laboratory Services, provide the complete test name on the Formal Quote Form.</p>
2.	<p>Submit the following information into the NACTRC system, ensuring Laboratory Services is selected:</p> <ul style="list-style-type: none"> Formal Quote Form Study Protocol Laboratory Manual (if available) Ethics Approval <p>If there is additional information or information was obtained after initial submission, email: Bertha.Woynorowski@albertahealthservices.ca</p> <p>Or submit a hard copy to: Clinical Trials Coordinator, Laboratory Services Room 1491 Laboratory Medicine Attn: Bertha Woynorowski Cross Cancer Institute 11560 University Ave. Edmonton, AB T6G 1Z2</p>	<p>Submissions are subject to a \$350.00 set-up fee. NACTRC does not subsidize this fee.</p> <p>The Set-up Fee will be invoiced once billing information is provided.</p> <p>Receipt Date in Laboratory will be recorded as the date that all the necessary information was received.</p> <p>Phone inquiries/price quote requests will not be accepted.</p>

*Printed copies are **UNCONTROLLED** unless signed by an authorized lab personnel below.*

(Authorized individuals are: lab personnel designated in their zone/program or provincial role to produce print copies)

Initials:

Site:

Date Printed:

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3.	Upon review, the Clinical Trials Coordinator will approve or deny the request in the NACTRC system.	If required, the Clinical Trials Coordinator will contact the Study Coordinators for additional information. If no response within 90 days, the study will be declined.
4.	If approved, a quote will be prepared and provided to the Study RN/Coordinator within 10 working days of receipt of all required information. <i>Alberta Health Services, Laboratory Services will review pricing every year and post new prices by April 1st of each year.</i>	For STAT requests (less than 10 working days), an Administration Fee of \$500.00 payable in advance is required. The administration fee is non-refundable and must be received before the STAT request is processed.
5.	A written response to the price quote is required and must be returned to the Clinical Trials Coordinator. The Study RN/Coordinator should review the formal quote for any errors or modifications.	The last page of the quote includes a section for a signature to indicate acceptance of the quote. The signed quote is returned.
6.	Draft requisitions, specific to the Study, will be created and forwarded to the Study RN/Coordinator. The Study RN/Coordinator should review the requisitions for any errors or modifications. The Study RN/Coordinator must notify the Clinical Trials Coordinator that the draft requisitions are acceptable.	Study RN/Coordinator must consult with Clinical Trials Coordinator to discuss required changes.
7.	Once all NACTRC approvals are obtained and ethics approval and billing account information is provided , the Clinical Trials Coordinator will assign a Laboratory Control Number and print master requisitions.	Please specify if you want the master requisitions left for pick-up in laboratory or sent by interoffice mail.

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