

**Protocol Training Log**  
 All site personnel involved in this clinical trial must complete this form.  
 The form should be filed in Trial Master File/Investigator Site File

Sponsor Reference number:	Principal Investigator:
Site Name:	Study Title:

Date	Training Topic*	Trainer name	Trainer signature	Trainee Name	Trainee signature
.././....					
.././....					
.././....					
.././....					
.././....					

I ..... (Name) confirm that I have read/received training in the areas as described above. I understand that any amendments to the protocol may result in the requirement for retraining in the relevant areas.

..... (Signature) ..... (Date)

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|---|--|-----------------|
| 1. Protocol (include version number)      | 7. GCP and regulatory requirements           | 12. Other ..... |
| 2. Investigator Brochure/SmPC             | 8. Maintenance of source documents           | 13. Other ..... |
| 3. Informed consent procedures            | 9. Maintenance of TMF/ISF                    | 14. Other.....  |
| 4. AE/SAE reporting procedures            | 10. Handling/storage/Shipping of Lab samples |                 |
| 5. CRF/eCRF/Data Entry                    | 11. Other .....                              |                 |
| 6. Electronic Case Report form/data entry |  |                 |