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| **SITE DETAILS** | | | | | | | | | |
| **Protocol Number:** | GI-IN26843-78376432 | | | | | **Site Name:** |  | | |
| **Principal Investigator:** |  | | | | | **Site Number:** |  | | |
| **Name** | **Study Role** | **Signature** | | **Initials** | **Study Task(s)\*** | **Training Date** | **Start Date** | **End Date**  **(complete only if prior to end of study)** | **Authorised by PI (Initials & Date)** |
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| **\*Key for study tasks** | | | | | | | | | |
| 1. Prepare study drug 2. Dispense study drug to bedside staff 3. Provide instructions to bedside staff on study drug administration | | | 1. Perform study drug accountability 2. Record and report any protocol deviations containing unblinded information 3. Maintain Unblinded Trial Folder | | | | | | |

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| **End of Study Declaration (to be completed by the Principal Investigator at the end of the study)**  **I confirm that the above information is accurate and complete, and that I authorised the delegation of study-related tasks to each of the appropriately trained, skilled and qualified individuals as listed above.** | | | | | |
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| **Principal Investigator Name** |  | **Signature** |  | **Date** |