|  |  |
| --- | --- |
| Study Title: |  |
| IRB Reference No: |  |
| Principal Investigator: |  |

|  |  |
| --- | --- |
| **MODIFIED MATERIALS**  Check the applicable options to indicate the modified materials: | |
| Research Submission application form |
| Informed Consent Form |
| Child Assent Form |
| Flyer, Advertisement |
| Questionnaire |
| Other, specify: |

1. **Study Status**

Please indicated what best reflects your research from the options below: [click the applicable checkbox]

The research study did not start yet.

The research involves pre-existing records or samples

New participant recruitment is still in progress.

Enrolment closed, but participants are still undergoing study procedures.

Enrolment closed, participants have completed study procedures, but are still in follow-up.

Remaining study activity is limited to analysis only, no further contact with participants.

## Summary of changes

*Briefly summarise the main changes proposed in this amendment. Explain the purpose of the changes and their significance to the study if applicable. [Please write a summary of the changes in the box provided below and submit a track changes copy of the revised documents.]*

|  |
| --- |
|  |

**Any other relevant information**

Indicate any specific ethical issues related to the amendment proposed, on which the opinion of the IRB is sought.

|  |
| --- |
|  |

# Declaration by the Principal Investigator:

*(please confirm by clicking the checkbox)*

**I confirm that the information in this form is accurate to the best of my knowledge, and I take full responsibility for it. In addition, I consider that it would be reasonable for the proposed amendment to be implemented.**