



Business Excellence Consulting, Inc.  
P.O. Box 8326, Bayamón, PR 00960-8326  
training@calidadpr.com  
info@calidadpr.com

Phone: 787.705.7272  
Fax 787.705.5272  
[www.bec-global.com](http://www.bec-global.com)  
[www.calidadpr.com](http://www.calidadpr.com)

## Calendario de Talleres

Horarios de Puerto Rico – AST (Atlantic Standard Time) – Actualizado 05-07-2026

<b>Fechas</b>	<b>VIRTUAL Training</b>
3 sábados (8:30 am – 5:00 pm) Mayo 30, Junio 6 & 13, 2026	<b>ASQ Certified Six Sigma Green Belt (CSSGB)</b> (21 horas)
3 días (8:30 am – 5:00 pm) Junio 10, 11 y 12, 2026	<b>ASQ Certified Quality Auditor (CQA)</b> (21 horas)
3 sábados (8:30 am – 5:00 pm) Agosto 15, 22 y 29, 2026	<b>ASQ Certified Supplier Quality Professional (CSQP)</b> (21 horas)
4 días (8:00 am – 12:00 pm) Agosto 25, 26 ,27 y 28, 2026	<b>Effective Compliance and Regulatory Writing</b> (14 horas)
4 días (1:00 pm – 5:00 pm) Agosto 25, 26 ,27 y 28, 2026	<b>Root Cause Analysis</b> (14 hours)
3 sábados (8:30 am – 5:00 pm) Septiembre 12, 19 y 26, 2026	<b>ASQ Certified Quality Auditor (CQA)</b> (21 horas)
4 días (8:00 am – 12:00 pm) Sept. 22, 23, 24 y 25, 2026	<b>Basic Applied Statistics</b> (14 hours)
4 días (1:00 pm – 5:00 pm) Sept. 22, 23, 24 y 25, 2026	<b>Quality Risk Management Certification</b> (14 hours)
5 sábados (8:30 am – 5:00 pm) Octubre 10, 17, 24, 31 y Nov 7, 2026	<b>ASQ Certified Quality Engineer CQE</b> (35 horas)
3 días (8:30 am – 5:00 pm) Octubre 19, 21 y 23, 2026	<b>Effective Internal Auditing Certification</b> (21 horas)
2 días (8:00 am – 12:00 pm) Octubre 20 y 22, 2026	<b>ISO 9001:2015</b> (7 horas)
2 días (1:00 pm – 5:00 pm) Octubre 20 y 22, 2026	<b>ISO 13485:2016</b> (7 horas)
2 días (8:00 am – 12:00 pm) Noviembre 10 y 12, 2026	<b>Data Trending Analysis</b> (7 horas)



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2 días (1:00 pm – 5:00 pm) Noviembre 10 y 12, 2026	<b>Sampling Best Practices (7 horas)</b>
2 días (1:00 – 5:00 pm) TBD	<b>21 CFR 820 Update: The New Quality Management System Regulation (8 horas)</b>
3 sábados (8:30 am – 5:00 pm) TBD	<b>ASQ Certified Manager of Quality and Organizational Excellence (CMQ-OE) (21 horas)</b>

### 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)  
 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)