



**Oncology Scientific Review  
Committee (OSRC)  
Clinical Trials Application Form**

**Introductory Information**

<b>Principal Investigator</b>	
<b>Address</b>	
<b>Email address</b>	
<b>Telephone number(s)</b>	
<b>Protocol Number</b>	
<b>Protocol Title (complete)</b>	

**Study Category (√)**

Yes No

<input type="checkbox"/>	<input type="checkbox"/>	Does this study involve <b>treatment</b> of patients with cancer?
<input type="checkbox"/>	<input type="checkbox"/>	Does this study involve <b>prevention of</b> cancer?
<input type="checkbox"/>	<input type="checkbox"/>	Does this study involve <b>detection</b> of cancer?
<input type="checkbox"/>	<input type="checkbox"/>	Does this study involve <b>cancer care delivery</b> ?

**Additional Information (√)**

Yes No

<input type="checkbox"/>	<input type="checkbox"/>	Is this part of a large cooperative group trial (SWOG, NRG, Alliance, etc)?
<input type="checkbox"/>	<input type="checkbox"/>	Is this part of a multi-center pharmaceutical sponsored trial?
<input type="checkbox"/>	<input type="checkbox"/>	Is this a TriHealth Investigator initiated trial?
<input type="checkbox"/>	<input type="checkbox"/>	Is this a retrospective study?
<input type="checkbox"/>	<input type="checkbox"/>	Does the study require collection of fresh tissue or tissue banking?
<input type="checkbox"/>	<input type="checkbox"/>	Does the study involve gene therapy?
<input type="checkbox"/>	<input type="checkbox"/>	Does the study include any quality of life components?
<input type="checkbox"/>	<input type="checkbox"/>	Does the study include any payment or stipend to subjects?

**National Clinical Trials 8-digit (NCT) Number: NCT\_\_**

**Primary site(s) of disease to be studied (√):**

Anus	Lip, Oral Cavity, Pharynx	Soft tissue
Bone	Liver	Stomach
Breast	Lung	Other:
Cervix	Ovary	Other:
Colon	Pancreas	Liquid Tumors
Endometrium	Peritoneal, primary	Leukemia: Sub-type-
Esophagus	Prostate	Lymphoma: Sub-type-
Fallopian Tube	Renal/kidney	MDS
Larynx	Sm. Intestine	Multiple Myeloma

**Duration: What is the expected duration of the study?**

- a. Enrollment period
- b. Data collection period

**Enrollment:**

- a. What is the expected **global enrollment** of the study?
- b. What is the expected **enrollment at TriHealth site(s)**?

**Will any satellite location(s) be used?**

Yes  No

If yes, list all locations:

\_\_\_\_\_

**What is the evidence for expected accrual estimated on this application?**

\_\_\_\_\_



**Protocol Submission Process:**

Oncology related clinical trials performed at TriHealth are reviewed for scientific merit and feasibility by the OSRC. Submission guidelines are outlined below:

1. Complete OSRC Application Form and transmit completed application to the OSRC Coordinator
2. Preferred method of transmission is e-mail to OSRC Coordinator at [yunmi\\_kwon@trihealth.com](mailto:yunmi_kwon@trihealth.com)  
Alternately, completed application can be faxed to 513-852-8719
3. You will receive an e-mail acknowledgment receipt of the application
4. After OSRC has reviewed the proposed study, and all pending issues have been resolved, an e-mail will be sent indicating approval or denial of your protocol submission

*Please contact Yun Mi Kwon at 513-853-1319 for any questions regarding your submission.*