



## Compliance4All Webinar

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Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501711	Pre-Control: Easier than SPC	Jd Marhevko	LIVE RECORDED	2018-01-22	\$150.000 \$390.000
501733	Accelerated Aging Techniques for Medical Device Products	Karl J Hemmerich	LIVE RECORDED	2018-01-22	\$150.000 \$390.000
501725	COSO New Fraud Principals	Lynn Fountain	LIVE RECORDED	2018-01-22	\$150.000 \$390.000
501734	Laboratory Investigation of Out-of-Specification Results	Jerry Lanese	LIVE RECORDED	2018-01-23	\$150.000 \$390.000
501737	GLPs: How are they Associated with GMPs and SOPs	Joy McElroy	LIVE RECORDED	2018-01-23	\$150.000 \$390.000
501722	GDPR - European Privacy Regulation what and how	Derk Yntema	LIVE RECORDED	2018-01-24	\$150.000 \$390.000
501742	Why Reputation Monitoring is Key to a Strong Know your Customer (KYC) Program	Jane Hennessy	LIVE RECORDED	2018-01-24	\$150.000 \$390.000
501727	Lean Validation: Cybersecurity and Computer Systems Validation	Valarie King Bailey	LIVE RECORDED	2018-01-24	\$150.000 \$390.000
501740	Root Cause Failure Analysis Closed Loop Corrective Action	Michael Abitz	LIVE RECORDED	2018-01-25	\$150.000 \$390.000
501658	GMP Expectations for Products Used in Early Phase IND Studies	Steven S. Kuwahara	LIVE RECORDED	2018-01-25	\$150.000 \$390.000
501676	Establishing an Effective Pharmaceutical Quality System (Per ICH Q10)	Stephanie Cooke	LIVE RECORDED	2018-01-25	\$150.000 \$390.000
501749	FDA QSR - a Practical Overview of its Implantation and Design Controls	Eyal Lerner	LIVE RECORDED	2018-01-26	\$150.000 \$390.000
501690	Complaint Handling and Management: From Receipt to Trending	David Dills	LIVE RECORDED	2018-01-26	\$150.000 \$390.000
501691	Excel Spreadsheet Validation for FDA 21 CFR Part 11	Angela Bazigos	LIVE RECORDED	2018-01-26	\$150.000 \$390.000
501692	Valid Statistical Rationales for Sample Sizes	John N. Zorich	LIVE RECORDED	2018-01-26	\$150.000 \$390.000
501701	Secrets to Writing Effective SOPs for Medical Device QMS	Susanne Manz	LIVE RECORDED	2018-01-26	\$150.000 \$390.000
501702	Mastering Excel Formulas and Functions	Mike Thomas	LIVE RECORDED	2018-01-26	\$150.000 \$390.000
501704	Water System Mythology: Common False Beliefs for Microbial Control and Monitoring	Teri C. Soli	LIVE RECORDED	2018-01-26	\$150.000 \$390.000
501719	Beta Lactam Facility Remediation: Decontamination for Repurposing the Facility	Paul Lorcheim	LIVE RECORDED	2018-01-29	\$150.000 \$390.000
501672	Evolution of the QMS - How to go from Surviving to Thriving	Susanne Manz	LIVE RECORDED	2018-01-30	\$150.000 \$390.000
501620	Software Validation for the New FDA Inspections	Edwin Waldbusser	LIVE RECORDED	2018-01-30	\$150.000 \$390.000
501685	The Dietary Supplement cGMP Rule (21 CFR part 111)	Thomas E. Colonna	LIVE RECORDED	2018-01-30	\$150.000 \$390.000

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501707	FDA's New Import Program for 2018	Casper Uldriks	LIVE RECORDED	2018-01-30	\$150.000 \$390.000
501708	Validation and Use of Excel® Spreadsheets in FDA Regulated Environments	Dr. Ludwig Huber	LIVE RECORDED	2018-01-30	\$150.000 \$390.000
501755	Supplier Evaluation and Assessment: How to Meet FDA QSR and ISO 13485 Requirements in a Cost-Effective Manner	Jeff Kasoff	LIVE RECORDED	2018-01-31	\$150.000 \$390.000
501651	Design History Files (DHF), Device Master Records (DMR), Device History Records (DHR), Technical Files , Design Dossiers - The Requirements	John E Lincoln	LIVE RECORDED	2018-01-31	\$150.000 \$390.000
501667	Performing Investigations in FDA Regulated Industry	Gary Miller	LIVE RECORDED	2018-01-31	\$150.000 \$390.000
501747	Proper Responses to FDA 483 Observations	Gary Bird	LIVE RECORDED	2018-01-31	\$150.000 \$390.000
501619	21 CFR Part 11 - Compliance for Electronic Records and Signatures	Edwin Waldbusser	LIVE RECORDED	2018-02-02	\$150.000 \$390.000
501694	3-hour Virtual Seminar on Analytical Method Validation Process	Angela Bazigos	LIVE RECORDED	2018-02-02	\$290.000 \$540.000
501625	Validation of FDA-Regulated Computer Systems	Carolyn Troiano	LIVE RECORDED	2018-02-02	\$150.000 \$390.000
501724	Analytical Method Validation Under Good Laboratory Practices - GLPs	John C. Fetzer	LIVE RECORDED	2018-02-05	\$150.000 \$390.000
501669	Master Excel: Introduction to Pivot Tables - Part 1	David Ringstrom	LIVE RECORDED	2018-02-05	\$150.000 \$390.000
501699	FDA's Enforcement of 21 CFR Part 11 Compliance	Angela Bazigos	LIVE RECORDED	2018-02-05	\$150.000 \$390.000
501705	21 CFR Part 11 - Compliance with SaaS/ Cloud Software Applications	David Nettleton	LIVE	2018-02-05	\$150.000
501759	Design History File (DHF), the Device Master Record (DMR) and the Device History Record (DHR) - Principles on Lean Documents and Lean Configuration	Jose Mora	LIVE RECORDED	2018-02-06	\$150.000 \$390.000
501751	Implementing ISO 9001:2015	Lena Cordie	LIVE RECORDED	2018-02-06	\$150.000 \$390.000
501593	How Accurate Adverse Event Reporting is the Key to Subject Safety of Approved Drugs?	Charles H Pierce	LIVE RECORDED	2018-02-06	\$150.000 \$390.000
501718	The RFP Process: How do I Change My Bank? Should I Even do it?	Ray Graber	LIVE RECORDED	2018-02-06	\$150.000 \$390.000
501681	Capturing Justifications in Change Control, Risk Assessment, Validations, and Investigation	Danielle DeLucy	LIVE RECORDED	2018-02-07	\$150.000 \$390.000
501363	Medical Device Hazard Analysis Following ISO 14971	Edwin Waldbusser	LIVE RECORDED	2018-02-07	\$150.000 \$390.000
501663	3-hour Virtual Seminar on How the FDA are Trained for Medical Device Inspections	Angela Bazigos	LIVE RECORDED	2018-02-07	\$290.000 \$540.000

