**FINAL REPORT (Form 3.4)**

**TO THE PRINCIPAL INVESTIGATOR:** *OBTAIN AN ELECTRONIC COPY OF THIS FORM AND ENCODE ALL INFORMATION REQUIRED IN THE SPACE PROVIDED. PRINT NAME, DATE AND SIGN THIS FORM BEFORE SUBMISSION. INDICATE WITH A (✓) CHECK MARK THE APPROPRIATE TICK BOX.*

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| **Date of Submission (MMM/DD/YYYY)** | Click here to enter text. | **IRB Protocol Number** | Click here to enter text. |

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| **Sponsor** | Click here to enter text. | **Sponsor’s Protocol Number** | Click here to enter text. |

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| **Principal Investigator** | Click here to enter text. | **Co-investigator(s)** **(if any)** | Click here to enter text. |

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| **Principal Investigator’s Signature** |  | **Principal Investigator’s Contact Number** | Click here to enter text. |

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| **Protocol Title**  | Click here to enter text. |

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| **Date of Initial Approval of Protocol (MMM/DD/YYYY)** | Click here to enter text. |

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| **Study site(s):** | Click here to enter text. | **No. of Study Arms** | Click here to enter text. |

1. **CUMULATIVE SUMMARY OF SUBJECTS ENROLLED TO DATE**

*(For studies involving record and/or specimen review only, skip and complete Section B)*

 *(For study designs utilizing multiple consent forms, this table may be replicated).*

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| 1. **Number of subjects accrued**
 | Click here to enter text. |

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| 1. **Number of subjects currently active/ on study**

*(For example, subjects receiving study interventions/ interactions or long-term follow-up)* | Click here to enter text. |

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| 1. **Number of subjects completed**

*(Without events leading to early termination/ withdrawal from the study)* | Click here to enter text. |

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| 1. **Number of subjects who voluntarily withdrew consent after enrolling**

*(For example, after signing the consent form, the subject changes his/her mind and decided not to participate or to stop participating after completing some of the study procedures)* | Click here to enter text. |

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| **Explanation**  | Click here to enter text. |

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| 1. **Number of subjects terminated/ withdrawn from the study by the investigator due to adverse event(s)**

*For example, subject met toxicity drop point or experienced a serious adverse event.*  | Click here to enter text. |

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| **Explanation**  | Click here to enter text. |

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| 1. **Number of subjects terminated/ withdrawn from the study by the investigator due to other reasons**

*For example, non-compliance with the protocol, pregnancy, etc.*  | Click here to enter text. |

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| **Explanation**  | Click here to enter text. |

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| 1. **Number of subjects lost to follow-up**
 | Click here to enter text. |

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| **Explanation**  | Click here to enter text. |

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| 1. **Number of subjects who are no longer participating for reasons other than those above**
 | Click here to enter text. |

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| **Explanation**  | Click here to enter text. |

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| 1. **Total of item nos. 2 to 8**

*(should be equal to item no. 1)* | Click here to enter text. |

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| 1. **Number of subjects approved at Makati Medical Center**
 | Click here to enter text. |

1. **RECORDS AND SPECIMENS**

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| 1. **Number of specimens and/or records approved by the IRB:**
 | Click here to enter text. |

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| 1. **Did you review medical records, patient charts, radiographs or other patient information for this study?**
 | **[ ]  No** **[ ]  Yes**  | **No. of records reviewed to date:** | dd |

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| 1. **Did you analyze specimens (e.g., archival tissue, blood, blood products, or body fluids) for this study?**
 | **[ ]  No** **[ ]  Yes**  | **No. of specimen analyzed to date:** |  |

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| **Duration of the Study (Date Initiated and Completed)****(MMM/DD/YYYY)** | Click here to enter text. |

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| **Objectives** | Click here to enter text. |

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| **Summary of Results:** |
| Click here to enter text. |

**TO THE PRIMARY REVIEWER/ INDEPENDENT CONSULTANT:***INDICATE WITH A (✓) CHECK MARK THE APPROPRIATE TICK BOX. SPECIFY ON THE SPACE PROVIDED OTHER COMMENTS, IF APPLICABLE.*

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| **Comments/ Recommendations of the Primary Reviewer**  |
| * Acknowledged
* Request for further information

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| **Primary Reviewer’s Name** | **Signature** | **Date (MMM/DD/YYYY)** |
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| **Date of IRB meeting the report was presented****(MMM/DD/YYYY)** |  |