**Please use this application if you are applying for a new research project involving human subjects that require Full or Expedited Review. *In order to check if your project might be Exempt from IRB review, please check the Exempt Criteria detailed in the IRB Protocol Exempt Application.***

**The Document Submission Checklist below provides a general guide regarding the required documents, as applicable, to be submitted with this application. For any assistance, please contact the IRB Office at** **irb@lau.edu.lb**

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| ***1. RESEARCH PROJECT INFORMATION*** |
| Name of Principal Investigator (PI):       |  |
| Research Project / Study Title:       |
| Location (s) where the study will be conducted:       *(identify all locations where the study will be conducted under your supervision)*  |
| Will data be collected at sites/locations other than LAU or its affiliates [ ]  Yes [ ]  NoIf yes, please provide necessary approvals, as applicable |
|  ***Study Administrator/Coordinator*** *(if different from PI and responsible for IRB submissions)* |
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| Name:        |  Email:        |  Phone:        | Phone #:         |

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|  ***Document submission Checklist — Initial Protocol Submission****One* ***Electronic Copy*** *of each of the following items are required, as applicable (study documents in Word editable form):**[ ]  Completed and signed* ***Research Project Submission Form*** *(GSR) – Only for Faculty Research Projects**[ ]  Completed* ***IRB Initial Protocol Application*** *– Signed by Advisor, if Submission is for a Student Project**[ ]  A detailed protocol or research project proposal**[ ]  LAU Medical Center-Rizk Hospital Signature page - For research to be conducted at the hospital**[ ]  Grant Application / study contract including detailed budget information* *[ ]  Informed consent form(s) including any short forms & translations**[ ]  Recruitment materials (cover letters, brochures, emails/referrals, advertisements including translations**[ ]  Research Instruments e.g. diaries, surveys, questionnaires, Quality of Life handouts, telephone scripts, etc.* *[ ]  Educational materials including information sheets, study guides**[ ]  Study material including case report form and data collection tools**[ ]  Human Subject Protection training certificate for all study personnel, valid within 3 years from date on certificate (*[*Protecting Human Subject Research Participants*](http://gsr.lau.edu.lb/irb/education-training.php)*)**[ ]  Curricular Vitae for the Principal Investigator and any Co-investigator involved in participant contact**[ ]  Investigator Financial Disclosure Form completed and signed for all PI and other investigators (only for funded/sponsored studies)* |

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| ***2. Type of Research*** *(check the one that applies from each column)* |
| [ ]  **Departmental Research***(not externally sponsored)*[ ]  **Graduate Study Research***(Graduate thesis/dissertation)*[ ]  **Undergraduate Research***(Senior thesis/independent study)*[ ]  **Externally Sponsored Research** |  **[ ]  *Epidemiological/ Observational Study*** *(including social/behavioral)***[ ]  *Other*** *(please specify)*:       |

