

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
502169	Fundamentals of Calibration and Validation of Laboratory and Process Analysis Instrumentation	Emil W Ciurczak	LIVE RECORDED	2018-10-19	\$150.000 \$390.000
502195	The New ISO 13485:2016 and Comparison with 21CFR820 - How to Comply with both in the Same Organization	Eyal Lerner	LIVE RECORDED	2018-10-19	\$150.000 \$390.000
502165	3-Hour Virtual Seminar - Good Pharmacovigilance Practice	Amer Alghabban	LIVE RECORDED	2018-10-19	\$290.000 \$540.000
502197	Understanding the USP Chapter 1224 for Transfer of Analytical Methods	Dr. Ludwig Huber	LIVE RECORDED	2018-10-22	\$150.000 \$390.000
502167	2-Hour Virtual Seminar on Data Integrity for GXP Compliance	Laura Brown	LIVE RECORDED	2018-10-22	\$190.000 \$440.000
502136	BREXIT - What's Changing for Life Science Product License Holders- Manufacturers and What you Need to do Right Now	Robert J. Russell	LIVE RECORDED	2018-10-22	\$150.000 \$390.000
502163	The 21st Century Cures Act - Expediting New Products and New Product Indications	Carolyn Troiano	LIVE RECORDED	2018-10-22	\$150.000 \$390.000
502140	HVAC and GMP Environmental Control - for Pharmaceutical Cleanrooms	Roger Cowan	LIVE RECORDED	2018-10-23	\$150.000 \$390.000
502183	3-Hour Virtual Seminar on Master Validation Plan - The Unwritten Requirements	John E Lincoln	LIVE RECORDED	2018-10-23	\$290.000 \$540.000
502178	3-Hour Virtual Seminar on Analytical Method Validation Process	Angela Bazigos	LIVE RECORDED	2018-10-23	\$290.000 \$540.000
502115	Computer System Validation	Edwin Waldbusser	LIVE RECORDED	2018-10-23	\$150.000 \$390.000
502095	Validation of Non-Product Software	Thomas Bento	LIVE RECORDED	2018-10-24	\$150.000 \$390.000
501982	Training in Human Error: Reducing Training Related Errors	Ginette Collazo	LIVE RECORDED	2018-10-24	\$150.000 \$390.000
502199	2-Hour Virtual Seminar - Biomarker Development and Validation - A Statistical Perspective	Dr. Shuguang Huang	LIVE RECORDED	2018-10-25	\$190.000 \$440.000
502191	Time-Saving Excel Tips, Tricks and 100 Shortcuts	Dennis Taylor	LIVE RECORDED	2018-10-25	\$150.000 \$390.000
502151	Essentials of Validation - IQ, OQ, PQ	Susanne Manz	LIVE RECORDED	2018-10-25	\$150.000 \$390.000
502160	FDA's Expectations from Supplier Management for GMP: Quality Agreements and More	Danielle DeLucy	LIVE RECORDED	2018-10-25	\$150.000 \$390.000
502023	3-Hour Virtual Seminar on Deciding When to Submit a 510(k) for a Change to an Existing Device	Angela Bazigos	LIVE RECORDED	2018-10-26	\$290.000 \$540.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
502187	Trial Master File Requirements and Essential Regulatory Documents	Marina Malikova	LIVE RECORDED	2018-10-26	\$150.000 \$390.000
502093	FDA Best Audit Behavior Practices - Do's and Don'ts	Charles H. Paul	LIVE RECORDED	2018-10-29	\$150.000 \$390.000
502220	2-Hour Virtual Seminar on Valid Statistical Rationales for Sample Sizes	John N. Zorich	LIVE RECORDED	2018-10-31	\$190.000 \$440.000
502176	How to Implement Lean Documents and Lean Document Control to Cut Costs, while Maintaining Compliance with Regulatory Authority Requirements	Angela Bazigos	LIVE RECORDED	2018-10-31	\$150.000 \$390.000
502204	Pharmaceutical Drug Registration in China	Yingying Liu	LIVE RECORDED	2018-11-01	\$150.000 \$390.000
502146	Becoming a Chief Audit Executive - Leading the Internal Audit Function	Lynn Fountain	LIVE RECORDED	2018-11-01	\$150.000 \$390.000
502235	Excel Spreadsheets - Step-By-Step Instructions for Ensuring Data Integrity	David Nettleton	LIVE RECORDED	2018-11-01	\$150.000 \$390.000
502236	Cultivating a Cybersecurity Continuity Culture: 10-Step Checklist for Compliance	Carly Souther	LIVE RECORDED	2018-11-01	\$150.000 \$390.000
502207	Handling Consumer ACH Disputes: Advanced ACH Sticky Situations	Rayleen M. Pirnie	LIVE RECORDED	2018-11-01	\$150.000 \$390.000
502117	3-Hour Virtual Seminar Computer System Validation (CSV) for FDA-Regulated Computers	Carolyn Troiano	LIVE RECORDED	2018-11-02	\$290.000 \$540.000
502180	Validation Sampling Plans for Process Validation, Design Verification and CAPA Effectiveness Checks	Wayne Taylor	LIVE RECORDED	2018-11-05	\$150.000 \$390.000
502216	Validation of Analytical Methods and Procedures	Dr. Ludwig Huber	LIVE RECORDED	2018-11-06	\$150.000 \$390.000
502239	FDA's New Import Program for 2018	Casper Uldriks	LIVE RECORDED	2018-11-06	\$150.000 \$390.000
502212	Laboratory Investigation of Out-of-Specification Results	Jerry Lanese	LIVE RECORDED	2018-11-06	\$150.000 \$390.000
502188	ICH GCP Guidelines E6 Revision, R2 Addendum- Changes Impacting Sponsors-CRO-Sites	Marina Malikova	LIVE RECORDED	2018-11-06	\$150.000 \$390.000
502205	3-Hour Virtual Seminar on GDPR - European Privacy Regulation what and how	Derk Yntema	LIVE RECORDED	2018-11-07	\$290.000 \$540.000
502218	Accelerated Aging Techniques for Medical Device Packaging	Karl J Hemmerich	LIVE RECORDED	2018-11-07	\$150.000 \$390.000
502231	21 CFR Part 11 - Compliance for Electronic Records and Signatures	Edwin Waldbusser	LIVE RECORDED	2018-11-07	\$150.000 \$390.000
502152	Root Cause Analysis for CAPA - Myths, Challenges, and Best Practices	Susanne Manz	LIVE RECORDED	2018-11-08	\$150.000 \$390.000
502158	Advanced Formulas & Functions in Excel	Tom Fragale	LIVE RECORDED	2018-11-09	\$150.000 \$390.000
502200	3-Hour Virtual Seminar - Assay Validation According to CLSI Guidelines	Dr. Shuguang Huang	LIVE RECORDED	2018-11-09	\$290.000 \$540.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
502168	Good Documentation and Record Keeping Best Practices (FDA & EMA)	Dr. Afsaneh Motamed Khorasani	LIVE RECORDED	2018-11-09	\$150.000 \$390.000
502161	How to Conduct Successful Supplier Audits	Danielle DeLucy	LIVE RECORDED	2018-11-09	\$150.000 \$390.000
502164	Disaster Recovery and Business Continuity Planning for Computer Systems Regulated by FDA	Carolyn Troiano	LIVE RECORDED	2018-11-09	\$150.000 \$390.000
502185	How to Write SOPs which are Compliant for Inspections for a GXP Compliance	Laura Brown	LIVE RECORDED	2018-11-12	\$150.000 \$390.000
502222	3-Hour Virtual Seminar on 2018 Updates on FDA's Refuse to Accept Policies for 510(k)s and PMAs Including Combination Products	Angela Bazigos	LIVE RECORDED	2018-11-12	\$290.000 \$540.000
502193	2-Hour Virtual Seminar on Mastering Excel Pivot Tables	Dennis Taylor	LIVE RECORDED	2018-11-12	\$190.000 \$440.000
502211	Current Challenges in Vaccine Pharmacovigilance and Risk Mitigation Strategies	Dr. Subodh Bhardwaj	LIVE RECORDED	2018-11-12	\$150.000 \$390.000
502024	What do the FDA, EMA and PMDA Look for When Conducting cGMP Regulatory Inspections	Angela Bazigos	LIVE RECORDED	2018-11-13	\$150.000 \$390.000
502221	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	LIVE RECORDED	2018-11-14	\$150.000 \$390.000
502238	3-Hour Virtual Seminar - The FDA Deeming Rule & the Current State of Electronic Nicotine Dispensing Systems (ENDS) Regulation	Carly Souther	LIVE RECORDED	2018-11-14	\$290.000 \$540.000
502223	Key Factors to Write an Effective Standard Operating Procedure (SOP) and Work Instructions (WIs)	Angela Bazigos	LIVE RECORDED	2018-11-14	\$150.000 \$390.000
502153	Risk Management Techniques for Medical Devices	Susanne Manz	LIVE RECORDED	2018-11-15	\$150.000 \$390.000
502172	3-Hour Virtual Seminar Data Integrity in Clinical Trials	Amer Alghabban	LIVE RECORDED	2018-11-16	\$290.000 \$540.000
502170	Control-Analysis Instrumentation used in PAT and Continuous Manufacturing	Emil W Ciurczak	LIVE RECORDED	2018-11-16	\$150.000 \$390.000
502208	The Board of Directors Role in Cybersecurity	Rayleen M. Pirnie	LIVE RECORDED	2018-11-16	\$150.000 \$390.000
502150	Secrets to Writing Effective SOPs for Medical Device QMS	Susanne Manz	LIVE RECORDED	2018-11-19	\$150.000 \$390.000
502224	3-Hour Virtual Seminar on Data Integrity - FDA's Latest Thinking	Angela Bazigos	LIVE RECORDED	2018-11-19	\$290.000 \$540.000
502217	FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	LIVE RECORDED	2018-11-20	\$150.000 \$390.000
502147	Auditing and Examining Travel and Entertainment Expenses: Expense Reimbursement Fraud	Lynn Fountain	LIVE RECORDED	2018-11-27	\$150.000 \$390.000
502250	How to Validate Software for SaaS-Cloud-Local Hosting	David Nettleton	LIVE	2018-11-27	\$150.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
502229	Block Chain Made Easy - Introduction and Basics for Non-Techies	Joe Lynch	LIVE RECORDED	2018-11-27	\$150.000 \$390.000
502201	3-Hour Virtual Seminar - 21 CFR Part 11 - Guidance for Electronic Records and Electronic Signatures used in FDA-Regulated Computer Systems	Carolyn Troiano	LIVE RECORDED	2018-11-27	\$290.000 \$540.000
502194	Excel Formulas and Functions	Dennis Taylor	LIVE RECORDED	2018-11-27	\$150.000 \$390.000
502213	Analytical Method Validation	Jerry Lanese	LIVE RECORDED	2018-11-28	\$150.000 \$390.000
502225	FDA's Enforcement of 21 CFR Part 11 Compliance	Angela Bazigos	LIVE RECORDED	2018-11-28	\$150.000 \$390.000
502219	Validating Radiation Sterilization for Medical Products	Karl J Hemmerich	LIVE RECORDED	2018-11-28	\$150.000 \$390.000
502251	The Modernization of "Milk": FDA Review of Dairy Terms in Plant-Based Product Labeling	Carly Souther	LIVE RECORDED	2018-11-28	\$150.000 \$390.000
502209	Cyber Incident Response Plans: Planning for the Inevitable	Rayleen M. Pirnie	LIVE RECORDED	2018-11-28	\$150.000 \$390.000
502214	Statistics in Non-Clinical Development	Dr. Shuguang Huang	LIVE RECORDED	2018-11-29	\$150.000 \$390.000
502154	Addressing CAPA within a Device Quality System	Susanne Manz	LIVE RECORDED	2018-11-29	\$150.000 \$390.000
501984	Controlling Human Error in the Manufacturing Floor	Ginette Collazo	LIVE RECORDED	2018-11-29	\$150.000 \$390.000
502203	3-Hour Virtual Seminar on Medical Device Single Audit Program (MDSAP) Preparation	Shep Bentley	LIVE RECORDED	2018-11-30	\$290.000 \$540.000
502162	Handling OOS Test Results and Completing Robust Investigations	Danielle DeLucy	LIVE RECORDED	2018-11-30	\$150.000 \$390.000
502206	How to Audit Against ICH GCP 2 Addendum to Ensure Compliance	Laura Brown	LIVE RECORDED	2018-11-30	\$150.000 \$390.000
502232	3-Hour Virtual Seminar on Writing Effective SOPs	Angela Bazigos	LIVE RECORDED	2018-12-03	\$290.000 \$540.000
502234	Batch Record Review and Product Release	Danielle DeLucy	LIVE RECORDED	2018-12-03	\$150.000 \$390.000
502230	Block Chain Transformation: Understanding the Impacts and Applications in the Logistics Industry	Joe Lynch	LIVE RECORDED	2018-12-04	\$150.000 \$390.000
502226	Valid Statistical Rationales for Sample Sizes	John N. Zorich	LIVE RECORDED	2018-12-05	\$150.000 \$390.000
502227	Statistical Quality Control in Pharmaceutical and IVD Industries	Dr. Shuguang Huang	LIVE RECORDED	2018-12-05	\$150.000 \$390.000
502215	How to Demonstrate Vendor-CRO-CMO Oversight in the Biopharma and Device Industry	Laura Brown	LIVE RECORDED	2018-12-05	\$150.000 \$390.000
502241	Getting Meaningful Results from your Data with Excel PowerPivot and PowerQuery	Tom Fragale	LIVE RECORDED	2018-12-07	\$150.000 \$390.000

