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	LIVE RECORDED	2018-07-18	$290.000 $540.000
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	LIVE RECORDED	2018-07-18	$150.000 $390.000
502025	Accelerated Aging Techniques for Medical Device Packaging	Karl J Hemmerich	LIVE RECORDED	2018-07-18	$150.000 $390.000
501955	Non-Conforming Material and Failure Investigation	Susanne Manz	LIVE RECORDED	2018-07-18	$150.000 $390.000
502032	FDA - Legal Writing Skills that Result in Effective Regulatory Submissions	Robert Michalik	LIVE RECORDED	2018-07-19	$150.000 $390.000
502033	3-Hour Virtual Seminar on Design Controls for Medical Devices	Susanne Manz	LIVE RECORDED	2018-07-19	$290.000 $540.000
502012	The Value of a Human Factors Program	Thomas Bento	LIVE RECORDED	2018-07-20	$150.000 $390.000
502030	Implementation and Management of GMP Data Integrity	Danielle DeLucy	LIVE RECORDED	2018-07-23	$150.000 $390.000
502036	HPLC Analytical Method Development and Validation	John C. Fetzer	LIVE RECORDED	2018-07-23	$150.000 $390.000
502038	Water System Mythology: Common False Beliefs for Microbial Control and Monitoring	Teri C. Soli	LIVE RECORDED	2018-07-24	$150.000 $390.000
502040	Fishbone Diagramming	Michael Abitz	LIVE RECORDED	2018-07-24	$150.000 $390.000
502044	Pre-Control: Easier than SPC	Jd Marhevko	LIVE RECORDED	2018-07-24	$150.000 $390.000
502064	Tools for Human Error Reduction	Ginette Collazo	LIVE RECORDED	2018-07-24	$150.000 $390.000
502000	3-Hour Virtual Seminar on DHF, DMR, DHR, TF , Design Dossiers - The Requirements	John E Lincoln	LIVE RECORDED	2018-07-25	$290.000 $540.000
501994	Drug Master Files: Understanding and Meeting your Regulatory and Processing Responsibilities	Robert J. Russell	LIVE RECORDED	2018-07-25	$150.000 $390.000
501762	Working Effectively with Customers & Suppliers	Charles H. Paul	LIVE RECORDED	2018-07-26	$150.000 $390.000
502007	Validation and Use of Excel® Spreadsheets in FDA Regulated Environments	Dr. Ludwig Huber	LIVE RECORDED	2018-07-26	$150.000 $390.000
502087	Combination Products: A Regulatory Perspective	Thomas E. Colonna	LIVE RECORDED	2018-07-26	$150.000 $390.000
501965	Key Factors to Write an Effective Standard Operating Procedure (SOP) and Work Instructions (WIs)	Angela Bazigos	LIVE RECORDED	2018-07-26	$150.000 $390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501951	Effective Design of Experiments (DOE) Strategies	Jose Mora	LIVE RECORDED	2018-07-26	$150.000 $390.000
502017	3-Hour Virtual Seminar on Mastering Excel Formulas and Functions	Mike Thomas	LIVE RECORDED	2018-07-27	$290.000 $540.000
502018	Excel Spreadsheets - Step-By-Step Instructions for Ensuring Data Integrity	David Nettleton	LIVE	2018-07-27	$150.000
502067	Mastering Excel Pivot Tables	Dennis Taylor	LIVE RECORDED	2018-07-27	$150.000 $390.000
502082	Project Management for Non-Project Managers - Achieving Results - Building the Foundation	Charles H. Paul	LIVE RECORDED	2018-07-30	$150.000 $390.000
502014	Purchasing Control Essentials for Medical Devices	Susanne Manz	LIVE RECORDED	2018-07-30	$150.000 $390.000
502066	21 CFR Part 11 - Compliance for Electronic Records and Signatures	David Nettleton	LIVE	2018-07-30	$150.000
502069	Understanding and Implementing USP 1058 Analytical Instrument Qualification	Dr. Ludwig Huber	LIVE RECORDED	2018-07-30	$150.000 $390.000
502004	FDA's New Draft Guidance on Software and Device Changes and the 510(k)	Carolyn Troiano	LIVE RECORDED	2018-07-31	$150.000 $390.000
502074	FDA's New Import Program for 2018	Casper Uldriks	LIVE RECORDED	2018-07-31	$150.000 $390.000
502045	Key Steps to Successful Project Management Principles	Carol Friedhoff	LIVE RECORDED	2018-07-31	$150.000 $390.000
502041	Medical Device Single Audit Program (MDSAP) Preparation	Shep Bentley	LIVE RECORDED	2018-07-31	$150.000 $390.000