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501712	Calibrating the Human Gage: Attribute Agreement Analysis	Jd Marhevko	LIVE RECORDED	2018-02-08	\$150.000 \$390.000
501677	Good Manufacturing Practices for Active Pharmaceutical Ingredients per ICH Q7	Stephanie Cooke	LIVE RECORDED	2018-02-08	\$150.000 \$390.000
501756	Complaint Handling in Compliance with FDA and ISO Regulations	Jeff Kasoff	LIVE RECORDED	2018-02-09	\$150.000 \$390.000
501739	Time-Saving Excel Tips, Tricks and 100 Shortcuts	Dennis Taylor	LIVE RECORDED	2018-02-09	\$150.000 \$390.000
501767	Combination Products Registration	Salma Michor	LIVE RECORDED	2018-02-09	\$150.000 \$390.000
501728	Automated Validation Testing: Principles & Best Practices	Valarie King Bailey	LIVE RECORDED	2018-02-09	\$150.000 \$390.000
501761	OneNote - Bringing Order to your Digital Chaos	Mike Thomas	LIVE RECORDED	2018-02-12	\$150.000 \$390.000
501764	Understanding and Implementing a Quality by Design Program	Steven Laurenz	LIVE RECORDED	2018-02-12	\$150.000 \$390.000
501689	Risk Management in Clinical Trials	Laura Brown	LIVE RECORDED	2018-02-12	\$150.000 \$390.000
501738	FDA Regulations for Environmental Monitoring(EM) Program	Joy McElroy	LIVE RECORDED	2018-02-13	\$150.000 \$390.000
501741	Process Reliability Modeling: Measure the Cause Not the Effect of Process Variations	Michael Abitz	LIVE RECORDED	2018-02-13	\$150.000 \$390.000
501779	FDA's Policies and Procedures for Food Recall	Michele Pfannenstiel	LIVE RECORDED	2018-02-13	\$150.000 \$390.000
501726	Understanding the Requirements of SSAE18	Lynn Fountain	LIVE RECORDED	2018-02-13	\$150.000 \$390.000
501781	Project Management Best Practices for Computer Systems Regulated by FDA	Carolyn Troiano	LIVE RECORDED	2018-02-14	\$150.000 \$390.000
501743	Understanding International ACH	Jane Hennessy	LIVE RECORDED	2018-02-14	\$150.000 \$390.000
501780	3 Hour Boot Camp Webinar on FDA's Food Facility Inspection	Angela Bazigos	LIVE RECORDED	2018-02-14	\$290.000 \$540.000
501678	Unique Device Identification (UDI) - GUDID: Final Rules, Implementation and Compliance	John E Lincoln	LIVE RECORDED	2018-02-14	\$150.000 \$390.000
501735	cGMPs in the Quality Control Laboratory	Jerry Lanese	LIVE RECORDED	2018-02-15	\$150.000 \$390.000
501776	2011 FDA Guideline on Process Validation	Louis Angelucci	LIVE RECORDED	2018-02-15	\$150.000 \$390.000
501688	The New ISO 13485:2016 and Comparison with 21CFR820 - How to Comply with both in the Same Organization	Eyal Lerner	LIVE RECORDED	2018-02-16	\$150.000 \$390.000
501748	Data Integrity: Finding and Resolving the Issues	Gary Bird	LIVE RECORDED	2018-02-16	\$150.000 \$390.000
501757	FDA Inspections - Do's and Don'ts	Jonathan M. Lewis	LIVE RECORDED	2018-02-21	\$150.000 \$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501763	Get Organised with Microsoft Outlook	Mike Thomas	LIVE RECORDED	2018-02-21	\$150.000 \$390.000
501673	Management Control - Underutilized and Underappreciated	Susanne Manz	LIVE RECORDED	2018-02-21	\$150.000 \$390.000
501731	Statistical Concepts of Process Validation	Dan OLeary	LIVE RECORDED	2018-02-21	\$150.000 \$390.000
501687	Medical Device Complaints & CAPA	Edwin Waldbusser	LIVE RECORDED	2018-02-21	\$150.000 \$390.000
501784	GCP/GLP/GMP : Comparison and Understanding of FDA's 3 Major Regulations	Dr. David Lim	LIVE RECORDED	2018-02-22	\$150.000 \$390.000
501787	Understanding the USP Chapter 1224 for Transfer of Analytical Methods	Dr. Ludwig Huber	LIVE RECORDED	2018-02-22	\$150.000 \$390.000
501768	Biosimilars from Development to Registration	Salma Michor	LIVE RECORDED	2018-02-23	\$150.000 \$390.000
501771	Excel Spreadsheets - Step-By-Step Instructions for Compliance	David Nettleton	LIVE RECORDED	2018-02-23	\$150.000 \$390.000
501720	Developing a Remediation Plan for MMV in a Pharmaceutical Biologic Facility	Paul Lorcheim	LIVE RECORDED	2018-02-23	\$150.000 \$390.000
501744	Managing Higher-Risk Business Customers	Jane Hennessy	LIVE RECORDED	2018-02-26	\$150.000 \$390.000
501785	Strategies to Enable SME Device Companies to Cope with the Challenges of the Upcoming European Medical Device Regulation (MDR)	Stefan Menzl	LIVE RECORDED	2018-02-26	\$150.000 \$390.000
501786	FSMA Compliance for Importing and Transporting Food	Joe Lynch	LIVE RECORDED	2018-02-27	\$150.000 \$390.000
501729	Lean Validation: Validating Microsoft Dynamics 365 For Operations & Azure: A Practical Approach	Valarie King Bailey	LIVE RECORDED	2018-02-27	\$150.000 \$390.000
501697	Controlling Human Error in the Manufacturing Floor	Ginette Collazo	LIVE RECORDED	2018-02-27	\$150.000 \$390.000
501754	Implementing an Effective CAPA System	Charles H. Paul	LIVE RECORDED	2018-02-28	\$150.000 \$390.000
501770	Tactical Problem Solving: Maximize Organizational Effectiveness, Efficiency, Reliability and Profit	Michael Abitz	LIVE RECORDED	2018-02-28	\$150.000 \$390.000
501682	Bioavailability and Bioequivalence Studies Submitted in NDAs and INDs for Orally-Administered Drug Products	Stephanie Cooke	LIVE RECORDED	2018-02-28	\$150.000 \$390.000
501679	Ensuring Data Integrity in Method Validations	Gary Miller	LIVE RECORDED	2018-02-28	\$150.000 \$390.000
501773	Bullet Proof 510(k) - Latest FDA Changes to the Process	David Dills	LIVE RECORDED	2018-02-28	\$150.000 \$390.000
501777	Similarities and Differences Between an FDA and MHRA Audit	Louis Angelucci	LIVE RECORDED	2018-03-01	\$150.000 \$390.000
501765	Understanding and Implementing a Technology Transfer Process	Steven Laurenz	LIVE RECORDED	2018-03-05	\$150.000 \$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501594	FDA Clinical Trial Auditing and the Due Diligence Companies Should Conduct as Part of their Monitoring Program	Charles H Pierce	LIVE RECORDED	2018-03-06	\$150.000 \$390.000
501782	Good Documentation Practices to Support FDA Computer System Validation	Carolyn Troiano	LIVE RECORDED	2018-03-07	\$150.000 \$390.000
501760	Risk Management Utilizing Lean Documents and Lean Configuration	Jose Mora	LIVE RECORDED	2018-03-07	\$150.000 \$390.000
501788	FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	LIVE RECORDED	2018-03-07	\$150.000 \$390.000
501703	FDA's 21 CFR 11 Add-On Inspections - Recent Updates	Angela Bazigos	LIVE RECORDED	2018-03-07	\$150.000 \$390.000
501674	Non-Conforming Material and Failure Investigation	Susanne Manz	LIVE RECORDED	2018-03-08	\$150.000 \$390.000
501713	Predictive Warranty Using Paynter Charts	Jd Marhevko	LIVE RECORDED	2018-03-08	\$150.000 \$390.000
501736	Analytical Method Validation	Jerry Lanese	LIVE RECORDED	2018-03-09	\$150.000 \$390.000
501769	The New EU Medical Device Regulation	Salma Michor	LIVE RECORDED	2018-03-12	\$150.000 \$390.000
501752	Software Verification & Validation	Lena Cordie	LIVE RECORDED	2018-03-13	\$150.000 \$390.000
501698	Tools for Human Error Reduction	Ginette Collazo	LIVE RECORDED	2018-03-13	\$150.000 \$390.000
501721	Eliminating Microbial Contamination in a Spiral Freezer	Paul Lorcheim	LIVE RECORDED	2018-03-13	\$150.000 \$390.000
501684	Dietary Supplements CGMPs - 21 CFR 111 Compliance	John E Lincoln	LIVE RECORDED	2018-03-14	\$150.000 \$390.000
501709	Key Factors to Write an Effective Standard Operating Procedure (SOP) and Work Instructions (WIs)	Angela Bazigos	LIVE RECORDED	2018-03-14	\$150.000 \$390.000
501778	Understanding Proper Application of Risk Assessment	Louis Angelucci	LIVE RECORDED	2018-03-15	\$150.000 \$390.000
501758	Building a Sustainable Vendor Qualification Program	Jonathan M. Lewis	LIVE RECORDED	2018-03-16	\$150.000 \$390.000
501732	Excel Workbooks and FDA Device Regulations	Dan OLeary	LIVE RECORDED	2018-03-21	\$150.000 \$390.000
501789	Understanding and Implementing USP 1058 Analytical Instrument Qualification	Dr. Ludwig Huber	LIVE RECORDED	2018-03-22	\$150.000 \$390.000
501683	Good Manufacturing Practices for Phase I Investigational Drug Products: GMPs Required for Drug Products Used in Phase I Clinical Trials	Stephanie Cooke	LIVE RECORDED	2018-03-22	\$150.000 \$390.000
501774	Conducting Successful Product Complaint Investigations	David Dills	LIVE RECORDED	2018-03-22	\$150.000 \$390.000
501775	Importing and Exporting Medical Devices: A Primer on Regulatory Strategy and Requirements	David Dills	LIVE RECORDED	2018-03-27	\$150.000 \$390.000

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501766	Pharmaceutical Quality Risk Management(QRM) Overview	Steven Laurenz	LIVE RECORDED	2018-03-28	\$150.000 \$390.000
501668	European Data Protection Regulation - 2018 Implementation	Angela Bazigos	LIVE RECORDED	2018-03-29	\$150.000 \$390.000
501783	Data Governance for Computer Systems Regulated by FDA	Carolyn Troiano	LIVE RECORDED	2018-03-30	\$150.000 \$390.000
501714	Understanding Autism Spectrum Disorder	Jd Marhevko	LIVE RECORDED	2018-03-30	\$150.000 \$390.000
501653	Key Factors to Write an Effective Standard Operating Procedure (SOP) and Work Instructions (WIs)	Angela Bazigos	LIVE RECORDED	2018-04-02	\$150.000 \$390.000
501772	21 CFR Part 11 - Compliance for Electronic Records and Signatures	David Nettleton	LIVE RECORDED	2018-04-06	\$150.000 \$390.000
501753	Differences Between ISO 13485 & ISO 9001 - and How to Choose Which One to Implement	Lena Cordie	LIVE RECORDED	2018-04-10	\$150.000 \$390.000
501715	Driving a Quality-Lean Steering Committee	Jd Marhevko	LIVE RECORDED	2018-04-20	\$150.000 \$390.000
501686	3-Hour Virtual Seminar on Off label Promotion of Drugs and Medical Devices - FDA's Latest	Angela Bazigos	LIVE RECORDED	2018-04-24	\$290.000 \$540.000
501746	How to Comply with 21 CFR 11 Requirements for Electronic Medical Records	Angela Bazigos	LIVE RECORDED	2018-05-02	\$150.000 \$390.000
501762	Working Effectively with Customers & Suppliers	Charles H. Paul	LIVE RECORDED	2018-07-26	\$150.000 \$390.000
500001	Introduction to Sales and Use Taxation	Miles Hutchinson	RECORDED	2014-11-12	\$149.000
500002	Hot Issues in Determining Multi-State Sales Tax Nexus	Miles Hutchinson	RECORDED	2014-11-25	\$149.000
500004	Complying with FATCA - the Foreign Accounts Tax Compliance Act	Miles Hutchinson	RECORDED	2014-05-22	\$149.000
500007	Compliance and Continual Improvement, Tools of the Trade	Rodriguez Gonzalez	RECORDED	2014-04-16	\$149.000
500010	Foodservice Sanitation	Norman G. Marriott	RECORDED	2014-07-15	\$149.000
500012	Foreign Corrupt Practices Act - Audits Role	Denise Cicchella	RECORDED	2014-05-27	\$149.000
500013	Operational Risk - Beyond Regulatory Constraints	Fred Vacelet	RECORDED	2014-07-02	\$149.000
500015	System Safety Engineering	Dev Raheja	RECORDED	2014-05-20	\$149.000
500016	Fundamental ATC Organization, Structure, and Concepts	Stephen Alvania	RECORDED	2014-05-20	\$149.000
500017	ATC Radar Approach Control (TRACON)	Stephen Alvania	RECORDED	2014-06-10	\$149.000
500018	Mastering the Power of Sensitivity Tools in Financial Modeling	Miles Hutchinson	RECORDED	2014-08-28	\$149.000

