Business Excellence Consulting, Inc.
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Calendario Talleres 2025

Fechas	VIRTUAL Training	Costo*
2 sábados (8:30 am – 5:00 pm) Mayo 31 & Junio 7, 2025	**Human Error Investigations** (14 horas)	$750
4 dias (8:30am – 5:00 pm) Junio 3, 5, 10 & 12, 2025	**Investigation/CAPA System and Human Errors Reduction Certification** (28 horas)	$1,100
4 days (8:30 am – 6:00 pm) June 9, 10, 11 & 12, 2025	**ISO 13485:2016 LEAD AUDITOR (VIRTUAL, 36 contact-hours) in English**	$1,990
Junio 27 (9:00 am – 5:00 pm)	**21 CFR 820 Update: The New Quality Management System Regulation**	$550
3 sábados (8:30 am – 5:00 pm) Agosto 9, 16 & 23, 2025	**ASQ Certified Quality Auditor CQA** (21 horas)	$950
5 sábados (8:30 am – 5:00 pm) Septiembre 6, 13, 20, 27 & Octubre 4, 2025	**ASQ Certified Quality Engineer CQE** (35 horas)	$1095
2 dias (8:30 am – 12:30 pm) TBD	**Knowledge Management** (7 horas)	$550
2 días (8:30 am – 12:30 pm) TBD	**How to Measure Training Effectiveness** (7 horas)	$550
2 días (8:30 am – 12:30 pm) TBD	**Train the Trainer** (7 horas)	$550
2 sábados (8:30 am – 5:00 pm) TBD	**Root Cause Analysis** (14 horas)	$750
2 días (8:30 am – 5:00 pm) TBD	**Quality Risk Management** (14 horas)	$750
1 día (8:30 am – 5:00 pm) TBD	**ISO 9001:2015 (7 horas)**	$550
1 día (8:30 am – 5:00 pm) TBD	**ISO 13485: 2016 (7 horas)**	$550
4 days (8:30 am – 6:00 pm) TBD	**ISO 9001:2015 LEAD AUDITOR (VIRTUAL, 36 contact-hours) in English**	$1,990
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 (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

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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
1 día (8:30am – 5:00 pm)	**Understanding FDA DRUGS Manufacturing Inspections** (7 horas)	$550
2 días (8:30 am – 5:00 pm)	**Basic Applied Statistics** (14 horas)	$750
1 día (9:00 am – 5:00 pm)	**ISO 17025: 2017 (7 horas)**	$550
1 día (9:00 am – 5:00 pm)	**ISO 14971:2019** (7 horas)	$550
SEMINARIOS GRATUITOS 2025		
Mayo 20, 2025 10 am – 11 am AST	**#9 MDSAP Overview**	
Mayo 29, 2025 10 am – 11 am AST	**#10 Laboratory OOS Invalidation**	
Junio 4, 2025 10 am – 11 am AST	**#11 Fraudulent Drugs: You're Taking What?**	
Junio 17, 2025 10 am – 11 am AST	**#12 An introduction to FDA Regulations of Medical Devices**	
Julio 15, 2025 10 am – 11 am AST	**#13 Drug Shortages: root Causes and Solutions**	
Agosto 6, 2025 10 am – 11 am AST	**#14 FDA Drug Regulation Pathways**	
Septiembre 16, 2025 10 am – 11 am AST	**#15 Questions & Answers on FDA Dietary Supplements**	
TBD	**#16 MedWatch: The FDA Safety Information & Adverse Event Reporting Program**	
TBD	**#17 FDA History**	
TBD	**#18 Quality Management Maturity and Quality Culture**	
TBD	**#19 Managerial Decision Making**	
TBD	**#20 FDA Regulatory Enforcement Fiscal Year 2025**	

más impuestos si aplica

Horarios de Puerto Rico – AST (Atlantic Standard Time) 05/19/2025

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

o puedes escribir a training@calidadpr.com