**[Insert SHORT TITLE] SYNOPSIS**

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| --- | --- | --- |
| **APPLICANT** | *Provide the name, email address of the applicant, institution.* |  |
| **STUDY TITLE** | *Provide the entire study title.* |  |
| **SHORT TITLE** | *Provide the short title given to the study.* |  |
| **INDICATION** | *pecify the medical condition, disease, or area of research the study addresses.* |  |
| **TARGET STUDY POPULATION** | *Describe the demographic group or specific patient population the study aims to include.* |  |
| **STUDY TYPE/PHASE** | *Specify the study phase (e.g. Pilot phase, Phase I to IV).* |  |
| **STUDY DESIGN** | *Provide a summary of the study methodology, including type, procedures, interventions, and expected workflow. A schematic diagram can be included.* |  |
| **STUDY BACKGROUND & RATIONALE** | *Explain the historical context and importance of the study. Justify why the project is needed and how it contributes to advancements in the field or benefits the community.* |  |
| **OBJECTIVES & OUTCOMES** | *Clearly define the primary and secondary objectives of the study, as well as expected outcomes.* |  |
| **NUMBER OF SUBJECTS** | *Estimate the total number of participants required and categorize them by relevant factors such as age groups.* |  |
| **INCLUSION CRITERIA** | *List the specific characteristics participants must meet to be eligible for the study.* |  |
| **EXCLUSION CRITERIA** | *List the specific conditions or characteristics that would prevent participation in the study.* |  |
| **ETHICAL CONSIDERATIONS** | *Describe ethical concerns, how informed consent will be obtained (if applicable), and measures to protect participants' rights and well-being.* |  |
| **METHODOLOGY** | *Outline the approach for data collection, intervention (if applicable), and analysis techniques.* |  |
| **STATISTICAL ANALYSES** | *Describe briefly the sample size calculation and the statistical analyses which will be performed with the data collected.* |  |
| **TRANSLATIONAL RESEARCH(ES)** | *(if applicable) Describe briefly the translational research(-es) foreseen in the study, the rationale and the biological samples to be collected.* |  |
| **STUDY WORKLPLAN** | *Provide a timeline for project completion, detailing key milestones.*   |  |  |  |  |  | | --- | --- | --- | --- | --- | | ***Task*** | ***Expected Timelines*** |  |  |  | | *Data Collection* | *XX months* |  |  |  | | *Analysis & Interpretation* | *XX months* |  |  |  | | *Report Drafting* | *XX months* |  |  |  | | *Dissemination* | *XX months* |  |  |  | |  |
| **BUDGET ESTIMATION** | *Provide an estimated budget, detailing major cost categories such as personnel, equipment, and operational expenses.* |  |
| **FUNDING AND MATERIAL SUPPORT** | *Specify the available funding sources, sponsors, and any material support provided for the study.* |  |
| **REFERENCES** | *List any relevant literature, studies, or guidelines that support the study proposal.* |  |

To apply, please submit the following documents via email to [Oncodistinct.CTC@hubruxelles.be](mailto:Oncodistinct.CTC@hubruxelles.be) by **June 1, 2025**

* Synopsis using the **Oncodistinct-Grant-Application-Template**
* A **budget forecast**
* A **project workplan**