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500020	Air Traffic Control and Airport Operation Coordination	Roger Nakata	RECORDED	2014-05-28	\$149.000
500022	Project Management - Auditing Problem Projects	Stuart Gardner	RECORDED	2014-06-19	\$149.000
500023	Understanding Next Gen (FAA New Technology)	Roger Nakata	RECORDED	2014-06-19	\$149.000
500024	Industrial Technical Benefits- Canada's New Approach to Offsets for Defence Procurement	Anthony Goode	RECORDED	2014-06-03	\$149.000
500025	Reducing Variation in Manufacturing Processes	Lynne Hare	RECORDED	2014-06-05	\$149.000
500026	Sanitation Chemical Testing - Best Practices	Donald Jones	RECORDED	2014-08-12	\$149.000
500027	Nonresident Alien W-8 & 1042-S Compliance Update	Miles Hutchinson	RECORDED	2014-06-18	\$149.000
500028	TIN Matching to Reduce Your B-Notices and Eliminate Proposed Penalties	Miles Hutchinson	RECORDED	2014-07-08	\$149.000
500029	Environmental Cleaning	Donald Jones	RECORDED	2014-07-09	\$149.000
500030	Basics of Compliance for Food Businesses	Rodriguez Gonzalez	RECORDED	2014-07-01	\$149.000
500031	Metrics and Compliance	Rodriguez Gonzalez	RECORDED	2014-07-31	\$149.000
500032	Planning for Resilience - Best Practices for Developing Reliable Disaster Recovery Plans	Ms. Michael Redmond	RECORDED	2014-07-15	\$149.000
500034	Airport Traffic Control Tower (ATCT)	Stephen Alvania	RECORDED	2014-07-09	\$149.000
500035	Air Route Traffic Control Center (ARTCC) Operations	Stephen Alvania	RECORDED	2014-07-30	\$149.000
500036	Traffic Flow Management (TFM) Operations-End to End Flight Scenario	Stephen Alvania	RECORDED	2014-08-20	\$149.000
500037	Accounting and Tax Procedures to Maximize your Deductions and Minimize Problems with The IRS	Reuven Rubinson	RECORDED	2014-08-26	\$149.000
500038	The National Shipbuilding Procurement Strategy- Changing the playing field for shipbuilding in Canada	Anthony Goode	RECORDED	2014-07-17	\$149.000
500039	The Business of Defence Procurement in Canada- A Market Over	Anthony Goode	RECORDED	2014-08-07	\$149.000
500040	ISO Standard for Business Continuity (ISO 22301)	Ms. Michael Redmond	RECORDED	2014-09-16	\$149.000
500041	Using Forensic Accountants Effectively in the Legal Process	Ray Kulzick	RECORDED	2014-08-07	\$149.000
500042	Detecting Frauds in Purchasing and Payables	Ray Kulzick	RECORDED	2014-09-09	\$149.000
500046	1099 and W-9 Update - Complying with IRS Information Reporting	Miles Hutchinson	RECORDED	2014-10-02	\$149.000
500051	Controlled Goods in Canada- A primer	Anthony Goode	RECORDED	2014-08-27	\$149.000



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500053	How to Write and Implement Effective Policies & Procedures	Miles Hutchinson	RECORDED	2014-10-21	\$149.000
500054	How to Design and Implement Outstanding KPI Performance Dash	Miles Hutchinson	RECORDED	2015-01-27	\$149.000
500056	One practical view of HACCP Plans	Rodriguez Gonzalez	RECORDED	2014-10-21	\$149.000
500058	Airport Traffic Control Tower (ATCT)	Stephen Alvania	RECORDED	2014-10-29	\$149.000
500060	Air Route Traffic Control Center (ARTCC) Operations	Stephen Alvania	RECORDED	2014-11-25	\$149.000
500061	Traffic Flow Management (TFM) Operations-End to End Flight Scenario	Stephen Alvania	RECORDED	2014-12-11	\$149.000
500064	What's Up? (and Down): Outlook for Construction Segments, Materials and Labor	Kenneth D. Simonson	RECORDED	2014-09-30	\$149.000
500066	Construction Shortages Ahead: Skilled Labor, Skilled Manager	Kenneth D. Simonson	RECORDED	2014-10-30	\$149.000
500067	How to Analyze Financial Statements	Miles Hutchinson	RECORDED	2014-12-10	\$149.000
500069	The Value of Hourly Cost Maintenance Programs	Anthony Kiousis	RECORDED	2014-10-29	\$149.000
500071	Canada's Defence Budget Crisis- Impact of recent Defense Budget Cuts on The Canadian Armed Forces- Opportunities for Industry	Anthony Goode	RECORDED	2014-11-19	\$149.000
500074	Contract Killer Clauses in Construction: And How to Neutralize Them	Brian Perlberg	RECORDED	2015-01-15	\$149.000
500075	The Master Validation Plan - The Unwritten Requirements	John E Lincoln	RECORDED	2014-11-04	\$189.000
500076	Quality by Design: Establishing a Systematic Approach to Pharmaceutical Development	Heath Rushing	RECORDED	2014-11-17	\$189.000
500077	21 CFR Part 11 - Compliance for Electronic Records and Signatures	David Nettleton		2014-11-04	
500079	FDA's Proposed Rule Regarding Device Establishment Registration and Listing and How to Register and List	David Dills	RECORDED	2014-11-24	\$189.000
500080	FDA Compliant HPLC Qualification and Performance Testing	Dr. Ludwig Huber	RECORDED	2014-11-06	\$189.000
500081	GMP Perspectives on Working with Contracting Laboratories	Steven S. Kuwahara	RECORDED	2014-11-11	\$189.000
500082	Cleaning and Sanitation Training for Food Processors	Melinda Allen	RECORDED	2014-11-13	\$189.000
500083	Auditing Analytical Laboratories for FDA Compliance	Steven S. Kuwahara	RECORDED	2014-11-18	\$189.000
500084	Compiling the Design History File, and Technical File, Design Dossier	John E Lincoln	RECORDED	2014-11-18	\$189.000
500086	FDA's 2011 Draft Guidance on Financial Disclosure by Clinical Investigators	David Dills	RECORDED	2014-11-19	\$189.000

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500088	The European Clinical Trial Directive (EUCTD)	Robert J. Russell	RECORDED	2014-12-16	\$189.000
500089	Quality by Design (QbD) for Development and Validation of Analytical Methods	Dr. Ludwig Huber	RECORDED	2014-11-20	\$189.000
500091	FDA Regulation of In Vitro Diagnostics	Thomas E. Colonna	RECORDED	2014-11-20	\$189.000
500092	Statistical Concepts of Process Validation	Dan OLeary	RECORDED	2015-01-22	\$189.000
500093	Change Control - Key to Successful cGMP Compliance	John E Lincoln	RECORDED	2014-12-02	\$189.000
500095	How to Manage a Medical Device Recall Efficiently and Effectively	David Dills	RECORDED	2014-12-03	\$189.000
500097	The FDA Drug Development Process: GLP, GMP and GCP Regulations	Albert A. Ghignone	RECORDED	2014-11-06	\$189.000
500102	Understanding Combination Products, Requests for Designation and Product Jurisdiction	Elisa Harvey	RECORDED	2014-11-05	\$189.000
500103	Using the Pre-Submission Process to Your Best Advantage	Elisa Harvey	RECORDED	2014-12-03	\$189.000
500105	Foreign Bodies in Foods - Effective Techniques for Prevention, Control and Detection	Rotimi Toki	RECORDED	2014-11-21	\$149.000
500106	Food Labelling: A Practical Guide for Unambiguous Labelling Information & Compliance with Regulatory Requirements	Rotimi Toki	RECORDED	2014-12-12	\$149.000
500108	EU Pharmacovigilance Directive and Regulations	Robert J. Russell	RECORDED	2014-12-10	\$189.000
500110	Developing an Efficient Relationship with FDA	Robert Kunka	RECORDED	2014-12-16	\$189.000
500111	Device Changes, FDA Changes, and the 510(k)	John E Lincoln	RECORDED	2014-12-16	\$189.000
500112	Maintaining an Effective CAPA Program and Using Risk Assessment Tools: Current Trends	David Dills	RECORDED	2014-12-17	\$189.000
500113	Device Corrections and Removals	Dan OLeary	RECORDED	2014-12-18	\$189.000
500114	GMP Expectations for Products Used in Early Phase IND Studies	Steven S. Kuwahara	RECORDED	2015-01-06	\$189.000
500115	IT Auditing - Principles and Practices (2nd Edition)	Robert E. Davis	RECORDED	2014-11-20	\$149.000
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500357	Using the Pre-Submission Process to Your Best Advantage	Elisa Harvey	RECORDED	2015-06-08	\$131.000
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500398	Calibration and Qualification in Analytical Laboratories	Dr. Ludwig Huber	RECORDED	2015-07-08	\$183.000
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500428	Medical Device Product Development Process	Karl Leinsing	RECORDED	2015-06-24	\$147.000
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500430	Stress-Testing for Financial Institutions, Beyond Regulatory Constraints	Fred Vacelet	RECORDED	2015-07-09	\$140.000
500432	Key Factors to Write an Effective Standard Operating Procedure (SOP) and Work Instructions (WIs)	Angela Bazigos	RECORDED	2015-08-04	\$163.000

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500440	Medical Device Recalls	Larry Spears	RECORDED	2015-08-26	\$148.000
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500443	Meeting Annual U.S. FDA cGMP Training Requirements	John E Lincoln	RECORDED	2015-08-26	\$162.000
500444	Device Changes, FDA Changes, and the 510(k)	John E Lincoln	RECORDED	2015-09-09	\$141.000
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500464	Capturing the Most Powerful Market - Moms	Shari Storm	RECORDED	2015-08-31	\$132.000
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500470	FDA Inspections – Do's and Don'ts	Jonathan M. Lewis	RECORDED	2015-09-25	\$161.000
500471	Key Steps to Successful Project Management Principles	Carol Friedhoff	RECORDED	2015-10-01	\$132.000
500472	What Contractors Need to Know About AIA A201 and Consensus Docs Contracts	Brian Perlberg	RECORDED	2015-08-12	\$146.000

