

Ontario Tumour Bank Sample and Data Application

Study Title						
Principal Investigator Information						
Salutation Given Name Surname						
Department and Institution						
Address City						
Province/State Postal/Zip Code Country						
Office Phone Office Fax Email						
I will attach a current CV in a standard scientific grant format (e.g., CIHR, NCIC, NIH)						
Laboratory Shipping Address						
Shipping Contact Room Number						
Department and Institution						
Address						
Province/State Postal/Zip Code Country						
Lab Phone Lab Fax Email						
Billing Information						
Same as shipping address Same as Principal Investigator address As below:						
Billing Contact						
Department and Institution						
Address City						
Province/State Postal/Zip Code Country						
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Courier o	Courier charges will be added to your invoice unless the following information is provided:						
O FedE	x OPurolator Shipping Account Number						
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	Please review the following statements and indicate your agreement to the conditions listed. In addition, provide a full signature and date on page 7 of this application.						
🔷 Agree	The Biological Materials, herein referring to blood plasma, buffy coat, tumour tissue, normal tissue, stained or unstained slides, wax curls and accompanying clinical data from human subjects provided by the Ontario Institute for Cancer Research (OICR), shall be used in the manner described on this Application Form. Any change in the project direction must be communicated in writing to and approved by OICR prior to implementation. OICR reserves the right to reject any changes.						
🔷 Agree	"Licensed Field of Use" means any use of the Biological Materials in the manner described on the Application Form. The Licensed Field of Use specifically excludes any use of Biological Materials in any <i>in vivo</i> use. Researcher shall not use the Biological Materials or any constituents or progeny thereof for transplantation purposes or for the genetic cloning of the donor of the Biological Materials.						
🔿 Agree	Neither the Biological Materials nor extracts from the Biological Materials shall be incorporated into any product that is intended for use in humans.						
🔷 Agree	The Biological Materials must not be sold, shared, distributed, or otherwise transferred (for consideration or otherwise) to third parties, including any other Researcher within the Researcher's organization, and may not be taken with the Researcher to another institution or company without specific prior written authorization of OICR.						
🔷 Agree	The Researcher understands and acknowledges that the identity of the donor of any Biological Materials, whether living or deceased, is confidential, and that the Researcher shall not attempt to establish the identity of a donor of any Biological Materials, but may relay any clinically significant findings to OICR.						
Agree	The screening of donors for the presence of such pathogens as HIV, tuberculosis, or Hepatitis B is not performed by OICR. The Researcher and its employees and assignees shall treat all Biological Materials as if they are contaminated and potentially infectious. The Researcher will ensure that its employees or others who on its behalf handle Biological Materials supplied by OICR are aware of the hazards and risks involved in handling Biological Materials. The Researcher will ensure that all necessary safety procedures and practices are in place and will ensure that its employees and all others with access to the Biological Materials will comply with all safety requirements necessary for their well being, including the use of universal precautions. Without prejudice to the provisions of aforesaid, OICR accepts no liability for harm caused to the Researcher's employees or others who handle the Biological Materials supplied by OICR.						
🔷 Agree	OICR will send links of digitized slide images of samples to Researchers before Biological Materials are shipped. It will be the Researcher's responsibility to use these slide images to make an informed decision about which Biological Materials to request from OICR.						
🔷 Agree	The Researcher is responsible for ensuring that all of his/her Research Organization's policies and guidelines are adhered to in using the requested Biological Materials for research.						
🔿 Agree	The researcher will ensure that appropriate physical and electronic measures are taken to ensure the security and confidentiality of the tissues and data. For example: restricted access and security of the tissue storage area, encryption of and restricted access to data, password protected computers, secure destruction of data, etc.						
🔿 Agree	Upon termination of the Material Transfer Agreement, the researcher agrees to promptly destroy any remaining Biological Materials in a safe and secure manner.						
Agree	The researcher agrees to acknowledge the contributions of OICR in all publications and presentations of studies using Biological Materials received from OICR as "Biological Materials were provided by the Ontario Tumour Bank, which is funded by the Ontario Institute for Cancer Research" and agrees to provide a copy to OICR.						
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Sample Request							
Disease	Disease detail Number of cases						
$\bigcirc \bigcirc $	h Frozen Samples Not required Fresh Frozen Tumour Fresh Frozen Normal Adjacent Fresh Frozen Plasma Fresh Frozen Buffy Coat	Number of vials required per case (250 mg (250 mg (1 mL/vi (250 µL/vi ed if more than one	y/vial) () Other sample type (describe here): vial)				
	Paraffin Embedded Sample Not required Paraffin Embedded Tumour Paraffin Embedded Normal Adjac Other (OTB pre-approval required Describe other:	ent i) Addit Wax Section	 Type of staining Charged Slides OR Uncharged Slides Baking temperature 37°C Baking time 60°C minutes Additional preparation details 				