| ***3. Study Site Personnel*** *(All personnel involved in the study including the* ***Principal Investigator/Co-Investigators*** *must complete the CITI Protection of Human Subjects as a condition for IRB Approval. Use the supplementary Study Personnel sheet, if necessary.)* ***NOTE: Include copies of each human subject research protection training certificate of completion.*** |
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| Study Personnel | Participant Interaction | ObtainsInformed Consent | Conducts data analysis, reviews medical records/ databases and/or handles biological specimens |
| Name:      School / Division      Name of Human Subject Research Training:        | [ ]  Yes [ ]  No | [ ]  Yes [ ]  No | [ ]  Yes [ ]  No |
| Name:      School / Division      Name of Human Subject Research Training:       | [ ]  Yes [ ]  No | [ ]  Yes [ ]  No | [ ]  Yes [ ]  No |
| Name:      School / Division      Name of Human Subject Research Training:       | [ ]  Yes [ ]  No | [ ]  Yes [ ]  No | [ ]  Yes [ ]  No |
| Name:      School / Division      Name of Human Subject Research Training:       | [ ]  Yes [ ]  No | [ ]  Yes [ ]  No | [ ]  Yes [ ]  No |
| Name:      School / Division      Name of Human Subject Research Training:       | [ ]  Yes [ ]  No | [ ]  Yes [ ]  No | [ ]  Yes [ ]  No |
| ***4. Research Protocol Summary***  |
| *This section should be written in lay terminology and must cover the following:** *Study Rationale*
* *Purpose and Objective*
* *Study population inclusion/exclusion criteria*
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| ***5. Human Participant Recruitment***  |
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| ***A. Number of Participants*** |
| i) Expected number of participants **that will be enrolled** under your supervision:      ii) For multi-centered trial, the **total number** of participants to be enrolled (including (i) above):       |
| ***Note: Please note that the expected number of participants noted above to be enrolled under your supervision will be for which the LAU IRB approval will be granted.***  |
| ***B. Targeted Population*** |
| **Age Range of Participants** *(check all that apply)*[ ]  Newborns/Infants (0-2yrs) [ ]  Children (2-7yrs) [ ]  Children (8-17yrs) [ ]  Adults (18 yrs+), Age range :      yrs to      yrs)  | **Gender**[ ]  Females/Males [ ]  Females only [ ]  Males only |
| **Type of Participants** *(check all that apply)* |
| [ ]  Inpatients[ ]  Outpatients [ ]  Healthy volunteers, Non LAU  | [ ]  LAU Students [ ]  LAU Faculty [ ]  LAU Staff  | [ ]  LAUMC-RH Staff[ ]  Other *(please specify)*:        |
| **Special Populations** (*indicate any special populations that will be targeted in the research)* |
| [ ]  Children *(<18 yrs incl. viable neonates)\** [ ]  Nonviable neonates [ ]  Fetus/Fetal tissue [ ]  Other *(please specify):*        | [ ]  Pregnant women [ ]  Impaired decision making [ ]  Illiterate [ ]  Employees of participating sites |
| *Please provide the rationale for using the special population(s) noted above:*      |
| ***C. Recruitment Methods*** *(check all that apply):* |
| [ ]  Brochures[ ]  Recruitment letter [ ]  Study personnel [ ]  Internet / Email | [ ]  Flyers/Posters[ ]  Physician referral [ ]  Medical records/Chart review [ ]  Media Outlet: TV, Radio ads  | [ ]  Telephone conversation text[ ]  Database/Registry[ ]  Other (please specify):       |
| ***Note: Please note that the text of all advertisement and letters used for recruiting participants must be submitted to the IRB for review and approval.***  |
| ***D. Payment to Participants taking part in the Research Project*** |
| Will participants receive any payments? [ ]  No [ ]  Yes🡪if yes, *please indicate the type and amount for each:* |
| **[ ]** Cash; LL:        | [ ]  Parking; LL:      ;  | [ ]  Transportation LL:      ;  | [ ]  Other, explain; LL:       |
| Please provide reason for payment:       |

| ***6. Informed Consent*** *(check all that apply)*  |
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| **Informed Consent will be obtained from:** [ ]  Participant [ ]  Parent/Guardian for child [ ]  Assent from Child [ ]  Legally Authorized Representative |
| **Written Informed Consent**[ ]  Informed consent will be obtained and documented through use of written **Informed Consent Form** approved by the IRB and signed by the participant or participant’s representative | **Request for Waiver Requirement to obtain a Signed written Informed Consent Form** Informed consent will be obtained and documented as follows (check all that apply):[ ]  Oral /Verbal Consent will be obtained and documented in the research record[ ]  A written statement or information sheet describing the research will be given to all participants and documented in the research record[ ]  Consent will be implied by voluntary completion of questionnaire or surveys[ ]  Request for waiver of informed consent for research that involves **No Contact With Participants**, e.g. retrospective/prospective chart review, excess tissue studies, some behavioral studies where data will remain anonymous |
| ***Note: Attach the relevant written Informed Consent Form, Information Sheet, or script for verbal consent to be used in this study and any translations. Verbatim translation is required. Back translations may be done to ensure accuracy*** |