502026	Validating Radiation Sterilization for Medical Products	Karl J Hemmerich	LIVE RECORDED	2018-07-31	$150.000 $390.000
501959	Protocol Deviation Reporting and Management	Grace Morgan Holmes	LIVE RECORDED	2018-07-31	$150.000 $390.000
502070	Controlling Human Error in the Manufacturing Floor	Ginette Collazo	LIVE RECORDED	2018-07-31	$150.000 $390.000
502071	Water System Biofilm Control and Microbial Monitoring Myths	Teri C. Soli	LIVE RECORDED	2018-07-31	$150.000 $390.000
502019	3-Hour Virtual Seminar on Project Management for Non-Project Managers	Charles H. Paul	LIVE RECORDED	2018-08-01	$290.000 $540.000
502107	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	LIVE RECORDED	2018-08-01	$150.000 $390.000
502043	Latin America Regulatory Compliance: Current Challenges in Argentina, Brazil and Mexico	Robert J. Russell	LIVE RECORDED	2018-08-01	$150.000 $390.000
502052	Food Safety Modernization Act	Thomas Perkins	LIVE RECORDED	2018-08-01	$150.000 $390.000
502054	Ethical Compliance Starts with Accountability	Ron Rael	LIVE RECORDED	2018-08-02	$150.000 $390.000
501986	3-Hour Virtual Seminar on Complaint Handling and Management: From Receipt to Trending	David Dills	LIVE RECORDED	2018-08-02	$290.000 $540.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
502053	GDPR - European Privacy Regulation what and how	Derk Yntema	LIVE RECORDED	2018-08-02	$150.000 $390.000
502072	Save your Food Business!! - Business Continuity & Disaster Recovery	Angela Bazigos	LIVE RECORDED	2018-08-02	$150.000 $390.000
502062	Safety and Motivation for EHS Practitioners	James Thatcher	LIVE RECORDED	2018-08-03	$150.000 $390.000
502021	Electronic Records & Electronic Signatures, 21 CFR Part 11, Basic Concepts	Angela Bazigos	LIVE RECORDED	2018-08-06	$150.000 $390.000
502005	Good Documentation Practices to Support FDA Computer System Validation	Carolyn Troiano	LIVE RECORDED	2018-08-07	$150.000 $390.000
502083	Project Management for Non-Project Managers - Building the Work Breakdown Structure	Charles H. Paul	LIVE RECORDED	2018-08-08	$150.000 $390.000
502108	Valid Statistical Rationales for Sample Sizes	John N. Zorich	LIVE RECORDED	2018-08-08	$150.000 $390.000
502056	Analytical Method Validation	Jerry Lanese	LIVE RECORDED	2018-08-08	$150.000 $390.000
502042	Root Cause Failure Analysis Closed Loop Corrective Action	Michael Abitz	LIVE RECORDED	2018-08-08	$150.000 $390.000
502085	Adaptive Designs for Medical Device Clinical Studies	Angela Bazigos	LIVE RECORDED	2018-08-09	$150.000 $390.000
502101	Packaging and Labeling in the Pharmaceutical Supply Chain	Michael Esposito	LIVE RECORDED	2018-08-09	$150.000 $390.000
502022	Excel Spreadsheets for 21 CFR 11 Compliance	Angela Bazigos	LIVE RECORDED	2018-08-10	$150.000 $390.000
502059	Understanding the Requirements of SSAE18	Lynn Fountain	LIVE RECORDED	2018-08-10	$150.000 $390.000
502037	Analytical Method Validation Under Good Laboratory Practices - GLPs	John C. Fetzer	LIVE RECORDED	2018-08-13	$150.000 $390.000
502090	Understanding the USP Chapter 1224 for Transfer of Analytical Methods	Dr. Ludwig Huber	LIVE RECORDED	2018-08-13	$150.000 $390.000
502020	Best Practices to Prepare for and Manage FDA Inspections	Vanessa Lopez	LIVE RECORDED	2018-08-14	$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	2018-08-14	$290.000 $540.000
502011	3-Hour Virtual Seminar on Data Integrity: Compliance with 21 CFR Part 11 and SaaS-Cloud Software Applications	David Nettleton	LIVE	2018-08-14	$290.000
502015	Design Control Basics	Susanne Manz	LIVE RECORDED	2018-08-14	$150.000 $390.000
502075	Determining Sample Size: What Sample Size Should I Use?	Steven Wachs	LIVE RECORDED	2018-08-15	$150.000 $390.000
