## **PRISM Delegation Log**

## **Principal Investigator:**

Site code/ID:





Name	Study Role	Signed Initials	Delegated Tasks (List duty categories)	PI confirmation		Study duration	
				Signature	Date	Start date	Date of leaving study
	Principal Investigator						

All those involved in the above study must read the protocol (and amendments if applicable) and understand their role as outlined in the protocol

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- 1. Obtaining informed consent
- 2. Physical exam / clinical evaluations
- 3. Source document entry (ie. medical notes)
- 4. CRF completion

- 5. Data entry
- 6. Resolving data queries
- 7. Review & reporting adverse events & SAEs

8.	Maintaining study file

9.	Archiving
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ther	duties	specific t	o above	study	Please spec	ify below

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Chief Investigator: Professor Rupert Pearse