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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** |