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502243	Essential Regulatory Documents and Regulatory Requirements for Biomedical Products	Marina Malikova	LIVE RECORDED	2018-12-07	\$150.000 \$390.000
502173	Theory and Practice of Near-Infrared Spectroscopy (NIRS)	Emil W Ciurczak	LIVE RECORDED	2018-12-07	\$150.000 \$390.000
502202	3-Hour Virtual Seminar - Best Practices to Help you Pass an FDA Computer System Validation Audit	Carolyn Troiano	LIVE RECORDED	2018-12-07	\$290.000 \$540.000
502252	Change Controls : How to Determine the Appropriate Levels and Reporting Categories	Lamont M. Fulton	LIVE RECORDED	2018-12-07	\$150.000 \$390.000
502242	3-Hour Virtual Seminar on Project Management for Non-Project Managers	Charles H. Paul	LIVE RECORDED	2018-12-10	\$290.000 \$540.000
502228	Seasonal Influenza Vaccines- Current Concepts and Future Trends for Effective Immunization	Dr. Subodh Bhardwaj	LIVE RECORDED	2018-12-10	\$150.000 \$390.000
502237	Data Integrity: Compliance with 21 CFR Part 11, SaaS-Cloud, EU GDPR	David Nettleton	LIVE	2018-12-11	\$150.000
502210	Handling Situations Where Regulation E and ACH Don't Align	Rayleen M. Pirnie	LIVE RECORDED	2018-12-11	\$150.000 \$390.000
502233	Excel Spreadsheets for 21 CFR 11 Compliance	Angela Bazigos	LIVE RECORDED	2018-12-12	\$150.000 \$390.000
502240	Sample Size Determination for Design Validation Activities	Steven Wachs	LIVE RECORDED	2018-12-12	\$150.000 \$390.000
502244	Audit Trail Generation and Review	Angela Bazigos	LIVE RECORDED	2019-01-08	\$150.000 \$390.000
502245	3-Hour Virtual Seminar on Excel Spreadsheet Validation for FDA 21 CFR Part 11	Angela Bazigos	LIVE RECORDED	2019-01-16	\$290.000 \$540.000
502246	If it Isn't Written Down, then it Didn't Happen: Complying with FDA's Good Documentation Practices	Angela Bazigos	LIVE RECORDED	2019-01-23	\$150.000 \$390.000
502247	Lean Document Control for Manufacturing in Life Sciences	Angela Bazigos	LIVE RECORDED	2019-02-06	\$150.000 \$390.000
502248	3-Hour Virtual Seminar on Off label Promotion of Drugs and Medical Devices - FDA's Latest	Angela Bazigos	LIVE RECORDED	2019-02-13	\$290.000 \$540.000
502249	European Data Protection Regulation - Latest	Angela Bazigos	LIVE RECORDED	2019-02-20	\$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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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
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

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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
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 Kioussis	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 R Dills	RECORDED	2014-11-24	\$189.000

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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 R Dills	RECORDED	2014-11-19	\$189.000
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 R 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 R Dills	RECORDED	2014-12-17	\$189.000
500113	Device Corrections and Removals	Dan OLeary	RECORDED	2014-12-18	\$189.000

