**PROGRESS REPORT**

**As of <date>**

**PROTOCOL GENERAL INFORMATION**

|  |  |
| --- | --- |
| **PSURERC Code** |  |
| **Protocol Title** |  |
| **Principal Investigator/Lead Researcher** |  |
| **Type of Initial Review** |  |
| **Protocol Approval Date/s** *(indicate dates of continuing review and/or protocol amendment dates, if applicable***)** |  |
| **Target Start Date** *(as stated in the approved protocol; start of study includes the start of all data collection-related activities post-approval, e.g., communication with recruitment party, invitation, etc. )* |  |
| **Target Completion Date** *(as stated in the approved protocol)* |  |
| **Study Site** |  |

**STATUS OF THE STUDY**

|  |  |
| --- | --- |
| **Current Stage** | **Details/Reasons** |
| [ ] Study has started and  is ongoing  [ ] Study has been initiated, but is  on hold  [ ] Study has not started | *<include actual start date (explain if date is different from target start date), and currently at what specific stage>*  *<include actual start date (explain if date is different from target start date), on hold since when (and reasons), and currently at what specific stage>*  *<reasons and planned steps to be taken>* |

**REPORT ON STUDY PARTICIPANTS**

|  |  |
| --- | --- |
| Target Number of Participants (*approved by RERC)* |  |
| Number of Participants Invited |  |
| Number of Participants Enrolled |  |
| Number of Participants Withdrawn *(include reasons for withdrawal)* |  |
| Number of Participants Who Completed |  |

**REPORT ON SPECIFIC PROGRESS**

| **Provide a description and details. Indicate NA if not applicable.** | |
| --- | --- |
| Approved amendments since the last review/report |  |
| Deviations from the approved protocol since the last review/report |  |
| Reportable negative events since the last review/report |  |
| Adverse events or reactions since the last review/report |  |
| Complaints |  |
| New information in recent literature or similar research that may change the risk/benefits ratio for participants |  |
| Other issues and problems encountered (aside from RNES and AER) |  |
| Benefits accrued so far |  |
| Risks exposures |  |

**DECLARATION**

[ ] I confirm that the study and its investigators and research personnel continue to abide

by the ethics standards and guidelines of the National Ethical Guidelines 2022.

[ ] 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 PSURERC to update on the status of the study.

**Principal Investigator/Lead Researcher: <Name and Signature>**

**Date of Report Submission: <Date>**

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*To be filled out by PSURERC Primary Reviewer*

|  |  |
| --- | --- |
| **RECOMMENDATIONS:** | [ ] Approve  [ ] Request additional information  [ ] Request further action  [ ] Amendment in the protocol  [ ] Amendment in the ICF/assent  [ ] Others  [ ] Pending, if major clarifications are needed before a decision can be  made  [ ] Suspend the study  [ ] Terminate the study |
| **Primary Reviewer** | *<Name and Signature>* |
| **Date of Recommendation** | *<Date>* |