



Business Excellence Consulting, Inc.  
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## Calendario Talleres 2025

| Fechas                                                    | VIRTUAL Training                                                                          | Costo*  |
|-----------------------------------------------------------|-------------------------------------------------------------------------------------------|---------|
| 3 sábados (8:30 am – 5:00 pm)<br>TBD                      | <b>ASQ Certified Supplier Quality Professional (CSQP)</b><br>(21 horas)                   | \$950   |
| 1 día (8:30 am – 5:00 pm)<br>Septiembre 8, 2025           | <b>ISO 9001:2015 (7 horas)</b>                                                            | \$550   |
| 4 horas (1:00 pm – 5:00 pm)<br>Septiembre 18, 2025        | <b>Laboratory OOS Investigation and Invalidation Best Practices (4 horas)</b>             | \$395   |
| 3 sábados (8:30 am – 5:00 pm)<br>Octubre 4, 11 & 18, 2025 | <b>ASQ Certified Manager of Quality and Organizational Excellence (CMQ-OE) (21 horas)</b> | \$950   |
| 2 días (8:30 am – 5:00 pm)<br>Octubre 7 & 8, 2025         | <b>Basic Applied Statistics (14 horas)</b>                                                | \$750   |
| 2 días (8:30 am – 5:00 pm)<br>Octubre 9 & 10, 2025        | <b>Effective Compliance and Regulatory Writing (14 horas)</b>                             | \$750   |
| 3 días (8:30 am – 5:00 pm)<br>Octubre 28, 29 & 30, 2025   | <b>Effective Internal Auditing (21 horas)</b>                                             | \$950   |
| 4 días (8:30am – 5:00 pm)<br>Noviembre 3, 4, 5 & 6, 2025  | <b>Investigation/CAPA System and Human Errors Reduction Certification (28 horas)</b>      | \$1,100 |
| 3 sábados (8:30 am – 5:00 pm)<br>TBD                      | <b>ASQ Certified Quality Auditor CQA (21 horas)</b>                                       | \$950   |
| 5 sábados (8:30 am – 5:00 pm)<br>TBD                      | <b>ASQ Certified Quality Engineer CQE (35 horas)</b>                                      | \$1095  |
| 2 días (8:30 am – 5:00 pm)<br>TBD                         | <b>Quality Risk Management (14 horas)</b>                                                 | \$750   |
| 2 días (8:30 am – 5:00 pm)<br>TBD                         | <b>Root Cause Analysis (14 horas)</b>                                                     | \$750   |
| 2 días (8:30 am – 5:00 pm)<br>TBD                         | <b>Human Error Investigations (14 horas)</b>                                              | \$750   |
| 4 days (8:30 am – 6:00 pm)<br>TBD                         | <b>ISO 13485:2016 LEAD AUDITOR (VIRTUAL, 36 contact-hours) in English</b>                 | \$1,990 |
| 1 día (9:00 am – 5:00 pm)<br>TBD                          | <b>Managerial Decision Making (7 horas)</b>                                               | \$550   |



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|------------------------------------------|--------------------------------------------------------------------------------------------|---------|
|                                          |                                                                                            |         |
| 1 día (9:00 am – 5:00 pm)<br>TBD         | <b>21 CFR 820 Update:<br/>The New Quality Management System Regulation</b>                 | \$550   |
| 1 día (9:00 am – 5:00 pm)<br>TBD         | <b>Complaint Investigation Best Practices (7 horas)</b>                                    | \$550   |
| 2 días (8:30 am – 12:30 pm)<br>TBD       | <b>Knowledge Management (7 horas)</b>                                                      | \$550   |
| 2 días (8:30 am – 12:30 pm)<br>TBD       | <b>How to Measure Training Effectiveness (7 horas)</b>                                     | \$550   |
| 2 días (8:30 am – 12:30 pm)<br>TBD       | <b>Train the Trainer (7 horas)</b>                                                         | \$550   |
| 4 days (8:30 am – 6:00 pm)<br>TBD        | <b>ISO 9001:2015 LEAD AUDITOR<br/>(VIRTUAL, 36 contact-hours) in English</b>               | \$1,990 |
| 2 días (9:00 am – 5:00 pm)               | <b>Best Practices Stability Programs for the<br/>Pharmaceutical Industry (14 horas)</b>    | \$750   |
| 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<br/>and CAPA Program (4 horas)</b> | \$395   |
| 1 día (8:30am – 5:00 pm)                 | <b>Understanding FDA Devices Manufacturing Inspections<br/>(7 horas)</b>                   | \$550   |
| 1 día (8:30am – 5:00 pm)                 | <b>Understanding FDA DRUGS Manufacturing Inspections<br/>(7 horas)</b>                     | \$550   |
| 1 día (9:00 am – 5:00 pm)                | <b>ISO 17025: 2017 (7 horas)</b>                                                           | \$550   |
| 1 día (9:00 am – 5:00 pm)                | <b>ISO 14971:2019 (7 horas)</b>                                                            | \$550   |
| <b>SEMINARIOS GRATUITOS 2025</b>         |                                                                                            |         |
| Septiembre 16, 2025<br>10 am – 11 am AST | <b>#15 Questions &amp; Answers on FDA Dietary Supplements</b>                              |         |
| TBD<br>10 am – 11 am AST                 | <b>#12 An introduction to FDA Regulations of Medical Devices</b>                           |         |
| TBD                                      | <b>#16 MedWatch: The FDA Safety Information &amp;<br/>Adverse Event Reporting Program</b>  |         |
| TBD                                      | <b>#17 FDA History</b>                                                                     |         |



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|     |                                                            |
|-----|------------------------------------------------------------|
| TBD | <b>#18 Quality Management Maturity and Quality Culture</b> |
| TBD | <b>#19 Managerial Decision Making</b>                      |
| TBD | <b>#20 FDA Regulatory Enforcement Fiscal Year 2025</b>     |

más impuestos si aplica

Horarios de Puerto Rico – AST (Atlantic Standard Time) 9/3/2025

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

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