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500475	Spreadsheet-Based Internal Controls	David Ringstrom	RECORDED	2015-09-17	\$153.000
500477	Sample Size and Statistical Rationale for Medical Device Packaging Validations	Karen Greene	RECORDED	2015-08-20	\$148.000
500478	Similarities and Differences Between an FDA and MHRA Audit	Louis Angelucci	RECORDED	2015-07-28	\$153.000
500479	Validation Master Planning and Regulatory Expectations	Louis Angelucci	RECORDED	2015-08-12	\$157.000
500480	Introduction to Risk Assessment	Louis Angelucci	RECORDED	2015-09-15	\$145.000
500481	Building Quality Systems for Pharmaceutical and Medical Device Firms	Louis Angelucci	RECORDED	2015-10-27	\$190.000
500487	Building a Successful Relationship with the US FDA: Key Steps	Robert Kunka	RECORDED	2015-08-19	\$127.000
500489	Introduction to Payroll Law	Miles Hutchinson	RECORDED	2016-01-04	\$190.000
500490	The Bayh-Dole Act and Its Implementing Regulations & Rule-Making	Ronald Kudla	RECORDED	2015-08-04	\$143.000
500492	Risk Management for Commissioning and Qualification for Pharmaceutical Industry	Majdi Ayoub	RECORDED	2015-09-24	\$151.000
500496	Quality Agreements and Annual Inspections	Jerry Dalfors	RECORDED	2015-09-14	\$158.000
500498	Design Considerations For Air Barriers	Leonard Anastasi	RECORDED	2015-08-14	\$148.000
500499	Avoiding Pitfalls With Air Barriers	Leonard Anastasi	RECORDED	2015-09-17	\$154.000
500501	Strategic Management of Corporate Governance: Setting the Right Tone at the Top about Risk, Part 1	Ron Rael	RECORDED	2015-08-24	\$128.000
500502	Ethical Compliance Starts with Accountability	Ron Rael	RECORDED	2015-09-23	\$132.000
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500511	Operational Risk for Financial Institutions - Beyond Regulatory Constraints	Fred Vacelet	RECORDED	2015-09-01	\$143.000
500517	Contract Killer Clauses in Construction: And How to Neutralize Them	Brian Perlberg	RECORDED	2015-09-24	\$148.000
500519	Menu Labelling: Knowing The Requirements and Available Options for Achieving Compliance	Rotimi Toki	RECORDED	2015-10-27	\$190.000
500520	Managing Client Expectations Without Losing Your Shirt or the Next Job	Heath Suddleson	RECORDED	2015-09-03	\$146.000
500521	From Project Manager to Project Leader	Heath Suddleson	RECORDED	2015-09-22	\$147.000
500534	21st Century Validation	Sam DeMarco	RECORDED	2015-10-30	\$190.000
500546	The New Clinical Trial Regulation	Adriaan Fruijtjer	RECORDED	2015-09-21	\$146.000
500554	Driving a Quality-Lean Steering Committee	Jd Marhevko	RECORDED	2015-11-24	\$152.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
500558	Pediatric Drug Development: Regulatory Expectations with New Laws	Robert Kunka	RECORDED	2015-11-09	\$190.000
500560	Good Documentation Practices for GxP Compliance	Alla Teresh	RECORDED	2015-10-14	\$190.000
500561	Analytical Instrument Qualification According The New Revision of USP 1058	Dr. Ludwig Huber	RECORDED	2015-11-05	\$190.000
500562	FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	RECORDED	2015-12-10	\$190.000
500563	FDA's Bioresearch Monitoring (BIMO) Program Biologic, Device, Animal Drug, and Food Areas	Thomas E. Colonna	RECORDED	2015-10-29	\$190.000
500564	FDA Regulation of Combination Products	Thomas E. Colonna	RECORDED	2015-11-19	\$190.000
500566	Cyber Security Risk Assessment	Ms. Michael Redmond	RECORDED	2015-12-01	\$190.000
500568	Cyber Security Incident Response Team Training Program	Ms. Michael Redmond	RECORDED	2015-12-09	\$190.000
500570	Verification or Validation of Methods in Food Microbiology	Michael Brodsky	RECORDED	2015-11-09	\$190.000
500573	Fundamentals of Lyophilization Technology	Jerry Dalfors	RECORDED	2015-11-10	\$190.000
500575	In Other Words- The Clinical and Construction Guide to Building Patient Centered Environments	Cathy Dolan-Schweitzer	RECORDED	2015-10-27	\$190.000
500576	Development and Implementation of an Internal Auditing Program as part of a Pharmaceutical Manufacturing Quality System	Joseph Habarta	RECORDED	2015-10-29	\$190.000
500578	Validating Radiation Sterilization for Medical Products	Karl J Hemmerich	RECORDED	2015-11-12	\$190.000
500579	Accelerated Aging Techniques for Medical Device Packaging	Karl J Hemmerich	RECORDED	2015-12-03	\$190.000
500580	Social Media Marketing & FDA Regulations	Casper Uldriks	RECORDED	2015-11-10	\$190.000
500581	Best Practices for Handling FDA Inspections	Casper Uldriks	RECORDED	2015-12-15	\$190.000
500582	The Bayh-Dole Act and Its Implementing Regulations & Rule-Making	Ronald Kudla	RECORDED	2015-11-03	\$190.000
500585	FSMA Impact to the Transportation of Perishables	John Ryan	RECORDED	2015-11-16	\$190.000
500586	USFDA Food Import and Export: Current and Future Safety Regulations	John Ryan	RECORDED	2015-12-07	\$190.000
500587	Operational Risk - Beyond Regulatory Constraints	Fred Vacelet	RECORDED	2015-11-09	\$190.000
500589	Root Cause Analysis: KEY to an Effective Corrective Actions System	Betty Lane	RECORDED	2015-12-07	\$190.000
500592	Electronic Informed Consent for Clinical Trials: Why, What and How	Susan Brink	RECORDED	2015-11-17	\$190.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
500598	Dealing with Performance Issues	Heath Suddleson	RECORDED	2015-11-30	\$190.000
500599	PMBA - Project Management Business Administration	Heath Suddleson	RECORDED	2015-12-15	\$190.000
500600	IFRS 6 for Oil, Gas and Mining Industries	Mike Morley	RECORDED	2016-01-20	\$190.000
500601	Does Anyone Care About SOX Anymore?	Mike Morley	RECORDED	2016-02-08	\$190.000
500602	SOX: Internal Controls for Accounts Payable	Mike Morley	RECORDED	2016-01-29	\$190.000
500603	Corporate Expense Accounts Review and Audit Best Practices	Mike Morley	RECORDED	2016-02-11	\$190.000
500604	B2B Payments in the US and Compliance Issues	Ray Graber	RECORDED	2015-12-09	\$190.000
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500607	Safety and Motivation for EHS practitioners	James Thatcher	RECORDED	2015-12-10	\$190.000
500610	SOPs for Clinical Trials: Regulatory Requirements	Harold Thibodeaux	RECORDED	2016-02-11	\$190.000
500612	Sterile Medical Packaging Design - 7 Essentials	Karen Greene	RECORDED	2016-01-14	\$190.000
500613	Sample Size and Statistical Rationale for Medical Device Packaging Validations	Karen Greene	RECORDED	2016-02-04	\$190.000
500615	Menu Labelling: Knowing The Requirements and Available Options for Achieving Compliance	Rotimi Toki	RECORDED	2016-01-20	\$190.000
500620	Combined HACCP Development & Risk-Based HACCP	Rotimi Toki	RECORDED	2016-06-22	\$390.000
500625	Understanding Autism Spectrum Disorder	Jd Marhevko	RECORDED	2016-01-08	\$190.000
500627	Post-approval changes for medicinal products in the EU	Adriaan Fruijtjer	RECORDED	2016-01-25	\$190.000
500628	Orphan Medicinal Product Designation in the EU	Adriaan Fruijtjer	RECORDED	2016-02-05	\$190.000
500630	Scientific Advice in the EU	Adriaan Fruijtjer	RECORDED	2016-03-04	\$190.000
500631	Pediatric Investigation Plans (PIP) in the EU	Adriaan Fruijtjer	RECORDED	2016-03-16	\$190.000
500633	Sanitizers for the Food Industry	Norman G. Marriott	RECORDED	2016-03-15	\$190.000
500634	The Marriage of the PFMEA and Control Plan, A Dynamic Control Plan	Jd Marhevko	RECORDED	2016-02-05	\$190.000
500635	Predictive Warranty using Paynter Charts	Jd Marhevko	RECORDED	2016-02-22	\$190.000
500636	Calibrating the Human Gage: Attribute Agreement Analysis	Jd Marhevko	RECORDED	2016-03-22	\$190.000
500638	Pre-Control: Easier than SPC	Jd Marhevko	RECORDED	2016-04-29	\$390.000

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500641	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	RECORDED	2016-01-27	\$190.000
500642	Introduction to SPC (Statistical Process Control)	John N. Zorich	RECORDED	2016-02-10	\$190.000
500643	Normality Tests and Normality Transformations: Explanations, Justifications, and Uses	John N. Zorich	RECORDED	2016-02-17	\$190.000
500644	Metrology: Statistical Analysis of Measurement Uncertainty	John N. Zorich	RECORDED	2016-03-02	\$190.000
500649	Construction - OSHA Construction Basics, for any Jobsite	Jason Teliszczak	RECORDED	2016-02-26	\$190.000
500657	Traceability and Recall Through Food Processes	John Ryan	RECORDED	2016-02-08	\$190.000
500659	Standardizing Transportation Procedures to Control Food Safety and Quality	John Ryan	RECORDED	2016-03-02	\$190.000
500660	Prepare Your Company to Meet the Final FSMA Subpart G Rule Requirements for the Receiver Liability for Supplier Preventive Controls	John Ryan	RECORDED	2016-03-22	\$190.000
500661	Validating Radiation Sterilization for Medical Products	Karl J Hemmerich	RECORDED	2016-04-07	\$190.000
500662	Accelerated Aging Techniques for Medical Device Packaging	Karl J Hemmerich	RECORDED	2016-01-29	\$190.000
500669	Outlook for Multifamily Housing	Bernard Markstein	RECORDED	2016-02-23	\$190.000
500670	Reclaiming Large Excel Spreadsheets	David Ringstrom	RECORDED	2016-01-15	\$190.000
500671	Spreadsheet-Based Internal Controls	David Ringstrom	RECORDED	2016-02-04	\$190.000
500672	Expedite Excel with Hidden Shortcuts	David Ringstrom	RECORDED	2016-03-24	\$190.000
500674	Transcend VLOOKUP in Microsoft Excel	David Ringstrom	RECORDED	2016-04-21	\$390.000
500677	Key Steps to Successful Project Management Principles	Carol Friedhoff	RECORDED	2016-02-18	\$190.000
500679	The Role of FDA in Health Care Software Regulations and Development	Anna Longwell	RECORDED	2016-01-19	\$190.000
500680	A second look at 510(k) changes	Anna Longwell	RECORDED	2016-02-11	\$190.000
500681	FDA's Medical Device Clinical Trials Program	Anna Longwell	RECORDED	2016-02-22	\$190.000
500682	Laboratory-Developed Tests: Why does FDA think They Can Regulate Them, and Why Do Others Think They Cannot	Anna Longwell	RECORDED	2016-03-15	\$190.000
500683	Clinical Trial Applications in Canada, and Comparison to the US and Europe	Anne Tomalin	RECORDED	2016-02-02	\$190.000
500684	Current Status of biosimilars in US, Europe and Canada	Anne Tomalin	RECORDED	2016-03-03	\$190.000
500687	Food Safety, Security and Fraud-Are You Ready	Michael Brodsky	RECORDED	2016-02-01	\$190.000