502001	3-Hour Virtual Seminar on Verification vs Validation-Product, Process or Equipment and QMS Software	John E Lincoln	LIVE RECORDED	2018-08-15	$290.000 $540.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
502024	What do the FDA, EMA and PMDA Look for When Conducting cGMP Regulatory Inspections	Angela Bazigos	LIVE RECORDED	2018-08-16	$150.000 $390.000
502051	Calibrating the Human Gage: Attribute Agreement Analysis	Jd Marhevko	LIVE RECORDED	2018-08-16	$150.000 $390.000
502099	3-Hour Virtual Seminar on Writing Effective SOPs	Angela Bazigos	LIVE RECORDED	2018-08-17	$290.000 $540.000
502095	Validation of Non-Product Software	Thomas Bento	LIVE RECORDED	2018-08-17	$150.000 $390.000
502089	3-Hour Virtual Seminar on Japan - Regulatory Filing Requirements and Compliance Processes for Life Sciences	Robert J. Russell	LIVE RECORDED	2018-08-17	$290.000 $540.000
502105	Time-Saving Excel Tips, Tricks and 100 Shortcuts	Dennis Taylor	LIVE RECORDED	2018-08-17	$150.000 $390.000
502098	Project Management for Computer Systems Validation and 21 CFR 11 - Annex 11	Angela Bazigos	LIVE RECORDED	2018-08-20	$150.000 $390.000
502102	FDA Regulation of Human Cells,Tissues and Cellular and Tissue-Based Products (HCT-Ps)	Marina Malikova	LIVE RECORDED	2018-08-20	$150.000 $390.000
502078	3-Hour Virtual Seminar: FDA Imports Require Precision in 2018	Casper Uldriks	LIVE RECORDED	2018-08-21	$290.000 $540.000
502047	3-Hour Virtual Seminar on How the FDA are Trained for Medical Device Inspections	Angela Bazigos	LIVE RECORDED	2018-08-21	$290.000 $540.000
502034	3-Hour Virtual Seminar on CAPA for Medical Devices	Susanne Manz	LIVE RECORDED	2018-08-22	$290.000 $540.000
502055	Strategic Management of Corporate Governance: Setting the Right Tone at the Top about Risk, Part 1	Ron Rael	LIVE RECORDED	2018-08-23	$150.000 $390.000
502048	Audit Trail Generation and Review	Angela Bazigos	LIVE RECORDED	2018-08-23	$150.000 $390.000
502079	The New ISO 13485:2016 and Comparison with 21CFR820 - How to Comply with both in the Same Organization	Eyal Lerner	LIVE RECORDED	2018-08-24	$150.000 $390.000
502110	Managing Excel Lists and Databases: Time-Saving Shortcuts for Sorting, Filtering, Analyzing and More	Tom Fragale	LIVE RECORDED	2018-08-24	$150.000 $390.000
502096	Behavior Based Safety System: A Guide to Building a Safety Culture	Sheldon Primus	LIVE RECORDED	2018-08-24	$150.000 $390.000
502103	Data Integrity in Compliance with GXP-GMP Regulations	Eleonora Babayants	LIVE RECORDED	2018-08-24	$150.000 $390.000
502049	How to Implement Good Clinical Practices for Successful FDA and International Regulatory Authority Inspections	Angela Bazigos	LIVE RECORDED	2018-08-24	$150.000 $390.000
502076	Computer System Validation (CSV) for FDA-Regulated Computers	Carolyn Troiano	LIVE RECORDED	2018-08-27	$150.000 $390.000
502063	EU General Data Protection Regulation (GDPR): Compliance for Clinical Trials -	Laura Brown	LIVE RECORDED	2018-08-27	$150.000 $390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
	are you Ready for Implementation for May 25th 2018?				
502050	Export Certificate for Medical Devices - Edge Out Your Competition!	Angela Bazigos	LIVE RECORDED	2018-08-28	$150.000 $390.000
502106	2-Hour Virtual Seminar on Mastering Excel Pivot Tables	Dennis Taylor	LIVE RECORDED	2018-08-28	$190.000 $440.000
501933	FDA's Revolutionary Change in Software Regulation	Casper Uldriks	LIVE RECORDED	2018-08-28	$150.000 $390.000
502057	Laboratory Investigation of Out-of-Specification Results	Jerry Lanese	LIVE RECORDED	2018-08-29	$150.000 $390.000
502060	COSO New Fraud Principals	Lynn Fountain	LIVE RECORDED	2018-08-29	$150.000 $390.000
502116	3-Hour Virtual Seminar on FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	LIVE RECORDED	2018-08-30	$290.000 $540.000
