



Trial-Ready GCP Training for Sponsors, Sites, and CROs


Drive ICH GCP E6(R3) compliance with trial-ready courses maintained by Veeva. This includes a standalone GCP course recognized by TransCelerate, as well as additional accredited courses for further GCP learning.

Automatically assign GCP training across sponsor, site, and CRO personnel in Veeva Study Training. Plus, enable site staff to demonstrate GCP training completion across sponsors to reduce repetitive training.

Benefits

- 

Drive GCP compliance
with tailored courses recognized by TransCelerate BioPharma.
- 

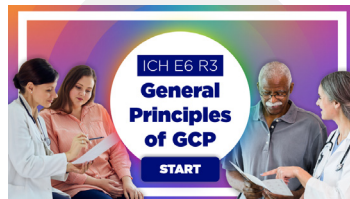
Future-proof your training
with Veeva-managed **courses** that evolve with regulatory updates.
- 

Streamline training
by assigning courses automatically through Veeva Study Training.

Courses

Standalone course for GCP compliance

- **Veeva GCP for Clinical Trials** (45 minutes)



Additional courses

- **ICH E6(R3) - General Principles of GCP** (20 minutes)
- **ICH E6(R3) - Sponsor** (25 minutes)
- **ICH E6(R3) - Data Governance** (15 minutes)
- **ICH E6(R3) - Investigator** (10 minutes)
- **ICH E6(R3) - IRB/IEC** (15 minutes)
- **ICH E6(R3) - Appendices** (15 minutes)
- **Investigator Oversight of Clinical Trials** (15 minutes)
- **Informed Consent** (15 minutes)
- **Clinical Trial Safety Reporting** (10 minutes)
- **Fundamentals of RECIST 1.1** (15 minutes)

About Veeva Study Training

Veeva Study Training brings sponsors, CROs, and research sites into a single platform to streamline and automate training while ensuring inspection readiness.

About Veeva LearnGxP

With **Veeva LearnGxP's** accredited GxP training library, organizations can significantly reduce deviations, reclaim valuable workforce time, and eliminate the burden of creating training in-house. View our **course catalog**.