

Clinical Trial Project Closeout Checklist

Project Information

Project Title	Protocol Number
Principal Investigator	Study Site(s)
Sponsor	Date of Closeout

Closeout Checklist

Task / Item	Completed	Notes
All data and source documents verified and archived		
All regulatory documents updated, signed and archived		
Study drug/device returned or destroyed as per protocol		
Final monitoring visit conducted		
Final report submitted to sponsor		
Final report/notification submitted to IRB/EC		
Site equipment/supplies returned (if applicable)		
Subject compensation and reimbursement completed		
Database locked and transferred to sponsor		
Essential documentation complete (Delegation logs, CVs, training logs, etc.)		
Investigator's final acknowledgment		

Comments / Additional Notes

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Prepared by (Name & Title):	Date:
Reviewed by (PI/Sponsor):	Date: