Business Excellence Consulting, Inc.
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Calendario Talleres 2025

Fechas	VIRTUAL Training	Costo*
2 días (8:30 am – 5:00 pm) TBD	**Effective Compliance and Regulatory Writing** (14 horas)	$750
4 dias (8:30am – 5:00 pm) Febrero 10, 12, 17 y 19 2025	**Investigation/CAPA System and Human Errors Reduction Certification** (28 horas)	$995
5 sábados (8:30 am–5:00 pm) Ene 25, Feb 1, 8, 15 y 22 2025	**ASQ Certified Quality Engineer CQE** (35 horas)	$1095
1 día (9:00 am – 5:00 pm) Enero 27, 2025	**A**ctualización de **MDSAP** **(**Medical Device Single Audit Program) 7 horas	$550
1 día (9:00 am – 5:00 pm) Enero 29, 2025	**Understanding FDA Drug Manufacturing Inspections:** (7 horas)	$195
1 día (9:00am – 5:00 pm) Febrero 11, 2025	**cGMP for Managers and Supervisors – Managerial Responsibilities** (7 horas)	$550
3 días (8:30 am – 5:00 pm) Febrero 24, 26 y 28, 2025	**Internal Audit Certification** (21 horas)	$800
3 sábados (8:30 am - 5:00 pm) Marzo 1, 8, y 15, 2025	**ASQ Certified Quality Auditor (CQA)** (21 horas)	$950
1 día (9:00 am – 5:00 pm) Marzo 3, 2025	**ISO 13485: 2016 (7 horas)**	$550
1 día (9:00 am – 5:00 pm) Marzo 4, 2025	**ISO 17025: 2017 (7 horas)**	$550
1 día (9:00 am – 5:00 pm) Marzo 6, 2025	**ISO 9001:2015 (7 horas)**	$550
2 días (8:30 am – 5:00 pm) Marzo 10 y 11, 2025	**Quality Risk Management** (14 horas)	$750
1 día (9:00 am – 5:00 pm) Marzo 13, 2025	**Process Validation - Lifecycle Approach for FDA Medical Products** (7 horas)	$550
3 sábados (8:30 am - 5:00 pm) Abril 5, 12 y 26, 2025	**ASQ Certified Six Sigma Green Belt** (21 horas)	$950
CURSOS PRESENCIALES en nuestras oficinas de Bayamón		
2 días (9:00 am – 5:00 pm) Febrero 13 y 14	**Escritura de Procedimientos para Minimizar los Errores Humanos** (14 horas)	$900
2 días (9:00 am – 5:00 pm) Febrero 17 y 18	**CSV – Computer System Validation (14 horas)**	$900
3 días (9:00 am – 5:00 pm) Marzo 3, 4 y 5	**Data Integrity Certification**	$1,200
4 days (8:30 am – 6:00 pm) TBD	**ISO 9001:2015 LEAD AUDITOR** **(VIRTUAL, 36 contact-hours) in English**	$1,990
4 days (8:30 am – 6:00 pm) TBD	**ISO 13485:2016 LEAD AUDITOR** **(VIRTUAL, 36 contact-hours) in English**	$1,990

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2 días (8:30 am – 5:00 pm) TBD	**Basic Applied Statistics** (14 horas)	$750
2 días (8:30 am – 5:00 pm)	**Root Cause Analysis** (14 horas)	$695
2 días (9:00 am – 5:00 pm)	**Best practices Stability Programs for the Pharmaceutical Industry** (14 horas)	$750
4 horas (9:00 am – 1:00 pm)	**Laboratory OOS Investigation and Invalidation Best Practices** (4 horas)	$395
1 día (9:00 am – 5:00 pm)	**ISO 14971:2019** (7 horas)	$550
1 día (8:30am – 5:00 pm)	**Critical Thinking** (7 horas)	$550
1 día (8:30am – 5:00 pm)	**Decision Making** (7 horas)	$550
1 día (8:30 am – 5:00 pm)	**Environmental Monitoring Deviations** (7 horas)	$550
1 día (8:30 am – 5:00 pm)	**Microbial Investigations Best Practices** (7 horas)	$550
4 horas (9:00 am – 1:00 pm)	**Implementing a Mature and sustainable Investigations and CAPA Program** (4 horas)	$395
1 día (8:30am – 5:00 pm)	**Understanding FDA Devices Manufacturing Inspections** (7 horas)	$550
SEMINARIOS GRATUITOS 2025		
Enero 30, 2025 (11 am –mediodia AST)	**#2 FY 2024 US FDA Drug and Devices Inspection Compliance**	
Marzo 5, 2024 (11 am-mediodia AST)	**#3 Data Integrity Investigations**	
Marzo 12, 2025 (11 am – mediodía AST)	**#4 Investigations and CAPA Pitfalls**	
Abril 2, 2025 10 am – 12 mediodia AST	**#5 Actualización de MOCRA+ Proposición 65 de California, PFAs, y Colorantes permitidos en Cosméticos por la FDA**	
TBD	**#6 Complaint Investigations Best Practices**	
TBD	**#7 21 CFR 820 Update: The New Quality Management System Regulation**	

más impuestos si aplica Horarios de Puerto Rico 1/20/2025

Para reservar https://bec-global.com/training-reservation-form/

o puedes escribir a training@calidadpr.com