**SITE RESPONSIBILITY (DELEGATION) LOG**

**Title of Protocol:** Study into the Reversal of Septic Shock with Landiolol (Beta Blockade) **Trial Site:**

**Trial Acronym/ Reference:** STRESS-L **Principal Investigator Name:**

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| Individual | | | | | | PI Authorisation | | Role Finished | |
| Name | Job title | Trial Responsibilities | Signature | Initial | Date | Signature of PI | Date | Initial (PI) | Date |
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| **Trial Responsibility Codes:** |  |  |
| **A** = Overall responsibility for trial at site *(PI only)* | **H** = Data query resolution and return | **O** = Investigator Site File maintenance |
| **B** = Subject screening and selection | **I** = SAE attribution assessment and sign off SAE form | **P** = Pharmacy Site File maintenance & pharmacy procedure |
| **C** = Obtaining Informed consent | **J** = SAE reporting | **Q** = Ensure IMP Inventory |
| **D** = Assisting in informed consent process | **K** = Prescribe medication and treatment | **V** = Prepare and be available for audit and inspections |
| **E** = Review eligibility and sign patient eligibility form | **L** = Dispense trial medication | **W** = Archiving of trial data |
| **F** = Medical care and supervision of trial participants | **M**= Perform randomisation |  |
| **G** = eCRF completion and correction | **N** = Sample collection, preparation and dispatch |  |

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