

WHO ARE WE?

Technology Ed is a leading provider of workforce development online training on pharmaceutical industry and related science, engineering, quality, regulatory, and management topics. Our training program is a subscription solution (with an unlimited company license).

Our programs are designed by a team of industry experts from each respective field, who work to provide your employees with an effective web-based learning experience. Instructors/mentors are available during your online learning experience.

1. *Learning modules (30 minutes each)*
2. *Self checks*
3. *Final situational assessment*
4. *Supplemental tools/resources*

We know you'll appreciate the quality as well as the convenience of anytime, anywhere learning!

We are a member of the American Association for Adult and Continuing Education.

WHO ARE OUR PARTNERS?

Educational Partners



Government and Industry

Employees from leading organizations such as the US Food & Drug Administration, Arnet Pharmaceutical, Boston Scientific, Boehringer Ingelheim, Pfizer, have benefited from our online training programs.



WHAT DO WE OFFER?

We offer 20+ pharmaceutical-related online training courses (30 minute modules) that can be provided on a subscription solution for your organization.

- Pharmaceutical Quality Assurance & Regulatory Affairs
- Pharmaceutical Validation Processes
- Pharmaceutical Manufacturing
- Pharmaceutical Drug Discovery
- Pharmaceutical Sales Training Program
- Medical Device Quality Assurance & Regulatory Affairs
- Medical Device Change Control
- Medical Device Technology
- Management of Change
- Good Clinical Practices
- Good Documentation Practices
- Good Laboratory Practices
- Good Manufacturing Practices (GMP)
- Hazards Analysis Critical Control Points (HACCP)
- Process Validation
- Equipment Validation
- Computer System Validation
- Analytical Equipment Validation
- Quality Audits
- Quality Concepts
- Quality Control
- Quality Investigations
- Quality & Statistical Tools
- Developing a Product Specification
- Developing a Corrective Action Report
- Phase Gate New Product Development
- Premarket Approval
- 510(K) Clearance to Market
- Design of Experiments
- Six Sigma

HOW TO REGISTER?