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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
500117	Curtailing Microbial Excursions and Investigations in Pharmaceutical Water Systems	Teri C. Soli	RECORDED	2014-11-06	\$189.000
500121	Regulatory Requirements for Medical Device Calibration Programs	Dan OLeary	RECORDED	2014-11-25	\$189.000
500124	Acceptance Sampling by Variables	Dan OLeary	RECORDED	2015-02-19	\$189.000
500125	Preventing Product Liability - Key Areas Manufactures Need to Control	Randy Goodden	RECORDED	2014-12-10	\$149.000
500126	Preventing Recalls & Product Liability in New Product Development	Randy Goodden	RECORDED	2015-01-21	\$149.000
500128	Drug Master Files: Understanding and Meeting your Regulatory and Processing Responsibilities	Robert J. Russell	RECORDED	2014-11-24	\$189.000
500129	Medical Device Product Development Process	Karl Leinsing	RECORDED	2014-12-03	\$189.000
500130	Conducting Observational Studies in US, Canada and Europe	Anne Tomalin	RECORDED	2015-02-24	\$189.000
500131	Clinical Trials in US, Europe and Canada	Anne Tomalin	RECORDED	2014-12-16	\$189.000
500132	International Regulatory Cooperation Among Agencies	Anne Tomalin	RECORDED	2015-01-08	\$189.000
500134	Managing FDA 483s: Before, During and After the Inspections	Casper Uldriks	RECORDED	2014-12-10	\$189.000
500136	Rational Predictions for FDA inspections	Casper Uldriks	RECORDED	2015-02-10	\$189.000
500140	Sterile Medical Packaging Design – 7 Essentials	Karen Greene	RECORDED	2015-02-03	\$189.000
500142	Bulletproof Supplier Management Program	Jeff Kasoff	RECORDED	2014-12-12	\$189.000
500143	Complaint Handling in Compliance with FDA and ISO Regulations	Jeff Kasoff	RECORDED	2015-02-09	\$189.000
500144	A CAPA Primer - Elements of a CAPA Program	Jeff Kasoff	RECORDED	2015-01-22	\$189.000
500145	FDA Inspections – Do's and Don'ts	Jeff Kasoff	RECORDED	2015-02-19	\$189.000
500146	The Strategy of Experimentation	Lynne Hare	RECORDED	2014-12-09	\$149.000
500147	Chemical Safety for Sanitation Workers	Donald Jones	RECORDED	2014-12-11	\$149.000
500148	What's Up? (and Down): Outlook for Construction Segments, Materials and Labor	Kenneth D. Simonson	RECORDED	2014-12-18	\$149.000
500149	Operational Risk - Beyond Regulatory Constraints	Fred Vacelet	RECORDED	2014-12-09	\$149.000
500150	ATC Radar Approach Control (TRACON)	Stephen Alvania	RECORDED	2015-01-07	\$149.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
500151	Air Route Traffic Control Center (ARTCC) Operations	Stephen Alvania	RECORDED	2015-02-04	\$149.000
500152	Evaporation Ponds: a Permitting Labyrinthine	Erin Hallenberg	RECORDED	2015-01-20	\$149.000
500153	Controlled Goods in Canada - A primer	Anthony Goode	RECORDED	2014-12-11	\$149.000
500154	Industrial Technical Benefits- Canada's New Approach to Offsets for Defence Procurement	Anthony Goode	RECORDED	2015-01-13	\$149.000
500157	Environmental Cleaning	Donald Jones	RECORDED	2015-02-11	\$149.000
500158	Building a Sustainable Vendor Qualification Program	Jonathan M. Lewis	RECORDED	2015-01-15	\$189.000
500160	FDA Inspections – Do's and Don'ts	Jonathan M. Lewis	RECORDED	2015-03-12	\$189.000
500159	Building a Validation Program From Top to Bottom	Jonathan M. Lewis	RECORDED	2015-02-12	\$189.000
500165	Drug Safety Risk Management Planning	Dr. Michael Forstner	RECORDED	2015-02-11	\$189.000
500168	Establishment of Quality Systems	Louis Angelucci	RECORDED	2015-05-21	\$189.000
500169	Introduction to Risk Assessment	Louis Angelucci	RECORDED	2015-03-30	\$189.000
500171	Foreign Corrupt Practices Act - Audits Role	Denise Cicchella	RECORDED	2015-02-12	\$149.000
500172	Project Management - Auditing Problem Projects	Stuart Gardner	RECORDED	2015-01-21	\$149.000
500173	Auditing Third Party Agreements - Common Pitfalls from IT to Construction	Stuart Gardner	RECORDED	2015-02-19	\$149.000
500174	Construction is Building, but Where will the Workers Come From?	Kenneth D. Simonson	RECORDED	2015-02-26	\$149.000
500175	Complaint Handling for Medical Device Manufacturers	Leo Lagrotte	RECORDED	2015-01-29	\$189.000
500177	Laboratory Accreditation: Getting there is just the beginning	Michael Brodsky	RECORDED	2015-01-13	\$189.000
500178	An Environmental Microbiologist's View on Estimation of Uncertainty of Measurement	Michael Brodsky	RECORDED	2015-02-03	\$189.000
500179	Verification or Validation of Methods in Food Microbiology	Michael Brodsky	RECORDED	2015-02-26	\$149.000
500181	Gluten - Free Product: How to Prove & Display It Correctly	Rotimi Toki	RECORDED	2015-01-15	\$149.000
500182	Proven Cleaning and Sanitation Techniques for Food Processors	Rotimi Toki	RECORDED	2015-02-12	\$149.000
500183	Ready- to-eat Foods: Pathogens of Concern and Intervention Controls	Rotimi Toki	RECORDED	2015-03-19	\$149.000
500184	FDA Regulation of Combination Products	Thomas E. Colonna	RECORDED	2015-01-22	\$189.000
500185	FDA Regulation of Dietary Supplements	Thomas E. Colonna	RECORDED	2015-02-19	\$189.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
500187	Better Alternatives to Sampling Plans	John N. Zorich	RECORDED	2015-03-11	\$189.000
500188	Introduction to SPC (Statistical Process Control)	John N. Zorich	RECORDED	2015-02-03	\$189.000
500189	Statistical Analysis of Gages	John N. Zorich	RECORDED	2015-02-18	\$189.000
500190	Confidence-Reliability Calculations and Statistically Valid Sample Sizes	John N. Zorich	RECORDED	2015-03-03	\$189.000
500191	Normality Tests and Transformations	John N. Zorich	RECORDED	2015-03-17	\$189.000
500192	Medical Devices and Radiation-Emitting Products: FDA Requirements	Leo Lagrotte	RECORDED	2015-02-19	\$189.000
500193	What does FDA require for Medical Devices Manufacturers to Do When Filing an Adverse Event Report (MDR)	Leo Lagrotte	RECORDED	2015-03-12	\$189.000
500195	Gold in Plant Tailings - Prevention and Recovery	Hannes Wagner	RECORDED	2015-02-26	\$149.000
500196	Developing a Cyber Incident Response Program	Ms. Michael Redmond	RECORDED	2015-02-19	\$149.000
500197	How to Analyze Financial Statements	Miles Hutchinson	RECORDED	2015-02-18	\$149.000
500198	Mastering the Power of Sensitivity Tools in Financial Modeling	Miles Hutchinson	RECORDED	2015-03-20	\$149.000
500199	How to Design and Implement Outstanding KPI Performance Dashboards	Miles Hutchinson	RECORDED	2015-03-31	\$149.000
500200	Airport Traffic Control Tower (ATCT)	Stephen Alvania	RECORDED	2015-02-25	\$149.000
500201	Terminal Radar Approach Control (TRACON)	Stephen Alvania	RECORDED	2015-03-11	\$149.000
500202	Air Route Traffic Control Center (ARTCC) Operations	Stephen Alvania	RECORDED	2015-04-08	\$149.000
500203	Traffic Flow Management (TFM) Operations-End to End Flight Scenario	Stephen Alvania	RECORDED	2015-05-06	\$149.000
500204	Pharma Contract Manufacturing: Managing Quality and Technical Agreements	Joseph Habarta	RECORDED	2015-04-24	\$189.000
500205	Using Statistics to Determine Sample Size	Steven Walfish	RECORDED	2015-02-19	\$189.000
500206	Complaint Handling and Management: From Receipt to Trending	David R Dills	RECORDED	2015-02-24	\$189.000
500207	Bullet Proof 510(k) – Latest FDA Changes to the Process	David R Dills	RECORDED	2015-03-19	\$189.000
500208	Combination Products: FDA's Final Rule for GMP Requirements and Introduction and Expectations for "Combo" Products	David R Dills	RECORDED	2015-03-24	\$189.000
500209	Conducting Successful Product Complaint Investigations	David R Dills	RECORDED	2015-04-14	\$189.000
500213	Biological Facility Design for Compliance	Herman Bozenhardt	RECORDED	2015-03-26	\$189.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
500212	Establish and Maintain an Effective Supplier Qualification Program	David R Dills	RECORDED	2015-05-28	\$139.000
500219	Reducing Variation in Manufacturing Processes	Lynne Hare	RECORDED	2015-03-17	\$149.000
500224	A second look at 510(k) changes	Anna Longwell	RECORDED	2015-04-07	\$189.000
500225	The New Clinical Trial Regulation	Adriaan Fruijtjer	RECORDED	2015-02-24	\$189.000
500229	Lineament analysis- The Modern Way to Look for Ore Deposits	Ricardo Valls	RECORDED	2015-03-10	\$189.000
500232	"Zero Defects" and the cGMPs - Pros and Cons	John E Lincoln	RECORDED	2015-02-24	\$189.000
500233	Avoid Warning Letters in View of the U.S. FDA's Stated Goal	John E Lincoln	RECORDED	2015-03-24	\$189.000
500236	Preventing Recalls & Product Liability in New Product Development	Randy Goodden	RECORDED	2015-03-17	\$149.000
500237	CAPA, Failure Investigation and Root Cause Analysis to Meet FDA Expectations	John E Lincoln	RECORDED	2015-04-07	\$189.000
500239	Design Controls - Requirements for Medical Device Developers	John E Lincoln	RECORDED	2015-04-29	\$189.000
500246	DHF, DMR, DHR, Technical File and Design Dossier - Key Requirements and Future Directions	John E Lincoln	RECORDED	2015-05-12	\$189.000
500250	Complying with FATCA - the Foreign Accounts Tax Compliance Act	Miles Hutchinson	RECORDED	2015-04-07	\$149.000
500251	Introduction to Sales and Use Taxation	Miles Hutchinson	RECORDED	2015-04-22	\$149.000
500252	1099 and W-9 Update - Complying with IRS Information Reporting	Miles Hutchinson	RECORDED	2015-09-15	\$149.000
500248	Developing the Master V&V Plan to Meet U.S. FDA, ISO 13485 and 14971 Requirements	John E Lincoln	RECORDED	2015-06-10	\$147.000
500249	Device Changes, FDA Changes, and the 510(k)	John E Lincoln	RECORDED	2015-06-24	\$189.000
500253	Introduction to Payroll Law	Miles Hutchinson	RECORDED	2015-06-17	\$138.000
500254	TIN Matching to Reduce Your B-Notices and Eliminate Proposed Penalties	Miles Hutchinson	RECORDED	2015-07-21	\$149.000
500255	Analytical Method Validation	Jerry Lanese	RECORDED	2015-03-25	\$189.000
500256	Understanding Next Gen (FAA New Technology)	Roger Nakata	RECORDED	2015-03-27	\$149.000
500258	Process Validation - Statistical Process Control	Jerry Dalfors	RECORDED	2015-03-25	\$189.000
500260	Lyophilization Technology	Jerry Dalfors	RECORDED	2015-04-29	\$189.000
500262	Impulse Sealing: Trials and Tribulations	Jan Gates	RECORDED	2015-03-19	\$189.000
500263	Medical Device Product Development Process	Karl Leinsing	RECORDED	2015-03-26	\$189.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
500264	Standardizing Transportation Procedures to Control Food Safety and Quality	John Ryan	RECORDED	2015-03-05	\$149.000
500267	The Social Cost of Carbon: The EPA's Stealth Imposition of a Carbon Tax That Will Make Everything More Expensive and Less Efficient	Roger Bezdek	RECORDED	2015-03-18	\$149.000
500268	Regulatory Aspects of Advanced Therapy Medicinal Products in the EU	Adriaan Fruijtjer	RECORDED	2015-05-19	\$189.000
500269	Scientific Advice in the EU	Adriaan Fruijtjer	RECORDED	2015-05-27	\$189.000
500270	Product Information in the EU	Adriaan Fruijtjer	RECORDED	2015-06-09	\$121.000
500272	cGMPs in the Quality Control Laboratory	Jerry Lanese	RECORDED	2015-04-09	\$189.000
500274	HACCP in Foodservice Establishment - Practical Design & Implementation	Rotimi Toki	RECORDED	2015-04-23	\$149.000
500277	FSMA Impact to the Transportation of Perishables	John Ryan	RECORDED	2015-04-28	\$149.000
500279	Understanding, Calculating, and Using Statistical Power	John N. Zorich	RECORDED	2015-04-08	\$189.000
500280	Process Capability Analysis of Extremely Non-Normal Data	John N. Zorich	RECORDED	2015-04-29	\$189.000
500281	Process Capability Analysis by means of Confidence - Reliability Calculations	John N. Zorich	RECORDED	2015-06-25	\$136.000
500282	Metrology: Statistical Analysis of Measurement Uncertainty	John N. Zorich	RECORDED	2015-05-13	\$189.000
500284	Foreign Bodies in Foods - Effective Techniques for Prevention, Control and Detection	Rotimi Toki	RECORDED	2015-05-05	\$149.000
500285	Food Allergen Programs: Management of Allergen Cross-Contamination and Validation of Cleaning Procedures to Ensure Effective Removal	Rotimi Toki	RECORDED	2015-05-27	\$189.000
500286	Proven Cleaning and Sanitation Techniques for Food Processors	Rotimi Toki	RECORDED	2015-06-11	\$149.000
500287	Gluten - Free Product: How to Prove & Display It Correctly	Rotimi Toki	RECORDED	2015-06-30	\$149.000
500288	Reclaiming Large Excel Spreadsheets	David Ringstrom	RECORDED	2015-04-22	\$149.000
500289	Spreadsheet-Based Internal Controls	David Ringstrom	RECORDED	2015-05-07	\$149.000
500290	Building a Validation Program From Top to Bottom	Jonathan M. Lewis	RECORDED	2015-04-10	\$189.000
500291	Building a Sustainable Vendor Qualification Program	Jonathan M. Lewis	RECORDED	2015-05-15	\$189.000
500292	FDA Inspections - Do's and Don'ts	Jonathan M. Lewis	RECORDED	2015-06-01	\$189.000
500295	IFRS 6 Oil, Gas, Mining and other Extractive Industries	Mike Morley	RECORDED	2015-04-28	\$149.000
500296	Does Anyone Care About SOX Anymore?	Mike Morley	RECORDED	2015-05-21	\$149.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
500297	Food Safety - Food Defense	Jason Teliszczak	RECORDED	2015-04-09	\$149.000
500298	Food Safety - Food Certifications - What Certification is Best for Your Company?	Jason Teliszczak	RECORDED	2015-04-29	\$149.000
500299	Construction - OSHA Construction Basics, for any Jobsite	Jason Teliszczak	RECORDED	2015-05-20	\$127.000
500301	SOX: Internal Controls for Accounts Payable	Mike Morley	RECORDED	2015-06-09	\$124.000
500302	Software FMEA for Medical Devices	Dev Raheja	RECORDED	2015-04-01	\$189.000
500303	Designing Medical Devices for Long Life at Lower Costs	Dev Raheja	RECORDED	2015-05-05	\$128.000
500304	Cyber Security Mitigation and Response Risk Assessment	Ms. Michael Redmond	RECORDED	2015-04-01	\$149.000
500305	BC ISO 22301: Business Continuity Planning	Ms. Michael Redmond	RECORDED	2015-04-20	\$149.000
500310	The Impact of Basel III on Export Finance	Buddy Baker	RECORDED	2015-04-08	\$149.000
500316	Contract Killer Clauses in Construction: And How to Neutralize Them	Brian Perlberg	RECORDED	2015-05-11	\$149.000
500317	Medical Devices - ISO 13485 – Do You Really Know What You Need To?	Jason Teliszczak	RECORDED	2015-06-10	\$189.000
500320	Your Social Media Marketing is Under FDA's Microscope	Casper Uldriks	RECORDED	2015-05-05	\$189.000
500324	Emerging Issues in Food Safety	Michael Brodsky	RECORDED	2015-06-15	\$149.000
500328	Manage Your FDA Inspection Before It Happens	Casper Uldriks	RECORDED	2015-06-16	\$189.000
500330	Aerospace - AS9100 vs ISO 9001	Jason Teliszczak	RECORDED	2015-07-16	\$149.000
500332	Exporting your Dietary Supplements to Canada	Karen Friedman	RECORDED	2015-04-30	\$149.000
500333	Facing the challenges of Canadian Consumer Health Product Registration	Karen Friedman	RECORDED	2015-05-28	\$149.000
500334	Designing Full Cyber Security Incident Response Team (CSIRT) Training Program as well as Table Top and Simulation Testing	Ms. Michael Redmond	RECORDED	2015-05-04	\$149.000
500339	Business Continuity and Disaster Recovery Management	Ms. Michael Redmond	RECORDED	2015-06-29	\$138.000
500340	Resiliency Individual, Community, Business	Ms. Michael Redmond	RECORDED	2015-07-29	\$149.000
500343	Statistical Methods for Process Validation	Heath Rushing	RECORDED	2015-05-22	\$189.000
500344	Quality by Design: Establishing a Systematic Approach to Pharmaceutical Development	Heath Rushing	RECORDED	2015-06-08	\$189.000
500345	Introduction to Industrial Microbiology	Neal Machtiger	RECORDED	2015-05-07	\$189.000

