ADMUREC Form 9 - Progress Report Form

INSTRUCTIONS TO THE PRINCIPAL INVESTIGATOR: The progress report is required by the UREC for research projects that are evaluated as entailing greater than minimal risk to human participants. The report should be submitted to the University Research Ethics Office no later than 30 days from the date indicated in the formal ethics approval letter issued by the UREC. Failure to submit a progress report might result in the project being suspended until it is reviewed.

Complete all the requested information. If the item is not applicable to your protocol, write “NA”. Submit the report in electronic format to univresearchethics@ateneo.edu and in hard copy (with signatures) to the University Research Ethics Office. Date and sign this form before submission.

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| **AdMUREC CODE (UREO only):** | |
| **Study Protocol Title** | |
| **Principal Investigator** | |
| **Email and Telephone Number** | **School / Department / Affiliation** |
| **Funding Source or Sponsor and Duration of Grant** | |
| **Funding Source or Sponsor Contact (name and contact information)** | |
| **Study Protocol Approval Date/s (indicate dates of continuing review and/or protocol amendment dates, if applicable)** | |
| 1. **Summary of Study Aims** | |
| 1. **Study Site/s** | |
| 1. **Status of the Research (Please check all that apply and elaborate as needed in space below):**   ☐ Recruiting participants / following up on (enrolled) participants  ☐ Data collection has begun and is ongoing  ☐ Research has not begun: <reason/s>  ☐ Research was initiated but on hold: <reason/s>  ☐ Other: <explain> | |
| 1. **Have there been any changes or deviations so far to your UREC-approved study protocol? Changes or deviations may be in study population/sites, selection criteria, recruitment or data collection methods, new instruments, new data collected, new personnel, and other changes that materially affect the risk-benefit ratio of the study or may increase risks to participants**    1. ☐ No, all procedures are in compliance with the UREC-approved protocols and materials.    2. ☐ Yes: submit Protocol Amendment Form or indicate date/s when Protocol Amendment Form was submitted to UREO. | |
| 1. **Report on Research participants** | |
| 5.1 target number of participants approved by UREC | |
| 5.2 New participants accrued since last review/approval | |
| 5.3 Total participants accrued since study began | |
| 5.4 Participants still involved in the study | |
| 5.5 Participants who discontinued the study | |
| 5.6 Participants who have completed the study | |
| 1. **Pls review the Informed Consent Form. From your experience in the conduct of the study, are the actual risks and procedures still adequately represented in the ICF? Have there been changes in the informed consent process or documentation since last review / approval?**     1. ☐ No (complete Protocol Amendment Form and submit revised ICFs)    2. ☐ Yes | |
| 1. **Has any information appeared in the literature, or evolved from this or similar research that might affect the committee’s evaluation of the risk/benefit assessment of human participants involved in the study?**     1. ☐ No    2. ☐ Yes (Describe briefly and provide copy of literature cited, if applicable, and submit amended informed consent documents) | |
| 1. **Have there been Problems, Adverse Events, and/or Serious Adverse Events[[1]](#endnote-1) documented in the course of the study so far? Describe if there were, and the corresponding responses and mitigating actions of the PIs.**    1. ☐ No problems or adverse events have arisen in the course of the study so far    2. ☐ Indicate date/s when Unanticipated Problems/Unanticipated Adverse Events report/s were submitted to UREO, if applicable. If not yet or previously reported to UREO, indicate reason for not reporting: | |
| 1. **Summary of Participants’ Complaints, Unfavorable comments, or Grievances documented in the course of the study so far**    1. ☐ There have been no complaints or unfavorable comments or grievances.    2. ☐ Indicate date/s when report/s submitted to UREO, if applicable. If not previously reported to UREO, indicate reason for not reporting: | |
| 1. **Summary of Benefits of the study so far (Direct or Indirect)** | |
| 1. **Are identifiable data being stored for this study?**   ☐ N/A  ☐ Yes – are you complying with your UREC-approved protocol? Describe what information is being stored, in what format, and the the measures you are taking to protect the confidentiality of records  ☐ No – data is or has been de-identified / anonymized | |
| **Declaration**  ☐ I confirm that the study and its principal investigators and research personnel continue to abide by the ethics standards and guidelines of the Ateneo de Manila University.  ☐ I confirm that, if necessary, I will submit the relevant, requisite forms and reports (e.g. Protocol Amendment Form, Continuing Ethics Review Application, Unanticipated Problems Report, etc.) to the University Research Ethics Office to update on the status of the project. | |
| **Signature of Principal Investigator**: | |
| **Progress Report Submission Date:** | |

RECOMMENDATIONS (for AdMUREC use only)

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| Comments of Reviewer(s)  (i.e. compliance with the terms of the approved protocol; overall assessment of risks against benefits in the conduct of study) | | | Details | |
| RECOMMENDED ACTION   * UPHOLD ETHICS APPROVAL UNTIL <dd/mm/yyyy>\_\_\_\_\_\_\_\_\_\_\_\_\_\_ * REQUEST INFORMATION * RECOMMEND FURTHER ACTION | | |  | |
| REVIEWER(S) |  | Signature: |  | Signature: |
| Date: |  | Name |  | Name |
| UREC/PANEL CHAIR |  | Signature: |  | |
| Date: |  | Name |  | |

Endnotes

1. Problems and adverse events may be anticipated (reported in the initial application for ethics clearance) or unanticipated. It is mandatory to report unanticipated problems and adverse events to UREO. Unanticipated problems are defined as any incident, experience, or outcome that meets all of the following criteria: a) it is unexpected, in terms of nature, severity, or frequency, given the research procedures that are described in the UREC-approved protocol and informed consent document, and the characteristics of the subject population being studied; b) related or possibly related to participation in the research; and c) suggests that the research places subjects or others at a greater risk of harm than was previously known or recognized.

   Adverse events include any event meeting the following definition: Any untoward or unfavorable medical occurrence in a human participant, including any abnormal sign, symptom, or disease, temporally associated with the subject’s participation in the research, whether or not considered related to the subject’s participation in the research.

   Serious adverse events are those temporally associated with the individual’s participation in the study that meets any of the following criteria:

   * results in death;
   * is life-threatenting;
   * requires inpatient hospitalization or prolongation of existing hospitalization;
   * results in persistent or significant disability or incapacity;
   * results in a congenital anomaly/birth defect; or
   * any adverse event that, based on appropriate medical judgment, may jeopardize the participant’s health and may require medical or surgical intervention to prevent any of the aforementioned outcomes

   [↑](#endnote-ref-1)