**DEVIATION/ NON-COMPLIANCE/ VIOLATION REPORT (Form 3.5)**

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

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| **Date of Submission (MMM/DD/YYYY)** |  | **IRB Protocol Number** |  |

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| **Sponsor** |  | **Sponsor’s Protocol Number** |  |

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| **Principal Investigator** |  | **Co-investigator(s)** **(if any)** |  |

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| --- | --- | --- | --- |
| **Principal Investigator’s Signature** |  | **Principal Investigator’s Contact Number** |  |

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| **Protocol Title**  |  |

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| **Reported by** |  | **Contact Number** |  |

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| **Description of Deviation/ Violation:** |
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| **Action(s) Taken** |  | **Date (MMM/DD/YYYY)** |  |

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| **Nature of the Protocol Deviation/ Violation:**  |
|  [ ]  Principal Investigator Deviation from the protocol [ ]  Participant Non Compliance [ ]  Others: Beyond the situations |   [ ]  Major[ ]  Minor |

**TO THE PRIMARY REVIEWER :***PUT A (√) MARK ON THE APPROPRIATE TICK BOX. PRINT NAME, SIGN AND DATE THIS FORM.*

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| **IRB DECISION** | [ ]  Continue study and monitor compliance [ ]  Request for further information[ ]  For site visit[ ]  Amend Protocol | [ ]  Amend Informed Consent Form[ ]  Suspend the study[ ]  Terminate approval of current study |

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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)** |  |