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## Calendario de Talleres

Horarios de Puerto Rico – AST (Atlantic Standard Time) – Actualizado 10-22-2026

| <b>Fechas</b>                                                           | <b>VIRTUAL Training</b>                                                                           |
|-------------------------------------------------------------------------|---------------------------------------------------------------------------------------------------|
| 3 días (8:30 am – 5:00 pm)<br>Noviembre 9, 16, 23 y 30, 2026            | <b>Investigation, CAPA, and Human Errors<br/>Reduction Certification (CERT-025)</b><br>(28 horas) |
| 2 días (8:00 am – 12:00 pm)<br>Noviembre 10 y 12, 2026                  | <b>Data Trending Analysis</b> (7 horas)                                                           |
| 2 días (1:00 pm – 5:00 pm)<br>Noviembre 10 y 12, 2026                   | <b>Sampling Best Practices</b> (7 horas)                                                          |
| 1 día (8:30 am – 5:00 pm)<br>Noviembre 17, 2026                         | <b>Critical Thinking for Investigators</b>                                                        |
| 1 día (9:00 am – 5:00 pm)<br>Noviembre 18, 2026                         | <b>New FDA Medical Device Inspection Model<br/>effective February 2, 2026</b>                     |
| 3 días (8:30 am – 5:00 pm)<br>Diciembre 14, 16 y 18, 2026               | <b>Effective Internal Auditing Certification</b><br>(21 horas)                                    |
| 3 sábados (8:30 am – 5:00 pm)<br>Enero 30, Febrero 6 y 13, 2027         | <b>ASQ Certified Quality Auditor CQA</b> (21 horas)                                               |
| 5 sábados (8:30 am – 5:00 pm)<br>Febrero 27, Marzo 6, 13, 20 y 27, 2027 | <b>ASQ Certified Quality Engineer CQE</b> (35 horas)                                              |
| 4 días (1:00 pm – 5:00 pm)<br>TBD                                       | <b>Quality Risk Management Certification</b> (14 horas)                                           |
| 9:30 am – 11:30 am<br>TBD                                               | <b>Drug GMP Overview/Refresher 2026</b> (2 horas)                                                 |
| 1 día (9:00 am – 5:00 pm)<br>TBD                                        | <b>Train-the Trainer</b>                                                                          |
| 1 día (9:00 am – 5:00 pm)<br>TBD                                        | <b>Knowledge Management</b>                                                                       |
| 2 días (1:00 pm – 5:00 pm)<br>TBD                                       | <b>ISO 13485:2016</b> (7 horas)                                                                   |
| 3 días (8:30 am – 5:00 pm)<br>TBD                                       | <b>ASQ Certified Six Sigma Green Belt (CSSGB)</b><br>(21 horas)                                   |
| 3 sábados (8:30 am – 5:00 pm)<br>TBD                                    | <b>ASQ Certified Manager of Quality and Organizational<br/>Excellence (CMQ-OE)</b> (21 horas)     |

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### **Cursos Gratuitos** (horario 10 am – 11 am PR / AST time)

|                    |                                                                            |
|--------------------|----------------------------------------------------------------------------|
| Octubre 27, 2026   | <b>Human Error Investigations</b>                                          |
| Noviembre 24, 2026 | <b>Cosmetics Regulations: Mandatory Recall and Adverse Event Reporting</b> |
| Diciembre 15, 2026 | <b>FDA Regulatory Enforcement FY 2026</b>                                  |
| Diciembre 22, 2026 | <b>IA and cGMP: Dos and Don'ts</b>                                         |

### **Otros cursos disponibles**

Managerial Decision Making (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 14971:2019 (7 horas)