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500346	FDA Regulation of Medical Device Software	Thomas E. Colonna	RECORDED	2015-05-28	\$189.000
500348	Fundamentals on FDA Regulations of Bioresearch Monitoring	Thomas E. Colonna	RECORDED	2015-07-30	\$189.000
500350	Clinical Trials in US, Europe and Canada	Anne Tomalin	RECORDED	2015-06-22	\$127.000
500351	Conducting Observational Studies in US, Canada and Europe	Anne Tomalin	RECORDED	2015-06-08	\$126.000
500352	Clinical Trial Applications in Canada, and Comparison to the US and Europe	Anne Tomalin	RECORDED	2015-05-18	\$127.000
500354	Current Status of Biosimilars in US, Europe and Canada	Anne Tomalin	RECORDED	2015-07-28	\$189.000
500355	Biocompatibility Testing of Medical Devices in the US and Use of ISO-10993	Elisa Harvey	RECORDED	2015-05-11	\$189.000
500357	Using the Pre-Submission Process to Your Best Advantage	Elisa Harvey	RECORDED	2015-06-08	\$131.000
500358	Orphan Devices: Humanitarian Device Exemptions and Humanitarian Use Designations	Elisa Harvey	RECORDED	2015-06-18	\$132.000
500359	Outlook for Multifamily Housing	Bernard Markstein	RECORDED	2015-08-19	\$127.000
500361	Outlook for Single-Family Housing	Bernard Markstein	RECORDED	2015-06-24	\$139.000
500367	Airport Traffic Control Tower (ATCT)	Stephen Alvania	RECORDED	2015-05-27	\$141.000
500368	Terminal Radar Approach Control (TRACON)	Stephen Alvania	RECORDED	2015-06-23	\$143.000
500369	Air Route Traffic Control Center (ARTCC) Operations	Stephen Alvania	RECORDED	2015-07-08	\$135.000
500373	Pest Control in Food Processing and Foodservice	Norman G. Marriott	RECORDED	2015-07-14	\$124.000
500376	Canadian Establishment Licenses: Drugs, Supplements and Medical Devices	Karen Friedman	RECORDED	2015-07-13	\$143.000
500377	The Bayh-Dole Act and Its Implementing Regulations & Rule-Making	Ronald Kudla	RECORDED	2015-06-03	\$131.000
500380	Drug Safety Risk Management Planning	Dr. Michael Forstner	RECORDED	2015-05-26	\$146.000
500382	Concepts of Hazard Analysis and Risk Assessment	Michael Allocco	RECORDED	2015-05-29	\$133.000
500384	Expedite Excel with Hidden Shortcuts	David Ringstrom	RECORDED	2015-05-26	\$145.000
500385	Leveraging Linked Workbooks in Excel	David Ringstrom	RECORDED	2015-06-22	\$144.000
500386	Transcend VLOOKUP in Microsoft Excel	David Ringstrom	RECORDED	2015-08-24	\$147.000
500392	Complaint Handling and Management: From Receipt to Trending	David R Dills	RECORDED	2015-07-23	\$155.000
500394	Bullet Proof 510(k) – Latest FDA Changes to the Process	David R Dills	RECORDED	2015-09-01	\$158.000