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500688	Quality Control for Analytical Materials used in Microbiology Laboratories	Michael Brodsky	RECORDED	2016-02-25	\$190.000
500690	Understanding the New USP Chapter 1224 for Transfer of Analytical Methods	Dr. Ludwig Huber	RECORDED	2016-02-03	\$190.000
500691	Validation and Use of Excel® Spreadsheets in FDA Regulated Environments	Dr. Ludwig Huber	RECORDED	2016-03-02	\$390.000
500692	Effective Training Practices for FDA Compliance	Dr. Ludwig Huber	RECORDED	2016-04-13	\$390.000
500693	Managing Client Expectations Without Losing Your Shirt or the Next Job	Heath Suddleson	RECORDED	2016-01-19	\$190.000
500694	From Project Manager to Project Leader	Heath Suddleson	RECORDED	2016-02-11	\$190.000
500695	Dealing with Performance Issues	Heath Suddleson	RECORDED	2016-02-29	\$190.000
500697	FDA Regulation of Dietary Supplements	Thomas E. Colonna	RECORDED	2016-02-25	\$190.000
500698	Introduction to FDA Regulation of Combination Products	Thomas E. Colonna	RECORDED	2016-03-31	\$190.000
500704	Establishment of Quality Systems	Louis Angelucci	RECORDED	2016-05-05	\$390.000
500705	Introduction to Risk Assessment	Louis Angelucci	RECORDED	2016-02-18	\$190.000
500706	Validation Master Planning and Regulatory Expectations	Louis Angelucci	RECORDED	2016-03-03	\$190.000
500707	2011 FDA Guideline on Process Validation	Louis Angelucci	RECORDED	2016-03-24	\$190.000
500708	Similarities and Differences Between an FDA and MHRA Audit	Louis Angelucci	RECORDED	2016-04-21	\$390.000
500731	Dietary Supplement Regulatory Compliance in the United States: Labeling, Product Claims & Updates from the FDA	James E. Russell	RECORDED	2016-03-29	\$190.000
500732	Japan - Regulatory Filing Requirements and Compliance Processes for Life Sciences	Robert J. Russell	RECORDED	2016-03-22	\$390.000
500737	Drug Master Files: Understanding and Meeting your Regulatory and Processing Responsibilities	Robert J. Russell	RECORDED	2016-04-20	\$390.000
500740	Medical Device Product Development Process	Karl Leinsing	RECORDED	2016-03-10	\$190.000
500745	FDA Regulations for Analytical Instrument Qualification and Validation Processes	Joy McElroy	RECORDED	2016-03-29	\$190.000
500746	Best Practices for an Effective Cleaning Validation Procedures	Joy McElroy	RECORDED	2016-04-14	\$190.000
500747	FDA Regulations for Environmental Monitoring(EM) Program	Joy McElroy	RECORDED	2016-04-28	\$390.000
500749	The New European Clinical Trial Regulation	Robert J. Russell	RECORDED	2016-04-06	\$190.000



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500750	Latin America Regulatory Compliance: Current Challenges in Argentina, Brazil and Mexico	Robert J. Russell	RECORDED	2016-09-26	\$390.000
500755	21 CFR Part 820 - Quality System Regulation - Applying Principles of Lean Documents and Lean Configuration	Jose Mora	RECORDED	2016-04-19	\$390.000
500758	Design History File (DHF), the Device Master Record (DMR) and the Device History Record (DHR) – Principles on Lean Documents and Lean Configuration	Jose Mora	RECORDED	2016-06-09	\$390.000
500759	Research Use Only Products - The Dos and Don'ts	Harold Thibodeaux	RECORDED	2016-03-15	\$190.000
500760	FDA New Drug Approval Process	Harold Thibodeaux	RECORDED	2016-04-12	\$190.000
500761	FDA - Legal Writing Skills that Result in Effective Regulatory Submissions	Robert Michalik	RECORDED	2016-04-06	\$190.000
500766	Understanding, Calculating, and Using Statistical Power	John N. Zorich	RECORDED	2016-04-05	\$190.000
500767	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	RECORDED	2016-04-20	\$390.000
500768	Process Capability Analysis by means of Confidence - Reliability Calculations	John N. Zorich	RECORDED	2016-05-03	\$390.000
500769	Process Capability Analysis of Extremely Non-Normal Data	John N. Zorich	RECORDED	2016-05-19	\$390.000
500770	Introduction to SPC (Statistical Process Control)	John N. Zorich	RECORDED	2016-06-16	\$390.000
500778	Modeling and Optimizing Process Behavior using Design of Experiments	Steven Wachs	RECORDED	2016-05-09	\$390.000
500779	Excel Spreadsheet Validation for FDA 21 CFR Part 11	Angela Bazigos	RECORDED	2016-04-14	\$390.000
500780	Key Factors to Write an Effective Standard Operating Procedure (SOP) and Work Instructions (WIs)	Angela Bazigos	RECORDED	2016-05-12	\$390.000
500781	FDA's 21 CFR 11 Add-On Inspections - Recent Updates	Angela Bazigos	RECORDED	2016-06-02	\$390.000
500785	Using Statistics to Determine Sample Size	Steven Walfish	RECORDED	2016-04-07	\$190.000
500787	Sample Size and Statistical Rationale for Medical Device Packaging Validations	Karen Greene	RECORDED	2016-04-28	\$390.000
500794	Accelerated Aging Techniques for Medical Device Packaging	Karl J Hemmerich	RECORDED	2016-04-27	\$390.000
500795	Payroll Updates 2016	David Sanders	RECORDED	2016-04-18	\$390.000
500798	Conducting Observational Studies in US, Canada and Europe	Anne Tomalin	RECORDED	2016-04-26	\$390.000
500799	Clinical Trials in US, Europe and Canada	Anne Tomalin	RECORDED	2016-05-26	\$390.000
500800	Current Status of Biosimilars in US, Europe and Canada	Anne Tomalin	RECORDED	2016-06-15	\$390.000

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500801	Carrier Food Safety Problems That May Occur during Transportation (FDA-FSMA)	John Ryan	RECORDED	2016-04-04	\$190.000
500806	Conducting Successful Product Complaint Investigations	David Dills	RECORDED	2016-04-05	\$190.000
500807	Complaint Handling and Management: From Receipt to Trending	David Dills	RECORDED	2016-05-04	\$390.000
500808	Construct and Manage the Technical File and Design Dossier	David Dills	RECORDED	2016-05-23	\$390.000
500809	The Master Validation Plan - The Unwritten Requirements	John E Lincoln	RECORDED	2016-04-13	\$190.000
500810	DHF, DMR, DHR, Technical File and Design Dossier - Key Requirements and Future Directions	John E Lincoln	RECORDED	2016-04-26	\$390.000
500811	Device Changes, FDA Changes, and the 510(k)	John E Lincoln	RECORDED	2016-05-04	\$390.000
500812	Drafting a Software V&V Documentation Package and Protocol	John E Lincoln	RECORDED	2016-05-25	\$390.000
500816	The Most Common Problems in FDA Software Validation & Verification	John E Lincoln	RECORDED	2016-10-19	\$390.000
500817	Change Control - The Achilles Heel of cGMP Compliance	John E Lincoln	RECORDED	2016-09-14	\$390.000
500821	CAPA, Failure Investigation and Root Cause Analysis to Meet FDA Expectations	John E Lincoln	RECORDED	2016-11-21	\$390.000
500824	Combination Products: FDA's Final Rule for GMP Requirements	David Dills	RECORDED	2016-05-26	\$390.000
500826	Importing and Exporting Medical Devices: A Primer on Regulatory Strategy and Requirements	David Dills	RECORDED	2016-06-22	\$390.000
500827	Update on Unique Device Identifier for Device Manufacturers	David Dills	RECORDED	2016-06-30	\$390.000
500831	Managing Your Medical Device Reporting (MDR) Program for Compliance Success	David Dills	RECORDED	2016-08-24	\$390.000
500836	Postmarketing Vigilance Reporting For Medical Device Manufacturers	David Dills	RECORDED	2016-11-03	\$390.000
500837	Maintaining an Effective CAPA Program and Using Risk Assessment Tools: Current Trends	David Dills	RECORDED	2016-11-10	\$390.000
500843	Setting the ideal cGMP HVAC Design Requirements for Pharmaceutical Sterile and Sterile Facilities	Majdi Ayoub	RECORDED	2016-06-29	\$390.000
500847	Laboratory-Developed Tests: Why does FDA think They Can Regulate Them, and Why Do Others Think They Cannot	Anna Longwell	RECORDED	2016-05-24	\$390.000
500848	FDA's Medical Device Clinical Trials Program	Anna Longwell	RECORDED	2016-06-08	\$290.000
500849	A second look at 510(k) changes	Anna Longwell	RECORDED	2016-06-28	\$390.000

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500855	Managing Your FDA "483" Inspectional Observations	Casper Uldriks	RECORDED	2016-05-24	\$390.000
500856	Contract Killer Clauses in Construction: And How to Neutralize Them	Brian Perlberg	RECORDED	2016-05-24	\$390.000
500860	FDA Export Certificates for Medical Devices	Larry Spears	RECORDED	2016-07-05	\$390.000
500864	SOPs for Clinical Trials: Regulatory Requirements	Harold Thibodeaux	RECORDED	2016-06-02	\$390.000
500867	Combination Products: A Regulatory Perspective	Thomas E. Colonna	RECORDED	2016-08-25	\$390.000
500868	Understanding, Calculating, and Using Statistical Power	John N. Zorich	RECORDED	2016-06-30	\$390.000
500871	Good Laboratory Practice Regulations - Introduction and Strategies for Implementation	Dr. Ludwig Huber	RECORDED	2016-05-18	\$390.000
500872	Understanding and Preparing for FDA's on-going Part 11 Inspection Program	Dr. Ludwig Huber	RECORDED	2016-06-22	\$390.000
500873	Japan Regulatory Approval Process for BioPharma and Medical Devices	Robert J. Russell	RECORDED	2016-08-11	\$390.000
500874	Water System Biofilm Control and Microbial Monitoring Myths	Teri C. Soli	RECORDED	2016-06-23	\$390.000
500875	GMP Expectations for Products Used in Early Phase IND Studies	Steven S. Kuwahara	RECORDED	2016-07-19	\$390.000
500877	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	RECORDED	2016-07-14	\$390.000
500878	Understanding the USP Chapter 1224 for Transfer of Analytical Methods	Dr. Ludwig Huber	RECORDED	2016-07-13	\$390.000
500879	Validation and Use of Excel® Spreadsheets in FDA Regulated Environments	Dr. Ludwig Huber	RECORDED	2016-08-10	\$390.000
500880	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2016-06-21	\$390.000
500881	Good Documentation Practice for GxP Environments	John R. Godshalk	RECORDED	2016-06-14	\$390.000
500883	Lyophilization: What you Need to Know, Validation and Regulatory Approaches	John R. Godshalk	RECORDED	2016-10-04	\$390.000
500884	Pharmaceutical and Biologics Facility Design: FDA and Regulatory Aspects	John R. Godshalk	RECORDED	2016-12-06	\$390.000
500891	Complaint Handling in Compliance with FDA and ISO Regulations	Jeff Kasoff	RECORDED	2016-06-15	\$390.000
500893	2011 FDA Guideline on Process Validation	Louis Angelucci	RECORDED	2016-06-23	\$390.000
500895	Validation Master Planning and Regulatory Expectations	Louis Angelucci	RECORDED	2016-08-25	\$390.000
500897	Applying Statistical Process Control Effectively	Steven Wachs	RECORDED	2016-07-11	\$390.000

