GCP Auditing:

Take a quality snapshot of your clinical study

Auditing brings an independent, quality assurance perspective to the clinical research landscape.

Investigational sites, sponsors, and vendors benefit from high-level process assessments and improvements. Auditors leverage extensive training and experience to help ensure subject safety, data integrity, and protocol and regulatory compliance.

When to Audit during a Clinical Study?	Why Audit?
During Early Enrollment	Snapshot of potential process-level problems (consenting, event reporting) in time to course-correct
When Hiring New Vendors	Qualify and verify product/service offerings
Shortly Before or After a PMA/NDA Submission	Assess inspection readiness, adequacy of processes, documentation, data integrity, and that subject safety has been ensured
Anytime!	With concerns of noncompliance, elevated rates or absence of AEs, high-risk sites or studies

Invest in the quality of your clinical program by
incorporating GCP Auditing into your studies today!

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