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500395	Draft Guidance for Device Industry and FDA - Postmarket Surveillance	David R Dills	RECORDED	2015-09-23	\$154.000
500396	Combination Products: FDA's Final Rule for GMP Requirements and Introduction and Expectations for "Combo" Products	David R Dills	RECORDED	2015-10-20	\$190.000
500397	Auditing Computer Systems for FDA and International Compliance	Dr. Ludwig Huber	RECORDED	2015-06-18	\$178.000
500398	Calibration and Qualification in Analytical Laboratories	Dr. Ludwig Huber	RECORDED	2015-07-08	\$183.000
500399	Postmarketing Vigilance Reporting For Medical Device Manufacturers	David R Dills	RECORDED	2015-10-29	\$190.000
500400	Conducting Successful Product Complaint Investigations	David R Dills	RECORDED	2015-11-18	\$190.000
500402	Conducting Safety Reviews for Systems, Products, Processes, facilities, and Procedures	Michael Allocco	RECORDED	2015-07-24	\$132.000
500405	Efficient and Effective FDA and ISO Management Reviews	Betty Lane	RECORDED	2015-08-04	\$152.000
500406	Corrective Actions - Current Expectation of ISO 13485 and FDA Auditors	Betty Lane	RECORDED	2015-09-22	\$148.000
500407	Effective Root Cause Analysis: The key to an Effective Corrective Actions System	Betty Lane	RECORDED	2015-10-09	\$147.000
500409	Good Documentation Practices for GxP Compliance	Alla Teresh	RECORDED	2015-07-30	\$144.000
500410	FDA Export Certificates for Medical Devices	Larry Spears	RECORDED	2015-07-22	\$146.000
500393	Importing and Exporting Medical Devices: A Primer on Regulatory Strategy and Requirements	David R Dills	RECORDED	2015-09-08	\$148.000
500412	Nonresident Alien W-8 & 1042-S Compliance Update	Miles Hutchinson	RECORDED	2015-06-30	\$142.000
500413	How to Design and Implement Outstanding KPI Performance Dashboards	Miles Hutchinson	RECORDED	2015-08-18	\$146.000
500414	Introduction to Sales and Use Taxation	Miles Hutchinson	RECORDED	2015-09-23	\$145.000
500415	Financial Statement Preparation and Analysis	Miles Hutchinson	RECORDED	2015-11-18	\$190.000
500417	FDA Regulation of In Vitro Diagnostics	Thomas E. Colonna	RECORDED	2015-06-09	\$134.000
500418	FDA Regulation of Combination Products	Thomas E. Colonna	RECORDED	2015-07-07	\$138.000
500419	FDA Regulation of Dietary Supplements	Thomas E. Colonna	RECORDED	2015-08-13	\$137.000
500424	Medical Device Registration & Listing, and Inspection Follow-up Activities	Larry Spears	RECORDED	2015-08-10	\$147.000
500422	Good Clinical Practice (GCP)	Omid Khodai	RECORDED	2015-07-06	\$146.000
500423	How to Survive an FDA Inspection	Omid Khodai	RECORDED	2015-08-03	\$148.000

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500425	Regulation of the Practice of Geology in the United States	Mark Brengelman	RECORDED	2015-07-07	\$131.000
500428	Medical Device Product Development Process	Karl Leinsing	RECORDED	2015-06-24	\$147.000
500429	Excel Spreadsheet Validation for FDA 21 CFR Part 11	Angela Bazigos	RECORDED	2015-07-07	\$157.000
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
500433	FDA's enforcement of 21 CFR part 11 compliance	Angela Bazigos	RECORDED	2015-09-15	\$167.000
500434	Introduction to Program-Project Management	Drex Rutledge	RECORDED	2015-08-06	\$124.000
500438	Anti-Money Laundering (AML) -Connecting the Dots	James Bone	RECORDED	2015-08-07	\$148.000
500439	Why Good Risk Management is so Elusive	James Bone	RECORDED	2015-08-20	\$152.000
500440	Medical Device Recalls	Larry Spears	RECORDED	2015-08-26	\$148.000
500441	Auditing Enterprise Risk Management	Denise Cicchella	RECORDED	2015-07-20	\$137.000
500442	Foreign Corrupt Practices Act - Audits Role	Denise Cicchella	RECORDED	2015-08-20	\$141.000
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
500445	Product Risk Management - ISO 14971:2007	John E Lincoln	RECORDED	2015-09-30	\$137.000
500446	Setting Up and Running a Tougher Supplier Audit Program	John E Lincoln	RECORDED	2015-10-21	\$190.000
500448	CAPA, Failure Investigation and Root Cause Analysis to Meet FDA Expectations	John E Lincoln	RECORDED	2015-12-02	\$190.000
500449	Electronic Informed Consent for Clinical Trials: Why, What and How	Susan Brink	RECORDED	2015-07-21	\$144.000
500450	Traceability and Recall Through Food Processes	John Ryan	RECORDED	2015-07-22	\$128.000
500451	Food Import Detections and Seizures: The Current and Future Regulatory Environment	John Ryan	RECORDED	2015-08-10	\$126.000
500455	Software FMEA for Medical Devices	Dev Raheja	RECORDED	2015-08-28	\$146.000
500462	Environmental Cleaning	Donald Jones	RECORDED	2015-08-18	\$127.000
500463	Sparking Innovation at Your Company	Shari Storm		2015-07-27	

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500464	Capturing the Most Powerful Market - Moms	Shari Storm	RECORDED	2015-08-31	\$132.000
500468	Owner's Guide to Understanding Financial Statements	Richard Melancon	RECORDED	2015-09-03	\$138.000
500469	Building a Sustainable Vendor Qualification Program	Jonathan M. Lewis	RECORDED	2015-08-21	\$158.000
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
500474	Reclaiming Large Excel Spreadsheets	David Ringstrom	RECORDED	2015-08-03	\$152.000
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
500504	Project Management for Internal Auditors	Danny M. Goldberg	RECORDED	2015-08-27	\$144.000
500511	Operational Risk for Financial Institutions - Beyond Regulatory Constraints	Fred Vacelet	RECORDED	2015-09-01	\$143.000

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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
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

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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
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
500606	Credit card surcharging What are The Stipulations for Compliance?	Ray Graber	RECORDED	2016-01-27	\$190.000
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

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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
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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
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

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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
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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
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 R Dills	RECORDED	2016-04-05	\$190.000
500807	Complaint Handling and Management: From Receipt to Trending	David R Dills	RECORDED	2016-05-04	\$390.000
500808	Construct and Manage the Technical File and Design Dossier	David R 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 R Dills	RECORDED	2016-05-26	\$390.000
500826	Importing and Exporting Medical Devices: A Primer on Regulatory Strategy and Requirements	David R Dills	RECORDED	2016-06-22	\$390.000
500827	Update on Unique Device Identifier for Device Manufacturers	David R Dills	RECORDED	2016-06-30	\$390.000
500831	Managing Your Medical Device Reporting (MDR) Program for Compliance Success	David R Dills	RECORDED	2016-08-24	\$390.000
500836	Postmarketing Vigilance Reporting For Medical Device Manufacturers	David R Dills	RECORDED	2016-11-03	\$390.000

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500837	Maintaining an Effective CAPA Program and Using Risk Assessment Tools: Current Trends	David R 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
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

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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
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

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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
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

