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Calendario de Talleres

Horarios de Puerto Rico – AST (Atlantic Standard Time) – Actualizado 11-20-2026

Fechas	VIRTUAL Training
2 días (8:30 am – 5:00 pm) Enero 25 y 27, 2027	**Root Cause Analysis** (14 horas)
2 días (8:30 am – 5:00 pm) Enero 26 y 28, 2027	**Effective Compliance and Regulatory Writing** (14 horas)
3 sábados (8:30 am – 5:00 pm) Enero 30, Febrero 6 y 13, 2027	**ASQ Certified Quality Auditor CQA** (21 horas)
4 días (8:00 am – 12:00 pm) Febrero 1, 3, 8 y 10, 2027	**Quality Risk Management Certification** (14 horas)
4 días (8:00 am – 12:00 pm) Febrero 2, 4, 9 y 11, 2027	**Basic Applied Statistics** (14 horas)
1 día (8:30 am – 5:00 pm) Febrero 5, 2027	**ISO 9001:2026** (7 horas)
3 sábados (8:30 am – 5:00 pm) Febrero 27, Marzo 6 y 13, 2027	**ASQ Certified Six Sigma Green Belt (CSSGB)** (21 horas)
3 sábados (8:30 am – 5:00 pm) Febrero 27, Marzo 6 y 13, 2027	**ASQ Certified Manager of Quality and Organizational Excellence (CMQ-OE)** (21 horas)
2 días (1:00 pm – 5:00 pm) Marzo 8 y 10, 2027	**Sampling Best Practices** (7 horas)
2 días (1:00 pm – 5:00 pm) Marzo 9 y 11, 2027	**Critical Thinking for Investigators** (7 horas)
5 sábados (8:30 am – 5:00 pm) Abril 3, 10, 17, 24 y Mayo 1, 2027	**ASQ Certified Quality Engineer CQE** (35 horas)
6 días (8:00 am – 12:00 pm) Abril 5, 7, 9, 12, 14 y 16, 2027	**Effective Internal Auditing Certification** (21 horas)
4 días (8:00 am – 12:00 pm) Abril 6, 8, 13 y 15, 2027	**Human Errors Investigation** (14 horas)
1 día (9:00 am – 5:00 pm) TBD	**New FDA Medical Device Inspection Model effective February 2, 2026** (7 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** (7 horas)

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

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1 día (9:00 am – 5:00 pm) TBD	**Knowledge Management** (7 horas)
2 días (1:00 pm – 5:00 pm) TBD	**ISO 13485:2016** (7 horas)

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)

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