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| --- | --- | --- | --- | --- | --- | --- |
| Full Name | Role in Study | Responsibility for | Signature | PI initials | Involvement Start date | Involvement End date |
|  | Principal Investigator | All tasks undertaken at site |  |  | DD-MMM-YYYY | DD-MMM-YYYY |

**\*By signing, the PI confirms that each team member is adequately trained appropriate to their trial related duties and functions.**

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| Full Name | Role in Study | Tasks delegated**\*\*** | Signature | Initials | Date of PI signature (Involvement start date of individual) | Signature of PI\* | Involvement  End date | PI Initials  (for end date) |
|  |  |  |  |  | DD-MMM-YYYY |  | DD-MMM-YYYY |  |
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**\*\* Please indicate delegated study tasks in this column using the task numbers from the list below. If amendments are made the PI must initial and date the changes.**

|  |  |
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| **01. Maintenance of Investigator Site File** | **08. Obtaining consent using optional pre-screening tumour PIS & ICF** |
| **02. Confirm patient eligibility *(decision must be made by a medically qualified person)*** | **09. Managing posting of ctDNA samples** |
| **03. Obtain informed consent using main PIS & ICF** | **10.** |
| **04. Screen trial patients** | **11.** |
| 1. **Register and/or randomise trial patients** |  |
| **06. e/CRF completion (including query resolution)** |  |
| **07. Clinical assessment of AEs & SAEs/SARs/SUSARs *(must be a medically qualified doctor)*** |  |

**By signing this form you are agreeing to allow SCTU to hold this data electronically.**

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| Full Name | Role in Study | Tasks delegated**\*\*** | Signature | Initials | Date of PI signature (Involvement start date of individual) | Signature of PI\* | Involvement  End date | PI Initials  (for end date) |
|  |  |  |  |  | DD-MMM-YYYY |  | DD-MMM-YYYY |  |
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| **01. Maintenance of Investigator Site File** | **08. Obtaining consent using optional pre-screening tumour PIS & ICF** |
| **02. Confirm patient eligibility *(decision must be made by a medically qualified person)*** | **09. Managing posting of ctDNA samples** |
| **03. Obtain informed consent** | **10.** |
| **04. Screen trial patients** | **11.** |
| 1. **Register and/or randomise trial patients** |  |
| **06. e/CRF completion (including query resolution)** |  |
| **07. Clinical assessment of AEs & SAEs/SARs/SUSARs *(must be a medically qualified doctor)*** |  |