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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
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 R 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:	Robert Michalik	RECORDED	2017-01-25	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
	Essential knowledge needed to Manage Drug&Device Combination Product Project				
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
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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
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 R Dills	RECORDED	2017-08-09	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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
501607	FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	RECORDED	2017-11-13	\$390.000
501608	Understanding the USP Chapter 1224 for Transfer of Analytical Methods	Dr. Ludwig Huber	RECORDED	2017-11-15	\$390.000
501609	New FDA, EMA and USP Guidelines for Transfer of Analytical Methods	Dr. Ludwig Huber	RECORDED	2017-11-30	\$390.000
501611	FDA's New Import Program for 2017	Casper Uldriks	RECORDED	2017-12-05	\$390.000
501612	Validation and Use of Excel® Spreadsheets in FDA Regulated Environments	Dr. Ludwig Huber	RECORDED	2017-12-14	\$390.000
501617	FDA Regulation of Dietary Supplements	Thomas E. Colonna	RECORDED	2017-12-12	\$390.000
501619	21 CFR Part 11 - Compliance for Electronic Records and Signatures	Edwin Waldbusser	RECORDED	2018-02-02	\$390.000
501620	Software Validation for the New FDA Inspections	Edwin Waldbusser	RECORDED	2018-01-30	\$390.000
501623	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	RECORDED	2017-12-12	\$390.000
501626	Drug Master Files: Understanding and Meeting your Regulatory and Processing Responsibilities	Robert J. Russell	RECORDED	2017-11-06	\$390.000
501627	Japan - Regulatory Filing Requirements and Compliance Processes for Life Sciences	Robert J. Russell	RECORDED	2017-12-08	\$390.000
501629	Complaint Handling in Compliance with FDA and ISO Regulations	Jeff Kasoff	RECORDED	2017-10-27	\$390.000
501633	Handling OOS Test Results and Completing Robust Investigations	Danielle DeLucy	RECORDED	2017-11-28	\$390.000
501634	How FDA Trains its Investigators to Review CAPA and What you Should do to Prepare	Jeff Kasoff	RECORDED	2017-12-04	\$390.000
501635	Time-Saving Excel Tips, Tricks and 100 Shortcuts	Dennis Taylor	RECORDED	2017-11-21	\$390.000
501636	Mastering Excel 2013 Pivot Tables	Dennis Taylor	RECORDED	2017-12-12	\$390.000
501638	The Value of a Human Factors Program	Thomas Bento	RECORDED	2018-01-12	\$390.000
501641	Effective Root Cause Analysis: The Key to an Effective Corrective Actions System	Betty Lane	RECORDED	2017-12-06	\$390.000
501645	Auditing Analytical Laboratories for FDA Compliance	Steven S. Kuwahara	RECORDED	2017-12-11	\$390.000
501647	Complaint Handling and Management: From Receipt to Trending	David R Dills	RECORDED	2017-11-28	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501648	CAPA, Failure Investigation and Root Cause Analysis to Meet FDA Expectations	John E Lincoln	RECORDED	2018-01-17	\$390.000
501651	Design History Files (DHF), Device Master Records (DMR), Device History Records (DHR), Technical Files , Design Dossiers - The Requirements	John E Lincoln	RECORDED	2018-01-31	\$390.000
501652	Excel Spreadsheet Validation for FDA 21 CFR Part 11	Angela Bazigos	RECORDED	2017-12-06	\$390.000
501653	Key Factors to Write an Effective Standard Operating Procedure (SOP) and Work Instructions (WIs)	Angela Bazigos	RECORDED	2018-04-02	\$390.000
501654	FDA's Enforcement of 21 CFR Part 11 Compliance	Angela Bazigos	RECORDED	2017-12-05	\$390.000
501655	2011 FDA Guideline on Process Validation	Louis Angelucci	RECORDED	2017-12-04	\$390.000
501657	Modeling and Optimizing Process Behavior Using Design of Experiments	Steven Wachs	RECORDED	2018-01-05	\$390.000
501658	GMP Expectations for Products Used in Early Phase IND Studies	Steven S. Kuwahara	RECORDED	2018-01-25	\$390.000
501659	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2017-12-14	\$390.000
501660	Laboratory-Developed Tests: Why does FDA think they Can Regulate them, and why do Others think they Cannot	Anna Longwell	RECORDED	2017-12-08	\$390.000
501661	Latin America Regulatory Compliance: Current Challenges in Argentina, Brazil and Mexico	Robert J. Russell	RECORDED	2017-11-28	\$390.000
501662	Controlling Human Error in the Manufacturing Floor	Ginette Collazo	RECORDED	2017-12-13	\$390.000
501668	European Data Protection Regulation - 2018 Implementation	Angela Bazigos	RECORDED	2018-03-29	\$390.000
501669	Master Excel: Introduction to Pivot Tables - Part 1	David Ringstrom	RECORDED	2018-02-05	\$390.000
501671	Death by CAPA - Does your Company have the Symptoms?	Susanne Manz	RECORDED	2018-01-18	\$390.000
501674	Non-Conforming Material and Failure Investigation	Susanne Manz	RECORDED	2018-04-17	\$390.000
501680	Batch Record Review and Product Release	Danielle DeLucy	RECORDED	2018-01-18	\$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	RECORDED	2018-04-18	\$390.000
501684	Dietary Supplements CGMPs - 21 CFR 111 Compliance	John E Lincoln	RECORDED	2018-05-01	\$390.000
501685	The Dietary Supplement cGMP Rule (21 CFR part 111)	Thomas E. Colonna	RECORDED	2018-01-30	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501686	3-Hour Virtual Seminar on Off label Promotion of Drugs and Medical Devices - FDA's Latest	Angela Bazigos	RECORDED	2018-06-08	\$540.000
501688	The New ISO 13485:2016 and Comparison with 21CFR820 - How to Comply with both in the Same Organization	Eyal Lerner	RECORDED	2018-02-23	\$390.000
501690	Complaint Handling and Management: From Receipt to Trending	David R Dills	RECORDED	2018-01-26	\$390.000
501691	Excel Spreadsheet Validation for FDA 21 CFR Part 11	Angela Bazigos	RECORDED	2018-01-26	\$390.000
501692	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2018-01-26	\$390.000
501693	FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	RECORDED	2018-01-09	\$390.000
501694	3-hour Virtual Seminar on Analytical Method Validation Process	Angela Bazigos	RECORDED	2018-02-26	\$540.000
501695	How FDA Trains its Investigators to Review CAPA and What you Should do to Prepare	Jeff Kasoff	RECORDED	2018-02-02	\$390.000
501696	Mastering Excel 2013 Pivot Tables	Dennis Taylor	RECORDED	2018-01-18	\$390.000
501697	Controlling Human Error in the Manufacturing Floor	Ginette Collazo	RECORDED	2018-02-27	\$390.000
501698	Tools for Human Error Reduction	Ginette Collazo	RECORDED	2018-03-20	\$390.000
501699	FDA's Enforcement of 21 CFR Part 11 Compliance	Angela Bazigos	RECORDED	2018-02-05	\$390.000
501700	Validation of Analytical Methods and Procedures	Dr. Ludwig Huber	RECORDED	2018-01-18	\$390.000
501701	Secrets to Writing Effective SOPs for Medical Device QMS	Susanne Manz	RECORDED	2018-01-26	\$390.000
501702	Mastering Excel Formulas and Functions	Mike Thomas	RECORDED	2018-01-26	\$390.000
501703	FDA's 21 CFR 11 Add-On Inspections - Recent Updates	Angela Bazigos	RECORDED	2018-03-07	\$390.000
501704	Water System Mythology: Common False Beliefs for Microbial Control and Monitoring	Teri C. Soli	RECORDED	2018-01-26	\$390.000
501707	FDA's New Import Program for 2018	Casper Uldriks	RECORDED	2018-01-30	\$390.000
501708	Validation and Use of Excel® Spreadsheets in FDA Regulated Environments	Dr. Ludwig Huber	RECORDED	2018-01-30	\$390.000
501709	Key Factors to Write an Effective Standard Operating Procedure (SOP) and Work Instructions (WIs)	Angela Bazigos	RECORDED	2018-03-14	\$390.000
501710	How to Audit Against ICH GCP 2 Addendum to Ensure Compliance	Laura Brown	RECORDED	2018-01-08	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501722	GDPR - European Privacy Regulation what and how	Derk Yntema	RECORDED	2018-01-24	\$390.000
501724	Analytical Method Validation Under Good Laboratory Practices - GLPs	John C. Fetzer	RECORDED	2018-02-05	\$390.000
501730	Understanding Attribute Acceptance Sampling Plans Including the Z1.4 and c=0 Plans	Dan OLeary	RECORDED	2018-01-17	\$390.000
501733	Accelerated Aging Techniques for Medical Device Products	Karl J Hemmerich	RECORDED	2018-01-22	\$390.000
501734	Laboratory Investigation of Out-of-Specification Results	Jerry Lanese	RECORDED	2018-01-23	\$390.000
501736	Analytical Method Validation	Jerry Lanese	RECORDED	2018-03-06	\$390.000
501737	GLPs: How are they Associated with GMPs and SOPs	Joy McElroy	RECORDED	2018-01-23	\$390.000
501739	Time-Saving Excel Tips, Tricks and 100 Shortcuts	Dennis Taylor	RECORDED	2018-02-09	\$390.000
501740	Root Cause Failure Analysis Closed Loop Corrective Action	Michael Abitz	RECORDED	2018-01-25	\$390.000
501745	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	RECORDED	2018-01-05	\$390.000
501752	Software Verification & Validation	Lena Cordie	RECORDED	2018-03-14	\$390.000
501757	FDA Inspections - Do's and Don'ts	Jonathan M. Lewis	RECORDED	2018-02-21	\$390.000
501761	OneNote - Bringing Order to your Digital Chaos	Mike Thomas	RECORDED	2018-02-12	\$390.000
501763	Get Organised with Microsoft Outlook	Mike Thomas	RECORDED	2018-05-14	\$390.000
501767	Combination Products Registration	Salma Michor	RECORDED	2018-02-09	\$390.000
501769	The New EU Medical Device Regulation	Salma Michor	RECORDED	2018-03-12	\$390.000
501776	2011 FDA Guideline on Process Validation	Louis Angelucci	RECORDED	2018-02-15	\$390.000
501778	Understanding Proper Application of Risk Assessment	Louis Angelucci	RECORDED	2018-03-23	\$390.000
