



Business Excellence Consulting, Inc.  
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## Calendario Talleres 2025

<b>Fechas</b>	<b>VIRTUAL Training</b>	<b>Costo*</b>
2 días (8:30 am – 5:00 pm) Enero 13 y 14, 2025	<b>Human Error Investigations</b> (14 horas)	\$750
1 día (9:00am – 4:00 pm) Enero 15, 2025	<b>Knowledge Management: Pitfalls of the Training Programs in the FDA-Regulated Companies</b> (6 horas)	Gratis
2 días (8:30 am – 5:00 pm) Enero 15 y 16, 2025	<b>Effective Compliance and Regulatory Writing</b> (14 horas)	\$750
4 días (8:30am – 5:00 pm) Enero 21, 23, 28 y 30, 2025	<b>Investigation/CAPA System and Human Errors Reduction Certification</b> (28 horas)	\$995
5 sábados (8:30 am–5:00 pm) Ene 25, Feb 1, 8, 15 y 22 2025	<b>ASQ Certified Quality Engineer CQE</b> (35 horas)	\$1095
1 día (9:00 am – 5:00 pm) Enero 27, 2025	<b>Actualización de MDSAP</b> (Medical Device Single Audit Program) 7 horas	\$550
1 día (9:00 am – 5:00 pm) Enero 29, 2025	<b>Understanding FDA Drug Manufacturing Inspections:</b> (7 horas)	\$195
1 día (9:00am – 5:00 pm) Febrero 11, 2025	<b>cGMP for Managers and Supervisors – Managerial Responsibilities</b> (7 horas)	\$550
3 días (8:30 am – 5:00 pm) Febrero 17, 19 y 21, 2025	<b>Internal Audit Certification</b> (21 horas)	\$800
3 sábados (8:30 am - 5:00 pm) Marzo 1, 8, y 15, 2025	<b>ASQ Certified Quality Auditor (CQA)</b> (21 horas)	\$950
1 día (9:00 am – 5:00 pm) Marzo 3, 2025	<b>ISO 13485: 2016 (7 horas)</b>	\$550
1 día (9:00 am – 5:00 pm) Marzo 4, 2025	<b>ISO 17025: 2017 (7 horas)</b>	\$550
1 día (9:00 am – 5:00 pm) Marzo 6, 2025	<b>ISO 9001:2015 (7 horas)</b>	\$550
2 días (8:30 am – 5:00 pm) Marzo 10 y 11, 2025	<b>Basic Applied Statistics</b> (14 horas)	\$750
2 días (8:30 am – 5:00 pm) Marzo 12 y 13, 2025	<b>Quality Risk Management</b> (14 horas)	\$750
1 día (9:00 am – 5:00 pm) Marzo 13, 2025	<b>Process Validation - Lifecycle Approach for FDA Medical Products</b> (7 horas)	\$550
3 sábados (8:30 am - 5:00 pm) Abril 5, 12 y 26, 2025	<b>ASQ Certified Six Sigma Green Belt</b> (21 horas)	\$950
<b>CURSOS PRESENCIALES en nuestras oficinas de Bayamón</b>		
2 días (9:00 am – 5:00 pm) Febrero 13 y 14	<b>Escritura de Procedimientos para Minimizar los Errores Humanos</b> (14 horas)	\$900
2 días (9:00 am – 5:00 pm) Febrero 17 y 18	<b>CSV – Computer System Validation</b> (14 horas)	\$900



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3 días (9:00 am – 5:00 pm) Marzo 3, 4 y 5	<b>Data Integrity Certification</b>	\$1,200
4 days (8:30 am – 6:00 pm) TBD	<b>ISO 9001:2015 LEAD AUDITOR (VIRTUAL, 36 contact-hours) in English</b>	\$1,990
4 days (8:30 am – 6:00 pm) TBD	<b>ISO 13485:2016 LEAD AUDITOR (VIRTUAL, 36 contact-hours) in English</b>	\$1,990
2 días (8:30 am – 5:00 pm)	<b>Root Cause Analysis (14 horas)</b>	\$695
3 días (9:00 am – 5:00 pm)	<b>Data Integrity Certification (21 horas)</b>	\$995
2 días (9:00 am – 5:00 pm)	<b>Best practices Stability Programs for the Pharmaceutical Industry (14 horas)</b>	\$750
4 horas (9:00 am – 1:00 pm)	<b>Laboratory OOS Investigation and Invalidation Best Practices (4 horas)</b>	\$395
1 día (9:00 am – 5:00 pm)	<b>ISO 14971:2019 (7 horas)</b>	\$550
1 día (8:30am – 5:00 pm)	<b>Critical Thinking (7 horas)</b>	\$550
1 día (8:30am – 5:00 pm)	<b>Decision Making (7 horas)</b>	\$550
1 día (8:30 am – 5:00 pm)	<b>Environmental Monitoring Deviations (7 horas)</b>	\$550
1 día (8:30 am – 5:00 pm)	<b>Microbial Investigations Best Practices (7 horas)</b>	\$550
4 horas (9:00 am – 1:00 pm)	<b>Implementing a Mature and sustainable Investigations and CAPA Program (4 horas)</b>	\$395
1 día (8:30am – 5:00 pm)	<b>Understanding FDA Devices Manufacturing Inspections (7 horas)</b>	\$550
<b>SEMINARIOS GRATUITOS 2025</b>		
Enero 30, 2025 (11 am –mediodía AST)	<b>#1 FY 2024 US FDA Drug Inspection Compliance</b>	
Febrero 12, 2025 (11:00 am-mediodía AST)	<b>#2 FY 2024 US FDA Device Inspection Compliance</b>	
Marzo 5, 2024 (11 am-mediodía AST)	<b>#3 Data Integrity Investigations</b>	
Marzo 12, 2025 (11 am – mediodía AST)	<b>#4 Investigations and CAPA Pitfalls</b>	
Abril 2, 2025 10 am – 12 mediodía AST	<b>#5 Actualización de MOCRA+ Proposición 65 de California, PFAs, y Colorantes permitidos en Cosméticos por la FDA</b>	
TBD	<b>#6 Complaint Investigations Best Practices</b>	
TBD	<b>#7 21 CFR 820 Update: The New Quality Management System Regulation</b>	

más impuestos si aplica

Horarios de Puerto Rico

12/06/2024

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o puedes escribir a [training@calidadpr.com](mailto:training@calidadpr.com)