502016	Preparing for an FDA Inspection - What you Need to Know	Susanne Manz	LIVE RECORDED	2018-08-30	$150.000 $390.000
502086	MDUFA IV - An Introduction	Angela Bazigos	LIVE RECORDED	2018-08-30	$150.000 $390.000
501980	Tools for Human Error Reduction	Ginette Collazo	LIVE RECORDED	2018-08-30	$150.000 $390.000
502080	Obscure But Effective Regulatory Pathways for Devices	Shep Bentley	LIVE RECORDED	2018-08-31	$150.000 $390.000
502084	Project Management for Non-Project Managers -Building the Project Schedule	Charles H. Paul	LIVE RECORDED	2018-08-31	$150.000 $390.000
502088	Preparing for Pre and Post Approval Inspections	Lamont M. Fulton	LIVE RECORDED	2018-08-31	$150.000 $390.000
502112	Pharmaceutical Drug Registration in China	Yingying Liu	LIVE RECORDED	2018-08-31	$150.000 $390.000
502109	Engaging with the FDA During New Drug Development	Angela Bazigos	LIVE RECORDED	2018-09-07	$150.000 $390.000
502100	Onboarding Employees in a GMP Environment: Best Practices for Foundational Employee Success	Michael Esposito	LIVE RECORDED	2018-09-10	$150.000 $390.000
501949	21 CFR Part 11 - Compliance for Electronic Records and Signatures	Edwin Waldbusser	LIVE RECORDED	2018-09-10	$150.000 $390.000
502114	3-Hours Virtual Seminar on Highlights of FDA GLP Regulations and the Roles and Responsibilities in a GLP Facility	Shib Mookherjea	LIVE RECORDED	2018-09-10	$290.000 $540.000
502061	Managing and Examining Travel and Entertainment Expense	Lynn Fountain	LIVE RECORDED	2018-09-11	$150.000 $390.000
502077	Medical Device Cybersecurity -Improving Compliance and your Company's Bottom Line	Carolyn Troiano	LIVE RECORDED	2018-09-11	$150.000 $390.000
502073	Compliance for Risk Based Approaches for Clinical Trials	Laura Brown	LIVE RECORDED	2018-09-12	$150.000 $390.000
502091	Project Management for Non-Project Managers - Estimating Resource Needs	Charles H. Paul	LIVE RECORDED	2018-09-12	$150.000 $390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
502104	Governance and Risk Control According to GxP-GMP Requirements	Eleonora Babayants	LIVE RECORDED	2018-09-14	$150.000 $390.000
502081	Quality Agreements Made Easy	Shep Bentley	LIVE RECORDED	2018-09-14	$150.000 $390.000
502058	cGMPs in the Quality Control Laboratory	Jerry Lanese	LIVE RECORDED	2018-09-14	$150.000 $390.000
502097	FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	LIVE RECORDED	2018-09-20	$150.000 $390.000
502113	Clinical Trial Applications in China	Yingying Liu	LIVE RECORDED	2018-09-21	$150.000 $390.000
501981	Controlling Human Error in the Manufacturing Floor	Ginette Collazo	LIVE RECORDED	2018-09-25	$150.000 $390.000
502094	Project Management for Non-Project Managers - Assessing and Planning for Risk	Charles H. Paul	LIVE RECORDED	2018-09-25	$150.000 $390.000
502111	PowerPoint Tips and Tricks	Tom Fragale	LIVE RECORDED	2018-09-28	$150.000 $390.000
502035	Medical Device Hazard Analysis Following ISO 14971	Edwin Waldbusser	LIVE RECORDED	2018-10-02	$150.000 $390.000
502092	Project Management for Non-Project Managers - Building the Project Team	Charles H. Paul	LIVE RECORDED	2018-10-09	$150.000 $390.000
502115	Computer System Validation	Edwin Waldbusser	LIVE RECORDED	2018-10-23	$150.000 $390.000
501982	Training in Human Error: Reducing Training Related Errors	Ginette Collazo	LIVE RECORDED	2018-10-24	$150.000 $390.000
502093	Project Management for Non-Project Managers -Managing and Controlling the Project	Charles H. Paul	LIVE RECORDED	2018-10-29	$150.000 $390.000
501984	Controlling Human Error in the Manufacturing Floor	Ginette Collazo	LIVE RECORDED	2018-11-29	$150.000 $390.000
500001	Introduction to Sales and Use Taxation	Miles Hutchinson	RECORDED	2014-11-12	$149.000
500002	Hot Issues in Determining Multi-State Sales Tax Nexus	Miles Hutchinson	RECORDED	2014-11-25	$149.000
