

ComplianceOnline to Host Seminar On 21 CFR Part 11 Compliance for SaaS/Cloud Applications

"21 CFR Part 11 Compliance for SaaS/Cloud Applications" Seminar has been added to ComplianceOnline.com's offering.

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EINPresswire.com/ --

ComplianceOnline, the leading provider of training for regulated companies, is all set to host an educational seminar on the topic 21 CFR Part 11 Compliance for SaaS/Cloud Applications geared toward professionals working in the health care, clinical trial, biopharmaceutical, and medical device sectors. It is also essential for software vendors, auditors, and quality staff involved in GxP applications. The December 8-9, 2021, includes two-day sessions featuring FDA Compliance Specialist David Nettleton discussing real-life examples and proven techniques for reducing costs, usually by two-thirds, associated with implementing and maintaining computer systems in regulated environments.



The graphic is a promotional poster for a seminar. It features a dark background with a keyboard and a headset. The text is white and yellow. At the top right is the ComplianceOnline logo. Below it is a yellow banner that says '2-Day Virtual Seminar'. The main title is '21 CFR Part 11 Compliance for SaaS/Cloud Applications'. Below that is the name 'David Nettleton, FDA Compliance Specialist, Computer System Validation'. At the bottom is a yellow banner with a calendar icon and the text 'December 8-9, 2021 (9:00 AM - 3:00 PM PST) Virtual Training Through WebEx'. The background also has 'SaaS' and 'ComplianceOnline' faintly visible.

21 CFR Part 11 Compliance for SaaS/Cloud Applications

- The seminar details the requirements for Part 11 and Annex 11: SOPs, software product features, infrastructure qualification, and validation.
- The instructor addresses the latest computer system industry standards for data security, data transfer, audit trails, electronic records and signatures, software validation, and computer system validation.
- Attendees will understand the specific requirements associated with local and SaaS/cloud hosting solutions.
- Nearly every computerized system used in laboratory, clinical, manufacturing settings and in the quality process has to be validated. Participants will learn how to decrease software implementation times and lower costs using a 10-step risk-based approach to computer system

validation.

- The instructor reviews recent FDA inspection trends and discusses how to streamline document authoring, revision, review, and approval.
- Participants will learn how to write a Data Privacy Statement to comply with the EU General Data Protection Regulation (GDPR).

The seminar's learning objectives according to the event overview are to:

- Understand what is expected in Part 11 and Annex 11 inspections so you are prepared
- Avoid 483 and Warning Letters
- Understand the responsibilities and specific duties of your staff including IT and QA
- Understand responsibilities and liabilities when using SaaS/cloud
- Learn how to perform risk-based Computer System Validation using fill-in-the-blank templates
- How to select resources and manage validation projects
- "Right size" change control methods that allow quick and safe system evolution
- Minimize the validation documentation to reduce costs without increasing regulatory or business risk
- Learn how to reduce testing time and write test cases that trace to elements of risk management
- Learn how to comply with the requirements for data privacy
- Learn how to buy COTS software and qualify vendors
- Protect intellectual property and keep electronic records safe

For more information or to register for this seminar, [please click here](#).

Virtual Training Through WebEx

Date: December 8-9, 2021 (9:00 AM - 3:00 PM PST)

About the Speaker:

Computer System Validation's principal, David Nettleton is an industry leader, author, and teacher for 21 CFR Part 11, Annex 11, HIPAA, software validation, and computer system validation. He is involved with the development, purchase, installation, operation, and maintenance of computerized systems used in FDA compliant applications.

He has completed more than 230 mission-critical laboratory, clinical, and manufacturing software implementation projects. His most popular book is Risk Based Software Validation - Ten easy Steps, which provides fill-in-the-blank templates for completing a COTS software validation project.

About ComplianceOnline.com:

ComplianceOnline is a leading provider of regulatory compliance training programs for companies and professionals in regulated industries. ComplianceOnline has successfully trained

over 55,000 professionals from 15,000 companies to comply with the requirements of regulatory agencies. ComplianceOnline is a MetricStream portal. MetricStream (www.metricstream.com) is a market leader in Enterprise-wide Governance, Risk, Compliance (GRC), and Quality Management Solutions for global corporations.

For more information on ComplianceOnline or to browse through our training programs, please [visit our website](#).

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