501782	Good Documentation Practices to Support FDA Computer System Validation	Carolyn Troiano	RECORDED	2018-05-01	\$390.000
501784	GCP-GLP-GMP : Comparison and Understanding of FDA's 3 Major Regulations	Dr. David Lim	RECORDED	2018-02-22	\$390.000
501787	Understanding the USP Chapter 1224 for Transfer of Analytical Methods	Dr. Ludwig Huber	RECORDED	2018-02-22	\$390.000
501788	FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	RECORDED	2018-03-07	\$390.000
501792	European Data Protection Regulation - 2018 Implementation	Angela Bazigos	RECORDED	2018-02-21	\$390.000
501796	Packaging and Labeling in the Pharmaceutical Supply Chain	Michael Esposito	RECORDED	2018-04-20	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501797	Onboarding Employees in a GMP Environment: Best Practices for Foundational Employee Success	Michael Esposito	RECORDED	2018-04-30	\$390.000
501800	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2018-02-26	\$390.000
501801	Mastering Excel 2013 Pivot Tables	Dennis Taylor	RECORDED	2018-02-23	\$390.000
501802	Validation of Analytical Methods and Procedures	Dr. Ludwig Huber	RECORDED	2018-02-20	\$390.000
501804	Mastering Excel Formulas and Functions	Mike Thomas	RECORDED	2018-03-02	\$390.000
501805	Water System Mythology: Common False Beliefs for Microbial Control and Monitoring	Teri C. Soli	RECORDED	2018-02-23	\$390.000
501806	GDPR - European Privacy Regulation what and how	Derk Yntema	RECORDED	2018-02-26	\$390.000
501807	FDA Regulation of Dietary Supplements	Thomas E. Colonna	RECORDED	2018-03-02	\$390.000
501808	Death By CAPA - Does your CAPA Program Need a CAPA?	Susanne Manz	RECORDED	2018-02-26	\$390.000
501810	Japan - Regulatory Filing Requirements and Compliance Processes for Life Sciences	Robert J. Russell	RECORDED	2018-03-20	\$390.000
501812	Auditing Analytical Laboratories for FDA Compliance	Steven S. Kuwahara	RECORDED	2018-03-08	\$390.000
501813	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	RECORDED	2018-03-27	\$390.000
501814	Accelerated Aging Techniques for Medical Device Products	Karl J Hemmerich	RECORDED	2018-03-14	\$390.000
501815	Latin America Regulatory Compliance: Current Challenges in Argentina, Brazil and Mexico	Robert J. Russell	RECORDED	2018-03-27	\$390.000
501817	Medical Device Software Validation Meeting FDA Regulations	Edwin Waldbusser	RECORDED	2018-06-14	\$390.000
501818	How to Conduct a Human Factors - Usability Validation Test following ISO 62366 and the 2016 FDA Guidance	Edwin Waldbusser	RECORDED	2018-04-18	\$390.000
501820	Medical Device Single Audit Program (MDSAP) Preparation	Shep Bentley	RECORDED	2018-04-06	\$390.000
501829	Best Practices to Prepare for and Manage FDA Inspections	Vanessa Lopez	RECORDED	2018-03-29	\$390.000
501830	Verification vs Validation-Product, Process or Equipment and QMS Software	John E Lincoln	RECORDED	2018-05-15	\$390.000
501831	GLPs: How are they Associated with GMPs and SOPs	Joy McElroy	RECORDED	2018-04-17	\$390.000
501832	Analytical Method Validation Under Good Laboratory Practices - GLPs	John C. Fetzter	RECORDED	2018-04-11	\$390.000
501833	HPLC Analytical Method Development and Validation	John C. Fetzter	RECORDED	2018-06-04	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501834	Understanding, Calculating, and Using Statistical Power	John N. Zorich	RECORDED	2018-04-17	\$390.000
501836	Root Cause Failure Analysis Closed Loop Corrective Action	Michael Abitz	RECORDED	2018-05-11	\$390.000
501839	Fishbone Diagramming	Michael Abitz	RECORDED	2018-04-19	\$390.000
501840	FDA's New Import Program for 2018	Casper Uldriks	RECORDED	2018-03-12	\$390.000
501841	Computer System Validation (CSV) for FDA-Regulated Computers	Carolyn Troiano	RECORDED	2018-05-21	\$390.000
501843	21 CFR Part 11 Guidance for Electronic Records and Electronic Signatures in FDA-Regulated Industries	Carolyn Troiano	RECORDED	2018-06-12	\$390.000
501844	Complaint Handling	Peggy Berry	RECORDED	2018-04-20	\$390.000
501849	Mastering Excel 2013 Pivot Tables	Dennis Taylor	RECORDED	2018-03-28	\$390.000
501850	Time-Saving Excel Tips, Tricks and 100 Shortcuts	Dennis Taylor	RECORDED	2018-04-05	\$390.000
501851	Mastering Excel Pivot Tables	Dennis Taylor	RECORDED	2018-04-24	\$390.000
501857	Complaint Handling and Management: From Receipt to Trending	David R Dills	RECORDED	2018-06-11	\$390.000
501858	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2018-03-26	\$390.000
501860	Controlling Human Error in the Manufacturing Floor	Ginette Collazo	RECORDED	2018-04-18	\$390.000
501861	FDA's Enforcement of 21 CFR Part 11 Compliance	Angela Bazigos	RECORDED	2018-05-10	\$390.000
501862	Validation of Analytical Methods and Procedures	Dr. Ludwig Huber	RECORDED	2018-03-28	\$390.000
501863	Validation and Use of Excel® Spreadsheets in FDA Regulated Environments	Dr. Ludwig Huber	RECORDED	2018-04-05	\$390.000
501865	Water System Mythology: Common False Beliefs for Microbial Control and Monitoring	Teri C. Soli	RECORDED	2018-03-27	\$390.000
501864	FDA Issues Final Rule on Symbols in Labeling	David R Dills	RECORDED	2018-06-13	\$390.000
501866	Design History Files (DHF), Device Master Records (DMR), Device History Records (DHR), Technical Files , Design Dossiers - The Requirements	John E Lincoln	RECORDED	2018-04-25	\$390.000
501867	Laboratory Safety: Don't be Caught Unaware of Laboratory Hazards	Sheldon Primus	RECORDED	2018-04-27	\$390.000
501869	3 Hours Virtual Seminar on Boot Camp: Data Integrity - FDA's Latest Thinking	Angela Bazigos	RECORDED	2018-04-17	\$540.000
501875	How to Implement Lean Documents and Lean Document Control to Cut Costs,	Angela Bazigos	RECORDED	2018-04-16	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
	while Maintaining Compliance with Regulatory Authority Requirements				
501876	Auditing Analytical Laboratories for FDA Compliance	Steven S. Kuwahara	RECORDED	2018-05-03	\$390.000
501877	GMP Expectations for Products Used in Early Phase IND Studies	Steven S. Kuwahara	RECORDED	2018-05-16	\$390.000
501879	GDPR - European Privacy Regulation what and how	Derk Yntema	RECORDED	2018-04-13	\$390.000
501883	Master Excel: Transcend the VLOOKUP Function	David Ringstrom	RECORDED	2018-06-08	\$390.000
501896	FDA's New Import Program for 2018	Casper Uldriks	RECORDED	2018-04-10	\$390.000
501898	Essentials of Validation - IQ, OQ, PQ	Susanne Manz	RECORDED	2018-04-19	\$390.000
501899	Quality is not an Organization	Susanne Manz	RECORDED	2018-04-30	\$390.000
501904	Understanding the USP Chapter 1224 for Transfer of Analytical Methods	Dr. Ludwig Huber	RECORDED	2018-04-12	\$390.000
501905	FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	RECORDED	2018-04-30	\$390.000
501909	Supervising a Human Error Free Environment: You can do a Lot More than you Think	Ginette Collazo	RECORDED	2018-07-10	\$390.000
501911	Batch Record Review and Product Release	Danielle DeLucy	RECORDED	2018-04-27	\$390.000
501915	Project Management for Non-Project Managers	Charles H. Paul	RECORDED	2018-04-27	\$390.000
501920	If it Isn't Written Down, then it Didn't Happen: Complying with FDA's Good Documentation Practices	Angela Bazigos	RECORDED	2018-05-09	\$390.000
501921	3-hour Virtual Seminar on Project Management for Non-Project Managers	Charles H. Paul	RECORDED	2018-05-25	\$540.000
501924	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2018-05-01	\$390.000
501925	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	RECORDED	2018-05-09	\$390.000
501928	HVAC and GMP Environmental Control for Pharmaceutical Clean Rooms	Roger Cowan	RECORDED	2018-06-12	\$390.000
501932	FDA's New Import Program for 2018	Casper Uldriks	RECORDED	2018-06-19	\$390.000
501933	FDA's Revolutionary Change in Software Regulation	Casper Uldriks	RECORDED	2018-08-28	\$390.000
501936	Water System Mythology: Common False Beliefs for Microbial Control and Monitoring	Teri C. Soli	RECORDED	2018-05-15	\$390.000
501939	Validation and Use of Excel® Spreadsheets in FDA Regulated Environments	Dr. Ludwig Huber	RECORDED	2018-05-14	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501940	Latin America Regulatory Compliance: Current Challenges in Argentina, Brazil and Mexico	Robert J. Russell	RECORDED	2018-05-22	\$390.000
501941	Japan - Regulatory Filing Requirements and Compliance Processes for Life Sciences	Robert J. Russell	RECORDED	2018-07-13	\$390.000
501943	Time-Saving Excel Tips, Tricks and 100 Shortcuts	Dennis Taylor	RECORDED	2018-05-30	\$390.000
501944	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	2018-07-18	\$390.000
501949	21 CFR Part 11 - Compliance for Electronic Records and Signatures	Edwin Waldbusser	RECORDED	2018-09-10	\$390.000
501953	Death By CAPA - Does your CAPA Program Need a CAPA?	Susanne Manz	RECORDED	2018-06-12	\$390.000
501954	Secrets to Writing Effective SOPs for Medical Device QMS	Susanne Manz	RECORDED	2018-06-28	\$390.000
501955	Non-Conforming Material and Failure Investigation	Susanne Manz	RECORDED	2018-07-18	\$390.000
501956	EU General Data Protection Regulation (GDPR): Compliance for Clinical Trials - are you Ready for Implementation for May 25th 2018?	Laura Brown	RECORDED	2018-06-11	\$390.000
501960	Mastering Excel Pivot Tables	Dennis Taylor	RECORDED	2018-06-01	\$390.000
501965	Key Factors to Write an Effective Standard Operating Procedure (SOP) and Work Instructions (WIs)	Angela Bazigos	RECORDED	2018-07-26	\$390.000
501966	GDPR - European Privacy Regulation what and how	Derk Yntema	RECORDED	2018-06-12	\$390.000
