



2026 BEC Course Catalog

ASQ Certification Academies

- ASQ Certified Quality Auditor
- ASQ Certified Quality Engineer
- ASQ Certified Six Sigma Green
- ASQ Certified Manager of Quality & Organizational Excellence
- ASQ Certified Supplier Quality Professional

Investigations and Effective CAPA Systems

- Investigation/CAPA System and Human Errors Reduction Certification
- Root Cause Analysis
- Human Error Investigations
- Effective CAPA Systems Management Overview
- Effective OOS/OOT Investigations for QC Analytical Laboratory Certification
- Microbiology Investigations and Environmental Monitoring Program
- Implementing a Mature and Sustainable Investigations/CAPA Program
- Data Integrity Investigations

Technical and Compliance Writing

- Effective Compliance and Regulatory Writing
- How to Write Procedures to Reduce Human Errors

Regulations and Standards

- MDSAP – Medical Device Single Audit Program
- 21 CFR 111: cGMP for Dietary Supplements
- 21 CFR 211: cGMP for Finished Drugs
- 21 CFR 820: QMSR for Medical Devices
- Understanding Combination Products
- ISO 9001:2015
- ISO 13485:2016
- ISO 14971:2019
- ISO 17025:2017
- ANVISA's GMP for Medical Device Regulation RDC No. 665

Auditing

- Internal Auditing Certification
- ISO 9001:2015 Lead Auditor (36-hour Exemplar Global Certified)
- ISO 13485:2016 Lead Auditor (36-hour Exemplar Global Certified)

Risk Management

- Quality Risk Management Certification
- ISO 14971:2019



Statistics

- Data Trending Analysis
- Basic Applied Statistics
- Sampling Best Practices for the FDA-Regulated Industry

Validations

- Computer System Validation for the FDA-Regulated Industry
- Cleaning Validation Lifecycle
- Facilities and Critical Utilities Systems Qualification
- Laboratory Equipment Qualification
- Sterilization
- Test Method Validation
- Process Validation for Medical Devices
- Cleaning validation for Medical Devices
- Process Validation for Pharmaceutical Products

Compliance

- Overview of Stability Programs for Drug, Biotech, and Combination Products
- Best Practices for Complaint Handling in the FDA Regulated Products
- Trending Analysis for the Stability Program
- Understanding FDA Drug Manufacturing Inspection
- Understanding FDA Medical Device Manufacturing Inspections

Training Program

- Train the Trainer
- How to Measure Training Effectiveness
- Knowledge Management

Organizational Behavior

- Managerial Decision Making and Critical Thinking
- Leadership Skills

Data Integrity

- Data Integrity Investigations
- Data Integrity Certification
- Data Integrity module for QC laboratory personnel
- Data Integrity module for manufacturing and operation personnel
- Data Integrity module for management

Management Controls

- Pharmaceutical cGMP for Leaders and Managers – Managerial Responsibilities
- How to Implement an Effective Change Management Control Program
- FDA Quality Management Maturity Model and **Quality Culture**
- Pharmaceutical Inspection Readiness Certification