| ***7. Data - How will participant data be collected*** *(Please include copies with submission) (check all that apply)*  |
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| [ ]  Interviews[ ]  Survey / Questionnaire Tools *(Please complete below section* ***Access for Use****)* | [ ]  Educational Training[ ]  Others, Please specify:       |
| **Access for Use of Survey / Questionnaire Tools**Please check the appropriate box for access to use the specific survey/questionnaire in your project. Documentation should be detailed and referenced in your research proposal or attached as part of this application[ ]  I created the survey / questionnaire[ ]  Survey / Questionnaire is available as Open Access [ ]  Survey / Questionnaire - I have received permission from the author to be able to use the Survey/ Questionnaire [ ]  Survey / Questionnaire - I have paid the author to be able to have access to the Survey / Questionnaire  |
| **Language to be used** *(check all that apply and provide copies of all translations):*[ ]  English [ ]  Arabic [ ]  Other:        |

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| ***8. COSTS*** |
| Will the participants be charged for research-related procedures? [ ]  No [ ]  Yes🡪*please explain:*        |

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| ***9. REASONABLY ANTICIPATED BENEFITS*** |
| 1. List the potential benefits that participants may expect as a result of this research study. State if there are no direct benefits to individual participants. ***Compensation is not to be considered a benefit.***

     1. List the potential benefits that society and/or others may expect as a result of this research study.

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| ***10. RISKS, HARMS, & DISCOMFORTS*** |
| 1. Describe all reasonably expected risks, harms, and/or discomforts that may apply to the research. Consider the range of risks, including physical, psychological, social, legal, and economic. As applicable, discuss severity and likelihood of occurrence.

     1. Describe how risks, harms, and/or discomforts will be minimized.

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| ***11. DATA SAFETY AND DATA HANDLING***  |
| Explain how information is handled, including storage, security measures (as necessary), and who will have access to the information. Include both electronic and hard copy records.      What will happen with the data after completion of the study?       |

| ***12. PRIVACY AND CONFIDENTIALITY*** |
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| a. Describe the provisions to protect the privacy of the participants     b. Does the research require access to personally identifiable private information?[ ]  No [ ]  Yes\* *If yes, please describe e.g., educational records, surveys, medical records, etc.*     c. What specific safeguards will be employed to protect confidentiality of data **during the course of the research study**, e.g. coding or removal of identifiers when sending participant related information to sponsor, limitation of access of data, use of locked file cabinets, and protection of computer based data systems etc.?[ ]  No [ ]  Yes\**\*If yes, please explain*:       |

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| ***13. PUBLICATION OR PRESENTATION OF STUDY RESULTS*** |
| Will research data be disseminated? (i.e. journal, dissertations, etc.) [ ]  Yes [ ]  No If yes, specify:       |

| ***14.* *PRINCIPAL* *INVESTIGATOR ASSURANCE*** |
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| As the Principal Investigator, by submitting this application:* I accept full responsibility for the protection of the rights and welfare of the human participants and the conduct of this study, including adherence to the ethical guidelines set forth in the Belmont Report, Declaration of Helsinki and Nuremberg Code
* I agree to comply with all applicable IRB policies and procedures, as well as with all relevant local and international laws regarding the protection of human participants in research
* I am responsible for the overall conduct of the study and will personally conduct or supervise this research within LAU and its affiliates and accept responsibility for adhering to the IRB-approved protocol
* I understand that I will not implement any modifications made to the protocol, study documents and/or informed consent documents prior to the IRB’s approval
* I understand that approval of this research could be suspended or terminated by the IRB
* I understand the requirement to report all serious and unexpected adverse events to the IRB within **10 calendar days** of the occurrence
* I understand that any research-related material is subject to an audit by the IRB
* I understand that I am required to submit a Continuing Review Application as an annual update if my research extends beyond the final approval period.
* If I am a student principal investigator, I am responsible for obtaining review and approval for this research proposal from my faculty advisor.
* I have completed the human subject protection training requirement and ensure that all investigators and personnel involved in this research have completed the human subject training requirements
* I certify that the information provided in this application is complete and accurate
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| ***15. PROJECT FACULTY ADVISOR ASSURANCE (if submission is by a Student Principal Investigator)*** |
| **As the Project Faculty Advisor for the above noted student, I have read the attached research project submitted to the LAU IRB and will ensure appropriate supervision of the student’s work.**  **Name of Project Faculty Advisor Signature Date** |