

Application Form:

- Departmental Research Committee Approval
- Pharmacy Sheet (If Applicable)
- Budget Sheet Details (if Sponsored or includes a non-routine test or procedure)
- Biological Hazards Section (If Applicable)
- To include in the detailed description box:

Introduction may include background information related to the research topic (Importance of the topic), the purpose in carrying out this research and the Importance of potential (expected) findings.

Methodology may include:

- 1. List of collaborating Centers and the coordinating center.*
- 2. Duration of the study*
- 3. Inclusion Criteria and Exclusion Criteria, which will be used in selecting the research participants*
- 4. Registration (If Applicable)*
- 5. Randomization Process*
- 6. Data gathering methods*
- 7. Procedures, Designated Central Laboratories (If Applicable)*
- 8. Follow-up (If Applicable)*
- 9. Safety and Efficacy Parameters*
- 10. Expected Outcome*
- 11. Sample Size*
- 12. Statistical Methods*
- 13. participant confidentiality*
- 14. References/ Literature Review Section Minimum 5 (Preferable recent ones)*
- 15. List of all investigators' Work Plan and Responsibilities*

Consent Form in word format:

- English Version
- Arabic Version
- Version Number and Date
- Contact Persons (Section J)
- Others: _____

Related Documents:

- Nursing Research Approval (for Nursing research project - jbeer@kfshrc.edu.sa)
- Ethical Approval Letters from Collaborating institution.
- Letter of Invitation to the KFSHRC-J Principal Investigator to Participate in study (from Research Owner or Collaborating Institution)
- Collaborative/Clinical Trial / Material transfer Agreement(s) with Collaborating /Funding Institution (To discuss with Sponsor Research Section MCD: 40530 – Email: lalsalmi@kfshrc.edu.sa)
- Copy of original protocol and other related documents e.g. consents, CRFs...etc. from collaborating institution
- Case Report Form (CRF) / Data Collection Sheet (including date and version number) must be validated if translated from another language
- CV of Principal Investigator (PI) & Co- Principal Investigator
- GCP Certificates of all Investigators including the PI from the collaborative center. Please find link: <https://gcp.nidatraining.org/>
- Copy of two (2) major literatures/references articles mentioned in the Literature Review Section
- Subject Recruitment Advertisements/Information (If Applicable)
- Invitation Letter to Participants if the study is questionnaire, interview, or survey (If Applicable)
- Questionnaire (including date and version Number) (If Applicable)
- Interview Questions (including date and version Number) (If Applicable)
- Bio-Medical Engineering Department Clearance If a medical device will be you used.

- Pathology and Laboratory Department clearance on the application If a lab technician will be assigned, or the study includes non-routine blood works.
- Pharmacy Department approval for medication interventions on pharmacy sheet.
- Deposit the IRB Funds Allocation (S.R 7,000) before the initial review for sponsoring research in the Research Centre account (If Applicable)
- Establishment of special Research Clinic to meet the patients (If Applicable)
- *For any study that involves sending biological samples, e.g., urine, blood, tissue, etc., outside the Kingdom of Saudi Arabia, you need to complete a form to notify the Saudi National committee of Medical and Bioethics and sign a Material Transfer Agreement before sending out any sample. (bioethics.kacst.edu.sa/intro_2_e.aspx)*
- *For any clinical trial sponsored by other than a Saudi Government Agency, you need to ensure that the sponsor has registered the trial and obtained the approval of SFDA, as required, before commencing the study. (sfda.gov.sa/Ar/home/Topis/regulations)*
- *For Biostatistician Assistance please contact (Ext. 66218) email- J1513915@kfshrc.edu.sa*
- *For Clinical Research Coordinator Assistance or IRB Clinic Services please fill attached form and send through email to (HawazinA@kfshrc.edu.sa)*

Thank you.

Office of Research Affairs

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