501968	Batch Record Review and Product Release	Danielle DeLucy	RECORDED	2018-06-08	\$390.000
501970	The New ISO 13485: 2016 and Comparison with 21CFR820 - how to Comply with Both in the same Organization	Eyal Lerner	RECORDED	2018-06-22	\$390.000
501972	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2018-06-06	\$390.000
501973	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	RECORDED	2018-06-20	\$390.000
501974	Understanding the USP Chapter 1224 for Transfer of Analytical Methods	Dr. Ludwig Huber	RECORDED	2018-06-11	\$390.000
501975	Validation of Analytical Methods and Procedures	Dr. Ludwig Huber	RECORDED	2018-06-27	\$390.000
501977	How to Implement Lean Documents and Lean Document Control to Cut Costs, while Maintaining Compliance with Regulatory Authority Requirements	Angela Bazigos	RECORDED	2018-07-23	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501979	Controlling Human Error in the Manufacturing Floor	Ginette Collazo	RECORDED	2018-06-12	\$390.000
501980	Tools for Human Error Reduction	Ginette Collazo	RECORDED	2018-08-30	\$390.000
501981	Controlling Human Error in the Manufacturing Floor	Ginette Collazo	RECORDED	2018-09-25	\$390.000
501986	3-Hour Virtual Seminar on Complaint Handling and Management: From Receipt to Trending	David R Dills	RECORDED	2018-08-02	\$540.000
501987	The New EU Medical Device Regulation	Salma Michor	RECORDED	2018-06-27	\$390.000
501988	Combination Products Registration	Salma Michor	RECORDED	2018-07-12	\$390.000
501994	Drug Master Files: Understanding and Meeting your Regulatory and Processing Responsibilities	Robert J. Russell	RECORDED	2018-09-11	\$390.000
501995	3-Hour Virtual Seminar on Japan - Regulatory Filing Requirements and Compliance Processes for Life Sciences	Robert J. Russell	RECORDED	2018-06-27	\$540.000
501998	Auditing Analytical Laboratories for FDA Compliance	Steven S. Kuwahara	RECORDED	2018-07-10	\$390.000
501999	3-Hour Virtual Seminar on Master Validation Plan - The Unwritten Requirements	John E Lincoln	RECORDED	2018-06-20	\$540.000
502000	3-Hour Virtual Seminar on DHF, DMR, DHR, TF , Design Dossiers - The Requirements	John E Lincoln	RECORDED	2018-07-25	\$540.000
502006	3-Hour Virtual Seminar on FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	RECORDED	2018-07-12	\$540.000
502007	Validation and Use of Excel® Spreadsheets in FDA Regulated Environments	Dr. Ludwig Huber	RECORDED	2018-07-26	\$390.000
502010	Project Management for Non-Project Managers	Charles H. Paul	RECORDED	2018-06-25	\$390.000
502016	Preparing for an FDA Inspection - What you Need to Know	Susanne Manz	RECORDED	2018-08-30	\$390.000
502017	3-Hour Virtual Seminar on Mastering Excel Formulas and Functions	Mike Thomas	RECORDED	2018-07-27	\$540.000
502019	3-Hour Virtual Seminar on Project Management for Non-Project Managers	Charles H. Paul	RECORDED	2018-08-01	\$540.000
502020	Best Practices to Prepare for and Manage FDA Inspections	Vanessa Lopez	RECORDED	2018-08-14	\$390.000
502022	Excel Spreadsheets for 21 CFR 11 Compliance	Angela Bazigos	RECORDED	2018-09-27	\$390.000
502025	Accelerated Aging Techniques for Medical Device Packaging	Karl J Hemmerich	RECORDED	2018-07-18	\$390.000
502029	Handling OOS Test Results and Completing Robust Investigations	Danielle DeLucy	RECORDED	2018-07-09	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
502031	3-Hour Virtual Seminar on Statistical Process Control (SPC): A Detailed Introduction	John N. Zorich	RECORDED	2018-07-18	\$540.000
502032	FDA - Legal Writing Skills that Result in Effective Regulatory Submissions	Robert Michalik	RECORDED	2018-07-19	\$390.000
502033	3-Hour Virtual Seminar on Design Controls for Medical Devices	Susanne Manz	RECORDED	2018-08-31	\$540.000
502034	3-Hour Virtual Seminar on CAPA for Medical Devices	Susanne Manz	RECORDED	2018-08-22	\$540.000
502037	Analytical Method Validation Under Good Laboratory Practices - GLPs	John C. Fetzer	RECORDED	2018-08-13	\$390.000
502038	Water System Mythology: Common False Beliefs for Microbial Control and Monitoring	Teri C. Soli	RECORDED	2018-07-24	\$390.000
502039	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2018-07-11	\$390.000
502040	Fishbone Diagramming	Michael Abitz	RECORDED	2018-07-24	\$390.000
502042	Root Cause Failure Analysis Closed Loop Corrective Action	Michael Abitz	RECORDED	2018-08-08	\$390.000
502043	Latin America Regulatory Compliance: Current Challenges in Argentina, Brazil and Mexico	Robert J. Russell	RECORDED	2018-08-01	\$390.000
502048	Audit Trail Generation and Review	Angela Bazigos	RECORDED	2018-09-12	\$390.000
502053	GDPR - European Privacy Regulation what and how	Derk Yntema	RECORDED	2018-08-02	\$390.000
502057	Laboratory Investigation of Out-of-Specification Results	Jerry Lanese	RECORDED	2018-08-29	\$390.000
502063	EU General Data Protection Regulation (GDPR): Compliance for Clinical Trials - are you Ready for Implementation for May 25th 2018?	Laura Brown	RECORDED	2018-08-27	\$390.000
502064	Tools for Human Error Reduction	Ginette Collazo	RECORDED	2018-07-24	\$390.000
502067	Mastering Excel Pivot Tables	Dennis Taylor	RECORDED	2018-07-27	\$390.000
502069	Understanding and Implementing USP 1058 Analytical Instrument Qualification	Dr. Ludwig Huber	RECORDED	2018-07-30	\$390.000
502070	Controlling Human Error in the Manufacturing Floor	Ginette Collazo	RECORDED	2018-07-31	\$390.000
502071	Water System Biofilm Control and Microbial Monitoring Myths	Teri C. Soli	RECORDED	2018-07-31	\$390.000
502074	FDA's New Import Program for 2018	Casper Uldriks	RECORDED	2018-08-31	\$390.000
502078	3-Hour Virtual Seminar: FDA Imports Require Precision in 2018	Casper Uldriks	RECORDED	2018-08-21	\$540.000
502079	The New ISO 13485:2016 and Comparison with 21CFR820 - How	Eyal Lerner	RECORDED	2018-08-24	\$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
	to Comply with both in the Same Organization				
502083	Project Management for Non-Project Managers - Building the Work Breakdown Structure	Charles H. Paul	RECORDED	2018-08-17	\$390.000
502084	3-Hour Virtual Seminar on Project Management for Non-Project Managers	Charles H. Paul	RECORDED	2018-08-31	\$540.000
502090	Understanding the USP Chapter 1224 for Transfer of Analytical Methods	Dr. Ludwig Huber	RECORDED	2018-08-13	\$390.000
502092	Project Management for Non-Project Managers	Charles H. Paul	RECORDED	2018-10-09	\$390.000
502097	FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	RECORDED	2018-09-20	\$390.000
502099	3-Hour Virtual Seminar on Writing Effective SOPs	Angela Bazigos	RECORDED	2018-08-22	\$540.000
502106	2-Hour Virtual Seminar on Mastering Excel Pivot Tables	Dennis Taylor	RECORDED	2018-08-28	\$440.000
502107	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	RECORDED	2018-09-12	\$390.000
502108	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2018-08-08	\$390.000
502112	Pharmaceutical Drug Registration in China	Yingying Liu	RECORDED	2018-08-31	\$390.000
502116	3-Hour Virtual Seminar on FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	RECORDED	2018-08-30	\$540.000
502118	FDA's New Draft Guidance on Software and Device Changes and the 510(k)	Carolyn Troiano	RECORDED	2018-10-12	\$390.000
502119	3-Hour Virtual Seminar on GDPR - European Privacy Regulation what and how	Derk Yntema	RECORDED	2018-09-07	\$540.000
502120	Handling OOS Test Results and Completing Robust Investigations	Danielle DeLucy	RECORDED	2018-08-24	\$390.000
502121	Batch Record Review and Product Release	Danielle DeLucy	RECORDED	2018-09-19	\$390.000
502124	2-Hour Virtual Seminar on Supplier and Service Provider Controls: FDA Expectations	Vanessa Lopez	RECORDED	2018-09-06	\$440.000
502126	2-Hour Virtual Seminar on Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2018-09-05	\$440.000
502128	Key Factors to Write an Effective Standard Operating Procedure (SOP) and Work Instructions (WIs)	Angela Bazigos	RECORDED	2018-09-18	\$390.000
502131	EU Medical Device Regulation (MDR) - Updated CE Marking Process & ISO 13485:2016 Expectations	Robert J. Russell	RECORDED	2018-10-09	\$390.000
502133	3-Hour Virtual Seminar on Excel Spreadsheet Validation for FDA 21 CFR Part 11	Angela Bazigos	RECORDED	2018-09-11	\$540.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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502156	Good Documentation Guideline (Chapter 1029 USP)	Dr. Afsaneh Motamed Khorasani	RECORDED	2018-10-11	\$390.000
502166	3-Hour Virtual Seminar on Validation, Verification and Transfer of Analytical Methods - Implementing Guidelines from FDA-EMA, USP and ICH	Dr. Ludwig Huber	RECORDED	2018-09-27	\$540.000
502177	3-Hour Virtual Seminar on Data Integrity - FDA's Latest Thinking	Angela Bazigos	RECORDED	2018-10-01	\$540.000
502179	Medical Devices - Complying With 820.250 Statistical Methods	Wayne Taylor	RECORDED	2018-10-12	\$390.000
502181	3-Hour Virtual Seminar on DHF, DMR, DHR, TF , Design Dossiers - The Requirements	John E Lincoln	RECORDED	2018-09-25	\$540.000
502184	Process Validation: Statistical Justification for Sample Size and the Use of Only 3 Lots	John N. Zorich	RECORDED	2018-10-03	\$390.000
502190	Mastering Excel Pivot Tables	Dennis Taylor	RECORDED	2018-10-02	\$390.000
502192	EU General Data Protection Regulation (GDPR): Compliance for Drug development and Clinical Trials?	Laura Brown	RECORDED	2018-10-12	\$390.000
502196	Validation and Use of Excel® Spreadsheets in FDA Regulated Environments	Dr. Ludwig Huber	RECORDED	2018-10-10	\$390.000