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500899	The Role of FDA in Health Care Software Regulations and Development	Anna Longwell	RECORDED	2016-07-13	\$390.000
500904	HPLC Analytical Method Development and Validation	John C. Fetzer	RECORDED	2016-08-31	\$390.000
500906	Direct-to-Consumer (DTC) Marketing: Impacts and Policy Implications	Casper Uldriks	RECORDED	2016-12-06	\$390.000
500914	Secrets to Writing Effective SOPs for Medical Device QMS	Susanne Manz	RECORDED	2016-07-06	\$390.000
500917	Addressing CAPA within a Device Quality System	Susanne Manz	RECORDED	2016-08-23	\$390.000
500919	Risk Management Techniques for Medical Devices	Susanne Manz	RECORDED	2016-09-27	\$390.000
500922	GLPs: How are they Associated with GMPs and SOPs	Joy McElroy	RECORDED	2016-07-20	\$390.000
500925	Responsibilities of the Carrier Under the Final FDA FSMA Rules on the Sanitary Transportation of Human and Animal Foods	John Ryan	RECORDED	2016-07-06	\$390.000
500929	Laboratory Accreditation: Getting There is Just the Beginning	Michael Brodsky	RECORDED	2016-07-11	\$390.000
500930	Is it Method Verification or Validation, or Just Semantics	Michael Brodsky	RECORDED	2016-07-28	\$390.000
500936	Laboratory-Developed Tests: Why does FDA think They Can Regulate Them, and Why Do Others Think They Cannot	Anna Longwell	RECORDED	2016-08-11	\$390.000
500937	Process Capability Analysis by means of Confidence Reliability Calculations	John N. Zorich	RECORDED	2016-07-26	\$390.000
500938	Process Capability Analysis Of Extremely Non-normal Data	John N. Zorich	RECORDED	2016-08-11	\$390.000
500939	Managing Your FDA "483" Inspectional Observations	Casper Uldriks	RECORDED	2016-07-12	\$390.000
500942	Modeling and Optimizing Process Behavior using Design of Experiments	Steven Wachs	RECORDED	2016-07-19	\$390.000
500943	FDA Perspective on International Clinical Trials: US, EU and Canada	Anne Tomalin	RECORDED	2016-07-21	\$390.000
500952	Accelerated Aging Techniques for Medical Device Packaging	Karl Leinsing	RECORDED	2016-08-30	\$390.000
500953	Fundamentals Of Technology Transfer And Intellectual Property Licensing	Ronald Kudla	RECORDED	2016-08-04	\$390.000
500954	Mastering Excel: How to Create ad hoc and Date Based Groupings within a PivotTable	Dennis Taylor	RECORDED	2016-07-18	\$390.000
500956	Why use IRS e-services TIN Matching Program	Greta Hicks	RECORDED	2016-08-22	\$390.000
500957	Determining Sample Size: How to Ensure You Get the Correct One	Steven Wachs	RECORDED	2016-08-08	\$390.000

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500963	Time-Saving Excel Tips, Tricks and 100 Shortcuts	Dennis Taylor	RECORDED	2016-08-16	\$390.000
500965	Excel's Data Management Tools	Dennis Taylor	RECORDED	2016-10-06	\$390.000
500966	Successful Through Wall Flashing Systems	Leonard Anastasi	RECORDED	2016-08-18	\$390.000
500969	Water System Biofilm Control and Microbial Monitoring Myths	Teri C. Soli	RECORDED	2016-09-12	\$390.000
500970	Using Statistics to Determine Sample Size	Steven Walfish	RECORDED	2016-09-20	\$390.000
500971	Statistical Power and Sample Size	John N. Zorich	RECORDED	2016-08-30	\$390.000
500972	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2016-09-19	\$390.000
500976	21 CFR Part 820 - Quality System Regulation - Applying Principles of Lean Documents and Lean Configuration	Jose Mora	RECORDED	2016-09-14	\$390.000
500977	Design History File (DHF), the Device Master Record (DMR) and the Device History Record (DHR) – Principles on Lean Documents and Lean Configuration	Jose Mora	RECORDED	2016-10-05	\$390.000
500978	Good Documentation Practice for GxP Environments	John R. Godshalk	RECORDED	2016-10-12	\$390.000
500979	Implementing the New FDA Data Integrity Guide	Dr. Ludwig Huber	RECORDED	2016-09-08	\$390.000
500980	Understanding and Preparing for FDA's on-going Part 11 Inspections	Dr. Ludwig Huber	RECORDED	2016-10-04	\$390.000
500981	Good Laboratory Practice Regulations - Introduction and Strategies for Implementation	Dr. Ludwig Huber	RECORDED	2016-11-02	\$390.000
500984	FDA's 21 CFR 11 Add-On Inspections - Recent Updates	Angela Bazigos	RECORDED	2016-10-05	\$390.000
500985	FDA's Medical Device Clinical Trials Program	Anna Longwell	RECORDED	2016-10-11	\$390.000
500988	Applying Statistical Process Control Effectively	Steven Wachs	RECORDED	2016-10-10	\$390.000
500992	Laboratory Investigation of Out-of-Specification Results	Jerry Lanese	RECORDED	2016-12-05	\$390.000
501003	Understanding FDA's Quality Metrics Draft Guidance and Its Impact	Louis Angelucci	RECORDED	2016-10-06	\$390.000
501019	Determining Sample Size: How to Ensure You Get the Correct	Steven Wachs	RECORDED	2016-11-14	\$390.000
501020	Time-Saving Excel Tips, Tricks and 100 Shortcuts	Dennis Taylor	RECORDED	2016-10-17	\$390.000
501022	The Bayh-Dole Act and Its Implementing Regulations & Rule-Making	Ronald Kudla	RECORDED	2016-10-19	\$390.000
501026	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	RECORDED	2016-10-11	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501030	FDA - Legal Writing Skills that Result in Effective Regulatory Submissions	Robert Michalik	RECORDED	2016-10-06	\$390.000
501047	FDA Issues Final Rule on Symbols in Labeling	David Dills	RECORDED	2016-10-26	\$390.000
501048	Understanding, Calculating, and Using Statistical Power	John N. Zorich	RECORDED	2016-11-01	\$390.000
501049	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2016-11-08	\$390.000
501050	Process Capability Analysis by means of Confidence Reliability Calculations	John N. Zorich	RECORDED	2016-12-06	\$390.000
501053	Device Changes, FDA Changes, and the 510(k)	John E Lincoln	RECORDED	2017-01-25	\$390.000
501054	The Master Validation Plan - The Unwritten Requirements	John E Lincoln	RECORDED	2017-02-15	\$390.000
501055	Water System Biofilm Control and Microbial Monitoring Myths	Teri C. Soli	RECORDED	2016-12-08	\$390.000
501056	Japan - Regulatory Filing Requirements and Compliance Processes for Life Sciences	Robert J. Russell	RECORDED	2016-11-28	\$390.000
501057	Drug Master Files: Understanding and Meeting your Regulatory and Processing Responsibilities	Robert J. Russell	RECORDED	2016-12-06	\$390.000
501060	Latin America Regulatory Compliance: Current Challenges in Argentina, Brazil and Mexico	Robert J. Russell	RECORDED	2017-02-14	\$390.000
501062	FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	RECORDED	2016-11-23	\$390.000
501063	Transfer of Analytical Methods and Procedures: FDA Requirements and Strategies and Tools for Implementation	Dr. Ludwig Huber	RECORDED	2017-01-12	\$390.000
501064	Validation and Use of Excel® Spreadsheets in FDA Regulated Environments	Dr. Ludwig Huber	RECORDED	2017-01-04	\$390.000
501075	FDA - Legal Writing Skills that Result in Effective Regulatory Submissions	Robert Michalik	RECORDED	2016-12-14	\$390.000
501077	Chemistry 101 for Medical Device Regulatory & Quality Professionals: Essential knowledge needed to Manage Drug&Device Combination Product Project	Robert Michalik	RECORDED	2017-01-25	\$390.000
501080	Design History File (DHF), Device Master Record (DMR), Device History Record (DHR)	Jose Mora	RECORDED	2017-01-17	\$390.000
501081	Risk Management Utilizing Lean Documents and Lean Configuration	Jose Mora	RECORDED	2017-02-08	\$390.000
501089	Effective Training Practices for FDA Compliance	Dr. Ludwig Huber	RECORDED	2017-01-26	\$390.000
501090	Validation of Analytical Methods and Procedures	Dr. Ludwig Huber	RECORDED	2017-03-15	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501092	Understanding and Implementing USP 1058 Analytical Instrument Qualification	Dr. Ludwig Huber	RECORDED	2017-04-19	\$390.000
501093	FDA Inspections – Do's and Don'ts	Jeff Kasoff	RECORDED	2016-12-14	\$390.000
501095	FDA Regulation of Dietary Supplements	Thomas E. Colonna	RECORDED	2017-02-28	\$390.000
501099	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	RECORDED	2017-01-24	\$390.000
501100	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2017-02-07	\$190.000
501102	Understanding, Calculating, and Using Statistical Power	John N. Zorich	RECORDED	2017-03-07	\$390.000
501106	Validating Radiation Sterilization for Medical Products	Karl J Hemmerich	RECORDED	2017-01-25	\$390.000
501107	Accelerated Aging Techniques for Medical Device Packaging	Karl J Hemmerich	RECORDED	2017-02-02	\$390.000
501110	Good Documentation Practice for GxP Environments	John R. Godshalk	RECORDED	2017-01-10	\$390.000
501114	Good Deviation Practice: What you Need to Know	John R. Godshalk	RECORDED	2017-05-10	\$390.000
501116	GLPs: How are they Associated with GMPs and SOPs	Joy McElroy	RECORDED	2017-01-20	\$390.000
501125	Verification or Validation of Methods in Food Microbiology	Michael Brodsky	RECORDED	2017-03-21	\$390.000
501128	Laboratory-Developed Tests: Why does FDA think They Can Regulate Them, and Why Do Others Think They Cannot	Anna Longwell	RECORDED	2017-01-25	\$390.000
501129	The Role of FDA in Health Care Software Regulations and Development	Anna Longwell	RECORDED	2017-02-22	\$390.000
501132	21 CFR Part 11 - Compliance for Electronic Records and Signatures	Edwin Waldbusser	RECORDED	2017-01-18	\$390.000
501133	Software Validation for the New FDA Inspections	Edwin Waldbusser	RECORDED	2017-02-02	\$390.000
501148	Get Organised with Microsoft Outlook	Mike Thomas	RECORDED	2017-03-09	\$390.000
501149	Mastering Excel Formulas and Functions	Mike Thomas	RECORDED	2017-03-28	\$390.000
501155	Death By CAPA - Does Your CAPA Program Need a CAPA?	Susanne Manz	RECORDED	2017-02-03	\$390.000
501157	Quality is not an Organization	Susanne Manz	RECORDED	2017-02-07	\$390.000
501158	Essentials of Validation -IQ,OQ,PQ	Susanne Manz	RECORDED	2017-02-22	\$390.000
501161	Verification vs. Validation-Product,Process or Equipment and QMS Software	John E Lincoln	RECORDED	2017-03-08	\$390.000
501162	Modeling and Optimizing Process Behavior using Design of Experiments	Steven Wachs	RECORDED	2017-01-27	\$390.000
501169	CAPA Training and Causes of Warning Letters due to Lack of Comprehension	Jerry Dalfors	RECORDED	2017-02-01	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501178	Time-Saving Excel Tips, Tricks and 100 Shortcuts	Dennis Taylor	RECORDED	2017-01-23	\$390.000
501184	FDA's Tougher Import Program	Casper Uldriks	RECORDED	2017-02-15	\$390.000
501185	Understanding FDA's Quality Metrics Draft Guidance and Its Impact	Louis Angelucci	RECORDED	2017-01-19	\$390.000
501186	2011 FDA Guideline on Process Validation	Louis Angelucci	RECORDED	2017-01-26	\$390.000
501187	Validation Master Planning and Regulatory Expectations	Louis Angelucci	RECORDED	2017-02-09	\$390.000
501188	Similarities and Differences Between an FDA and MHRA Audit	Louis Angelucci	RECORDED	2017-02-23	\$390.000
501190	Understanding Proper Application of Risk Assessment	Louis Angelucci	RECORDED	2017-03-16	\$390.000
501210	From Project Manager to Project Leader	Heath Suddleson	RECORDED	2017-03-21	\$390.000
501216	Implementing an Effective CAPA System	Charles H. Paul	RECORDED	2017-03-23	\$390.000
501217	FDA - Legal Writing Skills that Result in Effective Regulatory Submissions	Robert Michalik	RECORDED	2017-05-03	\$390.000
501236	Food Safety Modernization Act	Thomas Perkins	RECORDED	2017-04-19	\$390.000
501238	Determining Sample Size: What Sample Size Should I Use?	Steven Wachs	RECORDED	2017-04-18	\$390.000
501239	Design History File (DHF), the Device Master Record (DMR) and the Device History Record (DHR) - Principles on Lean Documents and Lean Configuration	Jose Mora	RECORDED	2017-05-18	\$390.000
501240	Japan - Regulatory Filing Requirements and Compliance Processes for Life Sciences	Robert J. Russell	RECORDED	2017-04-11	\$390.000
501243	Water System Biofilm Control and Microbial Monitoring Myths	Teri C. Soli	RECORDED	2017-04-13	\$390.000
501244	DHF, DMR, DHR, Technical File and Design Dossier - Key Requirements and Future Directions	John E Lincoln	RECORDED	2017-04-18	\$390.000
501248	The FDA Inspection: Preparation, Management, and Follow - up	Jeff Kasoff	RECORDED	2017-04-03	\$390.000
501250	Accelerated Aging Techniques for Medical Device Packaging	Karl J Hemmerich	RECORDED	2017-04-05	\$390.000
501255	Using Statistics to Determine Sample Size	Steven Walfish	RECORDED	2017-04-18	\$390.000
501264	Supplier Evaluation and Assessment: How to Meet FDA QSR and ISO 13485 Requirements in a Cost-Effective Manner	Jeff Kasoff	RECORDED	2017-05-01	\$390.000
501265	How FDA Trains its Investigators to Review CAPA and What You Should Do to Prepare	Jeff Kasoff	RECORDED	2017-05-15	\$390.000
501266	Equipment Validation, Tracking, Calibration and Preventive Maintenance	Jeff Kasoff	RECORDED	2017-06-02	\$390.000



Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501273	Auditing Analytical Laboratories for FDA Compliance	Steven S. Kuwahara	RECORDED	2017-05-03	\$390.000
501276	Validation and Control of Excel Spreadsheets in FDA Regulated Environments	Dr. Ludwig Huber	RECORDED	2017-04-27	\$390.000
501277	New FDA or EMA and USP Guidelines for Transfer of Analytical Methods	Dr. Ludwig Huber	RECORDED	2017-05-18	\$390.000
501278	Laboratory-Developed Tests: Why does FDA Think They Can Regulate Them, and Why Do Others Think They Cannot	Anna Longwell	RECORDED	2017-05-26	\$390.000
501280	FDA's New Import Program for 2017	Casper Uldriks	RECORDED	2017-06-20	\$390.000
501283	Drug Master Files: Understanding and Meeting your Regulatory and Processing Responsibilities	Robert J. Russell	RECORDED	2017-06-05	\$390.000
501284	Latin America Regulatory Compliance: Current Challenges in Argentina, Brazil and Mexico	Robert J. Russell	RECORDED	2017-05-11	\$390.000
501287	Controlling Human Error in the Manufacturing Floor	Ginette Collazo	RECORDED	2017-08-08	\$390.000
501288	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	RECORDED	2017-05-09	\$390.000
501289	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2017-05-30	\$390.000
501293	Master Excel: Spreadsheet Internal Controls	David Ringstrom	RECORDED	2017-06-12	\$390.000
501296	2011 FDA Guideline on Process Validation	Louis Angelucci	RECORDED	2017-07-13	\$390.000
501305	Time-Saving Excel Tips, Tricks and 100 Shortcuts	Dennis Taylor	RECORDED	2017-05-10	\$390.000
501306	21 CFR Part 11 - Compliance for Electronic Records and Signatures	Edwin Waldbusser	RECORDED	2017-05-17	\$390.000
501307	Software Validation for the New FDA Inspections	Edwin Waldbusser	RECORDED	2017-06-08	\$390.000
501323	Bullet Proof 510(k) - Latest FDA Changes to the Process	David Dills	RECORDED	2017-08-09	\$390.000
501333	Handling OOS Test Results and Completing Robust Investigations	Danielle DeLucy	RECORDED	2017-05-08	\$390.000
501336	Successful Supplier Audits	Danielle DeLucy	RECORDED	2017-07-12	\$390.000
501346	FDA Enforcement of 21 CFR 11 Compliance	Angela Bazigos	RECORDED	2017-05-09	\$390.000
501347	21 CFR 11 Compliance for Excel Spreadsheets	Angela Bazigos	RECORDED	2017-05-23	\$390.000
501354	Understanding, Calculating, and Using Statistical Power	John N. Zorich	RECORDED	2017-06-13	\$390.000
501355	Process Capability Analysis by means of Confidence Reliability Calculations	John N. Zorich	RECORDED	2017-06-27	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501361	Implementing Agile in an FDA-Regulated Environment	Brian Shoemaker	RECORDED	2017-07-13	\$390.000
501362	How to Prepare for and Host a FDA Inspection and Respond to 483's	Edwin Waldbusser	RECORDED	2017-08-08	\$390.000
501365	Statistical Concepts of Process Validation	Dan OLeary	RECORDED	2017-07-05	\$390.000
501369	Japan - Regulatory Filing Requirements and Compliance Processes for Life Sciences	Robert J. Russell	RECORDED	2017-06-29	\$390.000
501373	Understanding and Applying ICH Q3A and Q3B for Control of Impurities in Drug Substances and Drug Products	Greg Martin	RECORDED	2017-08-17	\$390.000
501380	FDA - Legal Writing Skills that Result in Effective Regulatory Submissions	Robert Michalik	RECORDED	2017-07-13	\$390.000
501381	Accelerated Aging Techniques for Medical Device Packaging	Karl J Hemmerich	RECORDED	2017-06-21	\$390.000
501384	Preventing and Detecting Fraudulent Vendor Disbursements	John E. Grimes	RECORDED	2017-07-27	\$390.000
501386	FDA Inspections - Do's and Don'ts	Jonathan M. Lewis	RECORDED	2017-07-14	\$390.000
501388	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	RECORDED	2017-07-07	\$390.000
501389	Understanding the USP Chapter 1224 for Transfer of Analytical Methods	Dr. Ludwig Huber	RECORDED	2017-06-27	\$390.000
501393	Latin America Regulatory Compliance: Current Challenges in Argentina, Brazil and Mexico	Robert J. Russell	RECORDED	2017-07-24	\$390.000
501394	Determining Sample Size: What Sample Size Should I Use?	Steven Wachs	RECORDED	2017-07-11	\$390.000
501395	Understanding and Implementing USP 1058 Analytical Instrument Qualification	Dr. Ludwig Huber	RECORDED	2017-08-07	\$390.000
501396	Auditing Analytical Laboratories for FDA Compliance	Steven S. Kuwahara	RECORDED	2017-06-30	\$390.000
501398	Device Changes, FDA Changes, and the 510(k)	John E Lincoln	RECORDED	2017-06-28	\$390.000
501399	Using Statistics to Determine Sample Size	Steven Walfish	RECORDED	2017-07-18	\$390.000
501400	Key Factors to Write an Effective Standard Operating Procedure (SOP) and Work Instructions (WIs)	Angela Bazigos	RECORDED	2017-06-29	\$390.000
501401	FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	RECORDED	2017-07-20	\$390.000
501402	Validation and Use of Excel® Spreadsheets in FDA Regulated Environments	Dr. Ludwig Huber	RECORDED	2017-06-23	\$390.000
501404	FDA Enforcement of 21 CFR 11 Compliance	Angela Bazigos	RECORDED	2017-07-07	\$390.000
501405	Water System Biofilm Control and Microbial Monitoring Myths	Teri C. Soli	RECORDED	2017-07-07	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501406	Time-Saving Excel Tips, Tricks and 100 Shortcuts	Dennis Taylor	RECORDED	2017-07-07	\$390.000
501407	Death By CAPA - Does your CAPA Program Need a CAPA?	Susanne Manz	RECORDED	2017-07-11	\$390.000
501410	Design History Files (DHF), Device Master Records (DMR), Device History Records (DHR), Technical Files , Design Dossiers - The Requirements	John E Lincoln	RECORDED	2017-07-11	\$390.000
501411	21 CFR Part 11 - Compliance for Electronic Records and Signatures	Edwin Waldbusser	RECORDED	2017-06-22	\$390.000
501419	Laboratory-Developed Tests: Why does FDA think they can Regulate them, and why do others think they Cannot	Anna Longwell	RECORDED	2017-08-10	\$390.000
501420	FDA's Medical Device Clinical Trials Program	Anna Longwell	RECORDED	2017-08-24	\$390.000
501422	GDPR - European Privacy Regulation what and how	Derk Yntema	RECORDED	2017-08-30	\$390.000
501423	Secrets to Writing Effective SOPs for Medical Device QMS	Susanne Manz	RECORDED	2017-08-04	\$390.000
501426	Essentials of Validation -IQ,OQ,PQ	Susanne Manz	RECORDED	2017-09-14	\$390.000
501424	Addressing CAPA within a Device Quality System	Susanne Manz	RECORDED	2017-08-09	\$390.000
501427	Quality is not an Organization	Susanne Manz	RECORDED	2017-11-02	\$390.000
501431	Medical Device Cybersecurity : The Landscape of Quicksand	Casper Uldriks	RECORDED	2017-07-12	\$390.000
501435	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2017-08-01	\$390.000
501436	Understanding, Calculating, and Using Statistical Power	John N. Zorich	RECORDED	2017-08-21	\$390.000
501437	HPLC Analytical Method Development and Validation	John C. Fetzer	RECORDED	2017-08-01	\$390.000
501439	Analytical Method Validation Under Good Laboratory Practices - GLPs	John C. Fetzer	RECORDED	2017-09-12	\$390.000
501440	FDA's New Import Program for 2017	Casper Uldriks	RECORDED	2017-08-01	\$390.000
501441	Dietary Supplement Regulatory Compliance in the United States: Labeling, Product Claims & Updates from the FDA	James E. Russell	RECORDED	2017-10-05	\$390.000
501442	CAPA - An important Element of the Pharmaceutical Quality System	Jerry Lanese	RECORDED	2017-10-04	\$390.000
501446	Verification vs Validation-Product, Process or Equipment and QMS Software	John E Lincoln	RECORDED	2017-09-05	\$390.000
501447	Mastering Excel Formulas and Functions	Mike Thomas	RECORDED	2017-08-03	\$390.000
501449	OneNote - Bringing Order to your Digital Chaos	Mike Thomas	RECORDED	2017-09-08	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501450	Excel - Business Intelligence with Power Pivot and Power Query	Mike Thomas	RECORDED	2017-09-22	\$390.000
501451	Project Management for 21 CFR 11	Angela Bazigos	RECORDED	2017-10-20	\$390.000
501452	Change Control,CAPA,Document Control, and Electronic Systems	Deb Simpson	RECORDED	2017-08-30	\$390.000
501458	Drug Master Files: Understanding and Meeting your Regulatory and Processing Responsibilities	Robert J. Russell	RECORDED	2017-09-08	\$390.000
501469	Implementation and Management of GMP Data Integrity	Danielle DeLucy	RECORDED	2017-10-23	\$390.000
501472	Batch Record Review and Product Release	Danielle DeLucy	RECORDED	2017-10-13	\$390.000
501479	Lean Validation: Leveraging the NIST Cybersecurity Framework for Computer Systems Validation	Valarie King Bailey	RECORDED	2017-10-25	\$390.000
501481	Design History Files (DHF), Device Master Records (DMR), Device History Records (DHR), Technical Files , Design Dossiers - The Requirements	John E Lincoln	RECORDED	2017-08-28	\$390.000
501483	Auditing Analytical Laboratories for FDA Compliance	Steven S. Kuwahara	RECORDED	2017-09-12	\$390.000
501484	Device Changes, FDA Changes, and the 510(k)	John E Lincoln	RECORDED	2017-08-22	\$390.000
501485	Using Statistics to Determine Sample Size	Steven Walfish	RECORDED	2017-08-14	\$390.000
501486	Key Factors to Write an Effective Standard Operating Procedure (SOP) and Work Instructions (WIs)	Angela Bazigos	RECORDED	2017-08-16	\$390.000
501487	Validation and Use of Excel® Spreadsheets in FDA Regulated Environments	Dr. Ludwig Huber	RECORDED	2017-08-29	\$390.000
501488	FDA's 21 CFR 11 Add-On Inspections - Recent Updates	Angela Bazigos	RECORDED	2017-10-03	\$390.000
501489	Water System Biofilm Control and Microbial Monitoring Myths	Teri C. Soli	RECORDED	2017-08-25	\$390.000
501490	Time-Saving Excel Tips, Tricks and 100 Shortcuts	Dennis Taylor	RECORDED	2017-08-25	\$390.000
501491	Death By CAPA - Does your CAPA Program Need a CAPA?	Susanne Manz	RECORDED	2017-08-28	\$390.000
501493	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	RECORDED	2017-08-11	\$390.000
501494	Understanding the USP Chapter 1224 for Transfer of Analytical Methods	Dr. Ludwig Huber	RECORDED	2017-09-07	\$390.000
501495	Similarities and Differences Between an FDA and MHRA Audit	Louis Angelucci	RECORDED	2017-09-11	\$390.000
501496	Mastering Excel 2013 Pivot Tables	Dennis Taylor	RECORDED	2017-09-01	\$390.000