Clinical data required

Please check all that apply:

- O Donor demographic information (e.g., age, sex, vital status)
- O Histology and diagnosis details (e.g., histologic type, stage, grade)
- Sample collection details
- O Patient history (e.g., prior cancers, history of smoking, risk factors)
- O Family history of cancer
- Surgery (e.g., procedure types and dates)
- Radiotherapy (e.g., intent, start and end dates, dose)
- O Systemic therapy (e.g., intent, start and end dates, regimen and agent details)
- Toxicities relating to treatment
- Outcome / Follow-up (e.g., progression/recurrence status, disease-free period)

Other:



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		Study D	Petails		
Time frame:	Proposed start date Duration of project				
		(months)			
	tific or REB approval? redite the OTB ethics	YesNo	If yes, please specify source *please attach supporting documentation		
	ient funding for the alysis of the requested	YesNo	If yes, please sp Approval date Grant number	Decify source	
	nalysis of tissue and/or d be and explain how secur			tained. If no, check here: 🚫 No	
f yes, please descrik	be and explain how secur	ity and confidential	ity of data will be main	tained. If no, check here: No	
f yes, please descrik	of other individuals who	ity and confidential	ity of data will be main		
f yes, please describ lease provide a list Name eason for	of other individuals who	vity and confidential	ity of data will be main	ason for them to have such access:	
f yes, please describ lease provide a list Name eason for	of other individuals who	vity and confidential	ity of data will be main	ason for them to have such access:	
f yes, please descrik lease provide a list Name leason for ccess	of other individuals who	vity and confidential will have access to c	ity of data will be main	ason for them to have such access: Institution	
f yes, please descrik lease provide a list Name Reason for ccess	of other individuals who	vity and confidential will have access to c	ity of data will be main	ason for them to have such access: Institution	



Research Proposal

Maximum 2 pages. Please include:

Objectives and significance of project

- Explain and justify
 - o The need for tissue;
 - o Inclusion/exclusion criteria for tissue (include criteria based on gender, age, etc.);
 - o The number of samples;
 - o Clinical data required (e.g., family history, staging chemotherapeutic treatments, pathology reports, etc.);
 - o Any potential conflicts of interest in this study involving any of the investigators.

A brief description of methods

Types of assays, markers to be measured, etc.;

• In the processing of data, will there be linkages to other databases? If yes, please describe the other databases, and comment on the possibility of identification of the source.

Please type your research proposal here. If you need additional space please use the form provided on the next page. If you are sending your research proposal separately, please indicate you are doing so in the form below.



I have read and ce	ertify that all the information provided in this	s form, together with any other	information that I may provide, is					
I have read and certify that all the information provided in this form, together with any other information that I may provide, is true and accurate to the best of my knowledge. I agree that a signed Material Transfer Agreement between OICR and my Research Organization is in place prior to the release of any tissue and data.								
Signature		Date:						
т	here are two options for submission of th	is form and your CV to the On	tario Tumour Bank:					
1. Email a copy of this form (without signature) and your CV to the Ontario Tumour Bank by pressing the Email Form button to the left (you can attach your CV after pressing the button). Please then fax the signed signature page to 416-977-7446, attention OTB Client Coordinator.								
OR 2. Print a copy of this form by pressing the Print Form button to the left. Please sign the printed copy and fax it, along with your CV, to the Ontario Tumour Bank at 416-977-7446, attention OTB Client Coordinator.								
Please save a copy of this form for your own records.								