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Calendario de Talleres

Horarios de Puerto Rico – AST (Atlantic Standard Time) – Actualizado 09-22-2026

Fechas	VIRTUAL Training
5 sábados (8:30 am – 5:00 pm) Octubre 10, 17, 24, 31 y Nov. 7, 2026	ASQ Certified Quality Engineer CQE (35 horas)
1 día (9:00 am – 5:00 pm) Octubre 13, 2026	New FDA Medical Device Inspection Model effective February 2, 2026
1 día (9:00 am – 5:00 pm) Octubre 15 (nueva fecha) Presencial oficinas BEC	Train the Trainer (8 horas)
1 día (9:00 am – 5:00 pm) Octubre 22 - Presencial	Knowledge Management (8 horas)
2 días (8:00 am – 12:00 pm) Octubre 20 y 22, 2026	ISO 9001:2026 (7 horas) New version of the standard
2 días (1:00 pm – 5:00 pm) Octubre 20 y 22, 2026	ISO 13485:2016 (7 horas)
3 días (8:30 am – 5:00 pm) Octubre 26, 28 y 30, 2026	ASQ Certified Six Sigma Green Belt (CSSGB) (21 horas)
3 días (8:30 am – 5:00 pm) Noviembre 2, 4 y 6, 2026	Effective Internal Auditing Certification (21 horas)
3 días (8:30 am – 5:00 pm) Noviembre 9, 11 y 13, 2026	ASQ Certified Supplier Quality Professional (CSQP) (21 horas)
2 días (8:00 am – 12:00 pm) Noviembre 10 y 12, 2026	Data Trending Analysis (7 horas)
2 días (1:00 pm – 5:00 pm) Noviembre 10 y 12, 2026	Sampling Best Practices (7 horas)
3 sábados (8:30 am – 5:00 pm) TBD	ASQ Certified Manager of Quality and Organizational Excellence (CMQ-OE) (21 horas)
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)

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



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2 días (1:00 pm – 5:00 pm) TBD	New 21 CFR 820 - Quality Management System Regulation (8 horas)
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Cursos Gratuitos (horario 10 am – 11 am PR / AST time)

Octubre 27, 2026	Human Error Investigations
Noviembre 24, 2026	Cosmetics Regulations: Mandatory Recall and Adverse Event Reporting
Diciembre 15, 2026	FDA Regulatory Enforcement FY 2026
TBD	IA and cGMP: Dos and don'ts

Otros cursos disponibles

Investigation, CAPA, and Human Error Reduction Certification (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)