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501497	Determining Sample Size: What Sample Size Should I Use?	Steven Wachs	RECORDED	2017-08-28	\$390.000
501498	Excel Spreadsheet Validation for FDA 21 CFR Part 11	Angela Bazigos	RECORDED	2017-09-01	\$390.000
501499	How FDA Trains its Investigators to Review CAPA and What you Should do to Prepare	Jeff Kasoff	RECORDED	2017-09-01	\$390.000
501505	FDA Compliance and Mobile Applications	Carolyn Troiano	RECORDED	2017-09-18	\$390.000
501506	21 CFR Part 11 Compliance for Computer Systems Regulated by FDA	Carolyn Troiano	RECORDED	2017-10-10	\$390.000
501507	FDA's New Draft Guidance on Software and Device Changes and the 510(k)	Carolyn Troiano	RECORDED	2017-10-30	\$390.000
501514	Medical Device Cybersecurity : The Landscape of Quicksand	Casper Uldriks	RECORDED	2017-11-14	\$390.000
501515	FDA's New Import Program for 2017	Casper Uldriks	RECORDED	2017-09-25	\$390.000
501521	Agile for Medical Devices: More than Just Software	Brian Shoemaker	RECORDED	2017-10-31	\$390.000
501525	Japan - Regulatory Filing Requirements and Compliance Processes for Life Sciences	Robert J. Russell	RECORDED	2017-09-21	\$390.000
501526	Latin America Regulatory Compliance: Current Challenges in Argentina, Brazil and Mexico	Robert J. Russell	RECORDED	2017-10-16	\$390.000
501530	Design History File (DHF), the Device Master Record (DMR) and the Device History Record (DHR) - Principles on Lean Documents and Lean Configuration	Jose Mora	RECORDED	2017-10-18	\$390.000
501532	How to Prepare for and Host a FDA Inspection and Respond to 483's	Edwin Waldbusser	RECORDED	2017-10-31	\$390.000
501534	Managing Client Expectations Without Losing your Shirt or the Next Job	Heath Suddleson	RECORDED	2017-09-18	\$390.000
501541	Accelerated Aging Techniques for Medical Device Products	Karl J Hemmerich	RECORDED	2017-09-20	\$390.000
501543	Key Steps to Successful Project Management Principles	Carol Friedhoff	RECORDED	2017-10-17	\$390.000
501552	Controlling Human Error in the Manufacturing Floor	Ginette Collazo	RECORDED	2017-10-17	\$390.000
501559	3-hour Virtual Seminar on Analytical Method Validation Process	Angela Bazigos	RECORDED	2017-11-03	\$540.000
501560	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	RECORDED	2017-09-25	\$390.000
501561	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2017-10-09	\$390.000
501570	Key Factors to Write an Effective Standard Operating Procedure (SOP) and Work Instructions (WIs)	Angela Bazigos	RECORDED	2017-10-11	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501571	Using Statistics to Determine Sample Size	Steven Walfish	RECORDED	2017-11-20	\$390.000
501573	Water System Biofilm Control and Microbial Monitoring Myths	Teri C. Soli	RECORDED	2017-09-29	\$390.000
501574	Time-Saving Excel Tips, Tricks and 100 Shortcuts	Dennis Taylor	RECORDED	2017-09-25	\$390.000
501578	Death By CAPA - Does your CAPA Program Need a CAPA?	Susanne Manz	RECORDED	2017-10-09	\$390.000
501576	Design History Files (DHF), Device Master Records (DMR), Device History Records (DHR), Technical Files , Design Dossiers - The Requirements	John E Lincoln	RECORDED	2017-10-04	\$390.000
501577	Validation and Use of Excel® Spreadsheets in FDA Regulated Environments	Dr. Ludwig Huber	RECORDED	2017-10-06	\$390.000
501580	Mastering Excel 2013 Pivot Tables	Dennis Taylor	RECORDED	2017-10-13	\$390.000
501581	Determining Sample Size: What Sample Size Should I Use?	Steven Wachs	RECORDED	2017-10-13	\$390.000
501582	Understanding and Implementing USP 1058 Analytical Instrument Qualification	Dr. Ludwig Huber	RECORDED	2017-10-23	\$390.000
501583	Essentials of Validation - IQ, OQ, PQ	Susanne Manz	RECORDED	2017-10-20	\$390.000
501587	Root Cause Analysis for CAPA - Myths, Challenges, and Best Practices	Susanne Manz	RECORDED	2018-01-08	\$390.000
501590	How to Prevent or Handle Protocol Deviations and Violations to be GCP and Regulatory Compliant	Charles H Pierce	RECORDED	2017-11-07	\$390.000
501596	Analytical Method Validation Under Good Laboratory Practices - GLPs	John C. Fetzer	RECORDED	2017-12-04	\$390.000
501597	HVAC and GMP Environmental Control for Pharmaceutical Clean Rooms	Roger Cowan	RECORDED	2017-10-17	\$390.000
501598	Pharmaceutical Compressed Air - Quality GMP Requirements - What you need to know to Meet FDA and International Quality Standards	Roger Cowan	RECORDED	2017-11-13	\$390.000
501603	Get Organised with Microsoft Outlook	Mike Thomas	RECORDED	2017-10-27	\$390.000
501604	Mastering Excel Formulas and Functions	Mike Thomas	RECORDED	2017-11-17	\$390.000
501605	OneNote - Bringing Order to your Digital Chaos	Mike Thomas	RECORDED	2017-12-08	\$390.000
501606	Accelerated Aging Techniques for Medical Device Products	Karl J Hemmerich	RECORDED	2017-11-08	\$390.000
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501609	New FDA, EMA and USP Guidelines for Transfer of Analytical Methods	Dr. Ludwig Huber	RECORDED	2017-11-30	\$390.000

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501612	Validation and Use of Excel® Spreadsheets in FDA Regulated Environments	Dr. Ludwig Huber	RECORDED	2017-12-14	\$390.000
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501635	Time-Saving Excel Tips, Tricks and 100 Shortcuts	Dennis Taylor	RECORDED	2017-11-21	\$390.000
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501648	CAPA, Failure Investigation and Root Cause Analysis to Meet FDA Expectations	John E Lincoln	RECORDED	2018-01-17	\$390.000
501652	Excel Spreadsheet Validation for FDA 21 CFR Part 11	Angela Bazigos	RECORDED	2017-12-06	\$390.000
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501671	Death by CAPA - Does your Company have the Symptoms?	Susanne Manz	RECORDED	2018-01-18	\$390.000
501680	Batch Record Review and Product Release	Danielle DeLucy	RECORDED	2018-01-18	\$390.000
501693	FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	RECORDED	2018-01-09	\$390.000
501696	Mastering Excel 2013 Pivot Tables	Dennis Taylor	RECORDED	2018-01-18	\$390.000
501700	Validation of Analytical Methods and Procedures	Dr. Ludwig Huber	RECORDED	2018-01-18	\$390.000
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