

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:

Individual						PI Authorisation		Role Finished	
Name	Job title	Trial Responsibilities	Signature	Initial	Date	Signature of PI	Date	Initial (PI)	Date

Trial Responsibility Codes:

- A** = Overall responsibility for trial at site (*PI only*)
- B** = Subject screening and selection
- C** = Obtaining Informed consent
- D** = Assisting in informed consent process
- E** = Review eligibility and sign patient eligibility form
- F** = Medical care and supervision of trial participants
- G** = eCRF completion and correction

- H** = Data query resolution and return
- I** = SAE attribution assessment and sign off SAE form
- J** = SAE reporting
- K** = Prescribe medication and treatment
- L** = Dispense trial medication
- M** = Perform randomisation
- N** = Sample collection, preparation and dispatch

- O** = Investigator Site File maintenance
- P** = Pharmacy Site File maintenance & pharmacy procedure
- Q** = Ensure IMP Inventory
- V** = Prepare and be available for audit and inspections
- W** = Archiving of trial data