500004	Complying with FATCA - the Foreign Accounts Tax Compliance Act	Miles Hutchinson	RECORDED	2014-05-22	$149.000
500007	Compliance and Continual Improvement, Tools of the Trade	Rodriguez Gonzalez	RECORDED	2014-04-16	$149.000
500010	Foodservice Sanitation	Norman G. Marriott	RECORDED	2014-07-15	$149.000
500012	Foreign Corrupt Practices Act - Audits Role	Denise Cicchella	RECORDED	2014-05-27	$149.000
500013	Operational Risk - Beyond Regulatory Constraints	Fred Vacelet	RECORDED	2014-07-02	$149.000
500015	System Safety Engineering	Dev Raheja	RECORDED	2014-05-20	$149.000
500016	Fundamental ATC Organization, Structure, and Concepts	Stephen Alvania	RECORDED	2014-05-20	$149.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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 Dills	RECORDED	2014-11-24	$189.000
500080	FDA Compliant HPLC Qualification and Performance Testing	Dr. Ludwig Huber	RECORDED	2014-11-06	$189.000
500081	GMP Perspectives on Working with Contracting Laboratories	Steven S. Kuwahara	RECORDED	2014-11-11	$189.000
500082	Cleaning and Sanitation Training for Food Processors	Melinda Allen	RECORDED	2014-11-13	$189.000
500083	Auditing Analytical Laboratories for FDA Compliance	Steven S. Kuwahara	RECORDED	2014-11-18	$189.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
500084	Compiling the Design History File, and Technical File, Design Dossier	John E Lincoln	RECORDED	2014-11-18	$189.000
500086	FDA's 2011 Draft Guidance on Financial Disclosure by Clinical Investigators	David Dills	RECORDED	2014-11-19	$189.000
500088	The European Clinical Trial Directive (EUCTD)	Robert J. Russell	RECORDED	2014-12-16	$189.000
500089	Quality by Design (QbD) for Development and Validation of Analytical Methods	Dr. Ludwig Huber	RECORDED	2014-11-20	$189.000
500091	FDA Regulation of In Vitro Diagnostics	Thomas E. Colonna	RECORDED	2014-11-20	$189.000
500092	Statistical Concepts of Process Validation	Dan OLeary	RECORDED	2015-01-22	$189.000
500093	Change Control - Key to Successful cGMP Compliance	John E Lincoln	RECORDED	2014-12-02	$189.000
500095	How to Manage a Medical Device Recall Efficiently and Effectively	David Dills	RECORDED	2014-12-03	$189.000
500097	The FDA Drug Development Process: GLP, GMP and GCP Regulations	Albert A. Ghignone	RECORDED	2014-11-06	$189.000
500102	Understanding Combination Products, Requests for Designation and Product Jurisdiction	Elisa Harvey	RECORDED	2014-11-05	$189.000
500103	Using the Pre-Submission Process to Your Best Advantage	Elisa Harvey	RECORDED	2014-12-03	$189.000
500105	Foreign Bodies in Foods - Effective Techniques for Prevention, Control and Detection	Rotimi Toki	RECORDED	2014-11-21	$149.000
500106	Food Labelling: A Practical Guide for Unambiguous Labelling Information & Compliance with Regulatory Requirements	Rotimi Toki	RECORDED	2014-12-12	$149.000
500108	EU Pharmacovigilance Directive and Regulations	Robert J. Russell	RECORDED	2014-12-10	$189.000
500110	Developing an Efficient Relationship with FDA	Robert Kunka	RECORDED	2014-12-16	$189.000
500111	Device Changes, FDA Changes, and the 510(k)	John E Lincoln	RECORDED	2014-12-16	$189.000
500112	Maintaining an Effective CAPA Program and Using Risk Assessment Tools: Current Trends	David Dills	RECORDED	2014-12-17	$189.000
500113	Device Corrections and Removals	Dan OLeary	RECORDED	2014-12-18	$189.000
500114	GMP Expectations for Products Used in Early Phase IND Studies	Steven S. Kuwahara	RECORDED	2015-01-06	$189.000
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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
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500554	Driving a Quality-Lean Steering Committee	Jd Marhevko	RECORDED	2015-11-24	$152.000

