|  |  |
| --- | --- |
| **Trial Name** |  |
| **Chief Investigator** |  | **IRAS No.**  |  |

|  |  |  |
| --- | --- | --- |
| *(If applicable) Participant Trial ID No. & initials (or other point of ID)* | *Initials (or other ID)*  | *Trial ID No.* |
| **This non-compliance is related to:** *Specify general category or type of N/C* |
| **Category of non-compliance:** *May be tailored according to trial (Excel log listing categories to match)*  |
| Missed safety test |  |
| Missed routine test |  |
| Missed research assessment / procedure |  |
| Missed study visit |  |
| Study visit outside of protocol window |  |
| IMP (\*incorrect doses / IMP stock / expired IMP / others) \**Delete where applicable*  |  |
| Consent  |  |
| Other aspect of trial |  |
| **Full details of non-compliance:** |
| **Relevant section(s) of Protocol/SOPs** *(number and version):*  |
| **Corrective action(s) taken:***Document any corrective action(s) taken at the time of non-compliance relating to this instance of the non-compliance. If none could be taken, enter the reason why***Preventative action(s) taken:** *Document any action(s) taken to prevent future similar occurrences of the non-compliance* **(If applicable) File Note reference: ……………………….** |
| **Report completed by: ………………………….Signature: ………………………Date: ………………………** **Reviewed by PI: …..……………………………..Signature: ……………………..Date: ………………………** *(If different from above)***Sites to send a copy of PI reviewed form to the Trial Coordinator** |

**CHIEF INVESTIGATOR (CI) ASSESSMENT**

|  |
| --- |
| **Categorisation of non-compliance** (*tick relevant box):*Type 2 Type 3Reportable minor Reportable major non-compliance\*\*Non-compliance \*\*Potential serious breach\*\*Refer to CCTU/SOP018 & R&D/SOP003.*Type 1 non-compliances do not require a N/C form to be completed, but should still be recorded on the N/C log* |
| **CI Justification of categorisation:** |
| **CI name………………………..………. Signature………………………Date…………………………** |

**Send a copy of CI reviewed form to the CCTU regulatory team**

|  |  |
| --- | --- |
| **For coordinating centre use only:**Date Coordinating Centre became aware of Non Compliance |  |
| Coordinating Centre Non-compliance No.  |  |
| Current number of similar non-compliances at this site |  |

**SPONSOR OVERSIGHT ASSESSMENT**

|  |  |
| --- | --- |
| **Date non-compliance report received by CCTU** |  |
| **Date of Regulatory Team Assessment** |  |
| **Categorisation of non-compliance** (*Tick relevant box):*Type 2 Type 3Reportable minor Reportable major non-complianceNon-compliance Potential serious breach |
| ReviewerName………………………………………….. Signature……………………………….. Date……………………………..  |
| **FOR POTENTIAL SERIOUS BREACH:**  |
| **This event is likely to:** | *(Please tick relevant box)* |
| Affect to a significant degree the safety, or physical or mental integrity of the trial subjects |  |
| Affect to a significant degree the scientific value of the trial |  |
| **Date of escalation to Sponsor** |  |
| **ADDITIONAL COMMENTS (if any):**  |

**Note: The Regulatory Team categorisation is the definitive categorisation for the purposes of escalation**

**Original Initial Sponsor oversight assessment to be retained by Sponsor, 1 copy to be sent to trial Coordinator**

**Date copy returned to trials team by the regulatory team ………………………………..**

**Signature…………………………………………………………………..**

**Date copy returned to the site team by the coordination team…………………………..**

 **Signature…………………………………………………………………..**