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Calendario de Talleres 2026

Horarios de Puerto Rico – AST (Atlantic Standard Time) – Actualizado 01-08-2026

Fechas	VIRTUAL Training
4 días (1:00 pm – 5:00 pm) Enero 19, 21, 26 y 28	**Quality Risk Management Certification** **(14 horas)**
4 días (1:00 pm – 5:00 pm) Enero 20, 22, 27 y 29	**Basic Applied Statistics** (14 horas)
3 sábados (8:30 am – 5:00 pm) Enero 24, 31 y Febrero 7	**ASQ Certified Quality Auditor CQA (21 horas)**
2 dias (1:00 – 5:00 pm) Enero 26 y 28	**21 CFR 820 Update: The New Quality Management System Regulation (8 horas)**
6 dias (1:00 pm – 5:00 pm) Febrero 2, 4, 6, 9, 11 y 13	**Effective Internal Auditing Certification** (21 horas)
4 días (1:00 pm – 5:00 pm) Febrero 10 y 12	**ISO 9001:2015** (7 horas)
1 día (9:00 am – 5:00 pm) Febrero 26	**How to Measure Training Effectiveness** (7 horas) **Presencial – oficina de BEC en Bayamón**
4 días (1:00 pm – 5:00 pm) **EST** March 2, 4, 9 & 11	**Human Error Investigation** (14 hrs) **Offered in English**
4 días (8:00 am – mediodia) Marzo 3, 5, 10 y 12	**Root Cause Analysis** (14 hours)
4 días (1:00 pm – 5:00 pm) Marzo 3, 5, 10 y 12	**Effective Compliance and Regulatory Writing** (14 horas)
5 sábados (8:30 am – 5:00 pm) Febrero 28, Marzo 7, 14, 21 y 28	**ASQ Certified Quality Engineer CQE** (35 horas)
3 sábados (8:30 am – 5:00 pm) Abril 11, 18 y 25	**ASQ Certified Manager of Quality and Organizational Excellence (CMQ-OE)** (21 horas)
3 sábados (8:30 am – 5:00 pm) Mayo 16, 23 y 30	**ASQ Certified Six Sigma Green Belt (CSSGB)**
Cursos Gratuitos	
Enero 21: 10 am – 11 am	**FY2025 Enforcement – FDA Inspectional Observations**

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

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Otros cursos disponibles

Investigation, CAPA, and Human Error Reduction Certification (28 horas)
Managerial Decision Making (7 horas) - Knowledge Management (7 horas)
Train the Trainer (7 horas) - ISO 13485: 2016 (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 17025: 2017 (7 horas) - ISO 14971:2019 (7 horas)