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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
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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
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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
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500613	Sample Size and Statistical Rationale for Medical Device Packaging Validations	Karen Greene	RECORDED	2016-02-04	$190.000
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500662	Accelerated Aging Techniques for Medical Device Packaging	Karl J Hemmerich	RECORDED	2016-01-29	$190.000
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500671	Spreadsheet-Based Internal Controls	David Ringstrom	RECORDED	2016-02-04	$190.000
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500684	Current Status of biosimilars in US, Europe and Canada	Anne Tomalin	RECORDED	2016-03-03	$190.000
500687	Food Safety, Security and Fraud-Are You Ready	Michael Brodsky	RECORDED	2016-02-01	$190.000

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500692	Effective Training Practices for FDA Compliance	Dr. Ludwig Huber	RECORDED	2016-04-13	$390.000
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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
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500745	FDA Regulations for Analytical Instrument Qualification and Validation Processes	Joy McElroy	RECORDED	2016-03-29	$190.000
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500747	FDA Regulations for Environmental Monitoring(EM) Program	Joy McElroy	RECORDED	2016-04-28	$390.000
500749	The New European Clinical Trial Regulation	Robert J. Russell	RECORDED	2016-04-06	$190.000

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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
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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
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500848	FDA's Medical Device Clinical Trials Program	Anna Longwell	RECORDED	2016-06-08	$290.000
500849	A second look at 510(k) changes	Anna Longwell	RECORDED	2016-06-28	$390.000

