**Canadian Cancer Clinical Trials Network**

**Network Portfolio**

**APPLICATION FORM**

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| A trial is eligible for the Network Portfolio if it meets all of the following eligibility criteria. Definitions are available on <https://3ctn.ca/for-researchers/trial-portfolio/>. For assistance in completing the form, contact [3CTN CC](mailto:info@oicr.on.ca).  Select all that apply: | |
| **Oncology interventional clinical trial** | **Funded independently of 3CTN** |
| **Academic sponsored** | **Multi-centered** |
| **\*External Peer reviewed by:** Enter peer reviewed by | |
| To apply for Portfolio status, complete the form and submit to [info@3ctn.ca](mailto:info@3ctn.ca) for review. All fields are mandatory. Please provide details of peer review in the form below. | |

**Date:** Enter a date.

**Section 1: Trial Information**

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| Official Title: Enter official title. | | | |
| Lay Summary: Enter lay summary (maximum of 350 words). | | | |
| NCT#/Study Registration #: Enter study registration #. | | | Study Acronym/Short Name: Enter study acronym. |
| Phase: Select one. | Trial Status: Select one. | | Date study open by sponsor to recruitment: Enter date. |
| Study Chair: Enter study chair name. | | | Intervention Type: Select one. |
| Sponsor: Enter academic sponsor. | | | |
| Is the Sponsor the Clinical Trials Application (CTA) holder? Select one. | | If no, CTA holder: Enter CTA holder. | |
| Open to pediatric patients? Select one. | | Is the study open to additional Canadian sites that are not listed below a) and b)? Select one. | |

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| List all Canadian sites that are: | |
| 1. Open to Recruitment (include dates if known):   List all sites open to recruitment. | 1. Planning to open to recruitment (include site status if known): List all sites planning to open to recruitment. |

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| Study Features (select all that apply): | | | | | | |
| Quality of Life | | Cost Effectiveness/ Economic Analysis | | | Biospecimen Collection | |
| Disease Site: | | | | | | |
|  | Brain/CNS | Breast | Gastrointestinal | Genito-Urinary | | Gynecological |
|  | Head and Neck | Hematology | Lung | Sarcoma | | Skin/Melanoma |
|  | Other: Describe. | | | | | |
| Anticipated length of treatment (longest arm): Select one. | | | | | | |
| Total study time (per patient; from enrollment to study exit, including follow up period): Select one. | | | | | | |
| Correlative science/lab/imaging (check all that may apply):  None  Blood and archival tumour at baseline  Imaging at baseline  Anything more than baseline collection (i.e., repeated blood or imaging)  Repeated tissue biopsies (i.e., on or after treatment)  Pharmacokinetics, pharmacodynamics | | | | | | |
| Does this trial have multiple stages for registrations/randomizations? Select one. | | | | | | |
| If yes, list the steps towards patient accrual:  List steps towards patient accrual. | | | | | | |
| Has the study drug received FDA approval?  Yes  No  N/A | | | | | | |
| \*Peer review:  Has the study previously failed peer review?  Yes  No  For peer review definitions, go to <https://3ctn.ca/for-researchers/trial-portfolio/>  Enter additional details of the peer review (e.g. type and group). | | | | | | |
| Comments:  Enter additional comments. | | | | | | |

**Section 2: Applicant Information**

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| Name: Enter applicant name. | Title: Enter applicant title. |
| Organization: Enter applicant organization. | Email: Enter applicant email. |

Save the form and submit to [info@3ctn.ca](mailto:info@3ctn.ca) for review.