Business Excellence Consulting, Inc.
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Calendario de Talleres

Horarios de Puerto Rico – AST (Atlantic Standard Time) – Actualizado 11-06-2026

Fechas	VIRTUAL Training
2 días (8:00 am – 12:00 pm) Noviembre 10 y 12, 2026	**Data Trending Analysis** (7 horas)
1 día (8:30 am – 5:00 pm) Noviembre 17, 2026	**Critical Thinking for Investigators**
3 días (8:30 am – 5:00 pm) Diciembre 14,16 y 18, 2026	**Effective Internal Auditing Certification** (21 horas)
3 sábados (8:30 am – 5:00 pm) Enero 30, Febrero 6 y 13, 2027	**ASQ Certified Quality Auditor CQA** (21 horas)
5 sábados (8:30 am – 5:00 pm) Febrero 27, Marzo 6, 13, 20 y 27, 2027	**ASQ Certified Quality Engineer CQE** (35 horas)
1 día (9:00 am – 5:00 pm) TBD	**New FDA Medical Device Inspection Model effective February 2, 2026**
4 días (1:00 pm – 5:00 pm) TBD	**Quality Risk Management Certification** (14 horas)
9:30 am – 11:30 am TBD	**Drug GMP Overview/Refresher 2026** (2 horas)
1 día (9:00 am – 5:00 pm) TBD	**Train-the Trainer**
1 día (9:00 am – 5:00 pm) TBD	**Knowledge Management**
2 días (1:00 pm – 5:00 pm) TBD	**Sampling Best Practices** (7 horas)
2 días (1:00 pm – 5:00 pm) TBD	**ISO 13485:2016** (7 horas)
3 días (8:30 am – 5:00 pm) TBD	**ASQ Certified Six Sigma Green Belt (CSSGB)** (21 horas)
3 sábados (8:30 am – 5:00 pm) TBD	**ASQ Certified Manager of Quality and Organizational Excellence (CMQ-OE)** (21 horas)

Para reservar https://bec-global.com/training-reservation-form/ o puedes escribir a training@calidadpr.com

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Cursos Gratuitos (horario 10 am – 11 am PR / AST time)

Noviembre 24, 2026	**Cosmetics Regulations: Mandatory Recall and Adverse Event Reporting**
Diciembre 15, 2026	**FDA Regulatory Enforcement FY 2026**
Diciembre 22, 2026	**IA and cGMP: Dos and Don'ts**

Otros cursos disponibles

Investigation, CAPA, and Human Errors Reduction Certification (CERT-025) (28 horas)
Managerial Decision Making (7 horas)
Best Practices Stability Programs for the Pharmaceutical Industry (14 horas)
Environmental Monitoring Deviations (7 horas) - Microbial Investigations Best Practices (7 horas)
Implementing a Mature and sustainable Investigations and CAPA Program (4 horas)
Understanding FDA Devices Manufacturing Inspections (7 horas)
Understanding FDA DRUGS Manufacturing Inspections (7 horas)
ISO 14971:2019 (7 horas)