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500860	FDA Export Certificates for Medical Devices	Larry Spears	RECORDED	2016-07-05	$390.000
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500867	Combination Products: A Regulatory Perspective	Thomas E. Colonna	RECORDED	2016-08-25	$390.000
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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
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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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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
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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
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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
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500953	Fundamentals Of Technology Transfer And Intellectual Property Licensing	Ronald Kudla	RECORDED	2016-08-04	$390.000
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500966	Successful Through Wall Flashing Systems	Leonard Anastasi	RECORDED	2016-08-18	$390.000
500969	Water System Biofilm Control and Microbial Monitoring Myths	Teri C. Soli	RECORDED	2016-09-12	$390.000
500970	Using Statistics to Determine Sample Size	Steven Walfish	RECORDED	2016-09-20	$390.000
500971	Statistical Power and Sample Size	John N. Zorich	RECORDED	2016-08-30	$390.000
500972	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2016-09-19	$390.000
500976	21 CFR Part 820 - Quality System Regulation - Applying Principles of Lean Documents and Lean Configuration	Jose Mora	RECORDED	2016-09-14	$390.000
500977	Design History File (DHF), the Device Master Record (DMR) and the Device History Record (DHR) – Principles on Lean Documents and Lean Configuration	Jose Mora	RECORDED	2016-10-05	$390.000
500978	Good Documentation Practice for GxP Environments	John R. Godshalk	RECORDED	2016-10-12	$390.000
500979	Implementing the New FDA Data Integrity Guide	Dr. Ludwig Huber	RECORDED	2016-09-08	$390.000
500980	Understanding and Preparing for FDA's on-going Part 11 Inspections	Dr. Ludwig Huber	RECORDED	2016-10-04	$390.000
500981	Good Laboratory Practice Regulations - Introduction and Strategies for Implementation	Dr. Ludwig Huber	RECORDED	2016-11-02	$390.000
500984	FDA's 21 CFR 11 Add-On Inspections - Recent Updates	Angela Bazigos	RECORDED	2016-10-05	$390.000
500985	FDA's Medical Device Clinical Trials Program	Anna Longwell	RECORDED	2016-10-11	$390.000
500988	Applying Statistical Process Control Effectively	Steven Wachs	RECORDED	2016-10-10	$390.000
500992	Laboratory Investigation of Out-of-Specification Results	Jerry Lanese	RECORDED	2016-12-05	$390.000
501003	Understanding FDA's Quality Metrics Draft Guidance and Its Impact	Louis Angelucci	RECORDED	2016-10-06	$390.000
501019	Determining Sample Size: How to Ensure You Get the Correct	Steven Wachs	RECORDED	2016-11-14	$390.000
501020	Time-Saving Excel Tips, Tricks and 100 Shortcuts	Dennis Taylor	RECORDED	2016-10-17	$390.000
501022	The Bayh-Dole Act and Its Implementing Regulations & Rule-Making	Ronald Kudla	RECORDED	2016-10-19	$390.000
501026	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	RECORDED	2016-10-11	$390.000

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

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

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

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

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

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

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

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

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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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 Dills	RECORDED	2017-11-28	$390.000
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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
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 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

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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
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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
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

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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. Fetzer	RECORDED	2018-04-11	$390.000
501833	HPLC Analytical Method Development and Validation	John C. Fetzer	RECORDED	2018-06-04	$390.000
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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 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 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, while Maintaining Compliance with Regulatory Authority Requirements	Angela Bazigos	RECORDED	2018-04-16	$390.000
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

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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
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
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
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
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

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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
501979	Controlling Human Error in the Manufacturing Floor	Ginette Collazo	RECORDED	2018-06-12	$390.000
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
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
502006	3-Hour Virtual Seminar on FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	RECORDED	2018-07-12	$540.000
502010	Project Management for Non-Project Managers	Charles H. Paul	RECORDED	2018-06-25	$390.000
502029	Handling OOS Test Results and Completing Robust Investigations	Danielle DeLucy	RECORDED	2018-07-09	$390.000
502039	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2018-07-11	$390.000