

THESIS ORGANIZATION CHECKLIST

Medical Thesis Data Management | AcadLabs ThesisLog | acadlabs.in/thesisLog

Study Title: _____ _____	Investigator: _____ _____	IEC No: _____ _____	Start Date: _____ _____
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■ PHASE 1 — Pre-Study (Before First Patient)

■	Study protocol approved by thesis guide and department head
■	IEC / IRB approval letter obtained and filed
■	Patient Information Sheet (PIS) prepared — English + local language
■	Informed Consent Form (ICF) designed and IEC-approved
■	Case Record Form (CRF) finalised (Version 1.0) and guide-approved
■	Data sheet template created with validation rules on all columns
■	Codebook created — every variable defined with allowed values
■	Patient Log set up (separate from data sheet)
■	Study ID numbering system decided (e.g., TL001–TL200)
■	Subject Identification Log created (secured, separate storage)
■	Backup system set up: laptop + cloud + external drive (3-2-1 rule)
■	Folder structure created: 01_Pre-Study / 02_Data-Collection / 03_Locked / 04_Analysis / 05_Manuscript

■ PHASE 2 — Data Collection (During Enrollment)

■	Consent obtained before any study procedures for every patient
■	Signed ICF filed under patient's Study ID
■	CRF completed same day as patient visit — no blank required fields
■	Data entered into data sheet within 24 hours of CRF completion
■	Patient Log updated after every patient event (enrollment, visit, exit)
■	QC check run every 20 patients: review missing values per column
■	Protocol Deviation Log updated if any deviation occurred
■	Serious Adverse Events reported to IEC within mandated timeframe
■	Backup confirmed after every data entry session
■	CRF scans saved as TL001.pdf, TL002.pdf in /CRFs folder
■	Monthly reconciliation: Patient Log numbers match Data Sheet rows

■ WEEKLY ROUTINE (10 minutes every Sunday)

1. Confirm all CRFs scanned and saved · 2. Patient Log updated · 3. Missing-values count run · 4. Cloud backup verified · 5. External drive copy made

■ PHASE 3 — Close-Out & Database Lock

■	All outstanding CRFs completed and data entered
■	Final QC check run — all missing values resolved or coded -9/NA
■	Query Log reviewed — all open queries resolved
■	Patient Log marked: every patient has a final status
■	CONSORT / STROBE numbers verified against Patient Log totals
■	Database lock performed: read-only copy saved as DataSheet_LOCKED_[date].xlsx
■	Codebook finalised and matches locked dataset columns exactly
■	Statistical Analysis Plan (SAP) written before analysis begins
■	Locked dataset shared with guide for review before statistical handoff

■ PHASE 4 — Analysis & Submission

■	All analysis performed on locked dataset only — never the live file
■	SPSS / R / Stata output files saved in /04_Analysis folder
■	Analysis scripts or syntax files archived (not just screenshots)
■	Results verified by guide before writing begins
■	Thesis manuscript drafts versioned: Draft_v1_[date].docx
■	Plagiarism report generated (Turnitin / iThenticate)
■	Final thesis signed by guide, co-guide, and department head
■	Data archiving plan confirmed: CRFs + consent forms + dataset stored per policy (min. 5 years)
■	If publishing: data availability statement prepared, de-identified dataset ready

■ FILE NAMING QUICK REFERENCE

Document Type	Correct Naming Format	Example
Protocol	Protocol_v[N]_[YYYY-MM-DD].docx	Protocol_v2.1_2025-08-10.docx
Live data sheet	DataSheet_LIVE.xlsx	— never date the live file —
Locked data sheet	DataSheet_LOCKED_[YYYY-MM-DD].xlsx	DataSheet_LOCKED_2026-01-15.xlsx
Patient CRF (scanned)	[StudyID]_CRF_[YYYY-MM-DD].pdf	TL047_CRF_2025-09-14.pdf
Consent form	[StudyID]_Consent_[YYYY-MM-DD].pdf	TL047_Consent_2025-09-14.pdf
Thesis draft	Draft_v[N]_[YYYY-MM-DD].docx	Draft_v3_2026-03-01.docx