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| **Adverse Event Log** |
| Study title |
| **R&D ID:**  | **IRAS number:** |
| **Chief Investigator:**  | **Principal Investigator:** |
| **Site:**  | **Participant ID:** |

| **Description of adverse event**(provide additional information on notes pages if required) | **Date of onset** | **Date reported to Investigator/team** | **Severity**1. Mild2. Moderate3. Severe | **Causality**1. Unrelated2. Possible3. Probable4. Definite  | **Action taken – please list all that apply**1. None2. Hospitalisation3. Intervention stopped4.Con Meds commenced (record on Con Meds Log)5. Other (specify) | **Outcome**1. Recovered2. Recovered with sequelae3. Recovering4. Not recovered5. Unknown6. Fatal | **Is this a Serious AE?** **YES\*** **or NO** | **Date resolved**(Enter date resolved or tick if ongoing at end of study) | **Signature and Date** |
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|  |  |  |  |  |  |  | \*complete an SAE form and email to the Sponsortay.pharmacovigilance@nhs.scot  |
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