Compliance4All Webinar

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Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501947	FDA Review of 510(k)s - How to Speed the Process	Shep Bentley	LIVE RECORDED	2018-05-25	$150.000 $390.000
501921	3-hour Virtual Seminar on Project Management for Non-Project Managers	Charles H. Paul	LIVE RECORDED	2018-05-25	$290.000 $540.000
501934	How Should you Prepare for a BSA Regulatory Examination	Thomas Nollner	LIVE RECORDED	2018-05-29	$150.000 $390.000
501927	Sterile Filtration of Pharmaceutical Products	Roger Cowan	LIVE RECORDED	2018-05-29	$150.000 $390.000
501943	Time-Saving Excel Tips, Tricks and 100 Shortcuts	Dennis Taylor	LIVE RECORDED	2018-05-30	$150.000 $390.000
501902	Risk Management Techniques for Medical Devices	Susanne Manz	LIVE RECORDED	2018-05-30	$150.000 $390.000
501956	EU General Data Protection Regulation (GDPR): Compliance for Clinical Trials - are you Ready for Implementation for May 25th 2018?	Laura Brown	LIVE RECORDED	2018-05-31	$150.000 $390.000
501935	Understanding the Complexity of OSHA's Recordkeeping Regulation	Michael Aust	LIVE RECORDED	2018-05-31	$150.000 $390.000
501960	Mastering Excel Pivot Tables	Dennis Taylor	LIVE RECORDED	2018-06-01	$150.000 $390.000
501963	GLPs: How are they Associated with GMPs and SOPs	Joy McElroy	LIVE RECORDED	2018-06-04	$150.000 $390.000
501833	HPLC Analytical Method Development and Validation	John C. Fetzer	LIVE RECORDED	2018-06-04	$150.000 $390.000
501929	Criteria for IRB Approval - Essential Training for IRB Members and Staff	George Gasparis	LIVE RECORDED	2018-06-05	$150.000 $390.000
501686	3-Hour Virtual Seminar on Off label Promotion of Drugs and Medical Devices - FDA's Latest	Angela Bazigos	LIVE RECORDED	2018-06-05	$290.000 $540.000
501962	Advanced Legal Writing Methodologies for Successful FDA 'Breakthrough Therapy' Submissions	Robert Michalik	LIVE RECORDED	2018-06-06	$150.000 $390.000
501972	Valid Statistical Rationales for Sample Sizes	John N. Zorich	LIVE RECORDED	2018-06-06	$150.000 $390.000
501952	FDA Export Certificates for Medical Devices	Larry Spears	LIVE RECORDED	2018-06-06	$150.000 $390.000
501942	The Safety Maturity Curve - Where does your Organization Stand?	Michael Aust	LIVE RECORDED	2018-06-07	$150.000 $390.000
501890	Corrective and Preventive Action (CAPA) per FDA Requirements	Eleonora Babayants	LIVE RECORDED	2018-06-07	$150.000 $390.000
501914	Successful Supplier Audits	Danielle DeLucy	LIVE RECORDED	2018-06-07	$150.000 $390.000
501968	Batch Record Review and Product Release	Danielle DeLucy	LIVE RECORDED	2018-06-08	$150.000 $390.000
501948	Risk Management Simplified	Shep Bentley	LIVE RECORDED	2018-06-08	$150.000 $390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501971	Discussion Regarding the Substitution of a New Inactive Ingredient in an Approved Parenteral Drug Product	Lamont M. Fulton	LIVE RECORDED	2018-06-08	$150.000 $390.000
501895	Oversight of CROs-Vendors-CMOs	Laura Brown	LIVE RECORDED	2018-06-08	$150.000 $390.000
501918	Data Integrity Compliance with 21 CFR Part 11 and SaaS/Cloud Software Applications	David Nettleton	LIVE	2018-06-08	$150.000
501883	Master Excel: Transcend the VLOOKUP Function	David Ringstrom	LIVE RECORDED	2018-06-08	$150.000 $390.000
501974	Understanding the USP Chapter 1224 for Transfer of Analytical Methods	Dr. Ludwig Huber	LIVE RECORDED	2018-06-11	$150.000 $390.000
501957	Developing and Implementing Effective Corrective Action Plans in Clinical Trials	Grace Morgan Holmes	LIVE RECORDED	2018-06-11	$150.000 $390.000
501857	Complaint Handling and Management: From Receipt to Trending	David Dills	LIVE RECORDED	2018-06-11	$150.000 $390.000
501922	Introduction to Good Manufacturing Practices (GMP)	Michael Esposito	LIVE RECORDED	2018-06-11	$150.000 $390.000
501979	Controlling Human Error in the Manufacturing Floor	Ginette Collazo	LIVE RECORDED	2018-06-12	$150.000 $390.000
501961	Verification vs. Validation in Regulated Industries for Processes	Jack Dhuwalia	LIVE RECORDED	2018-06-12	$150.000 $390.000
501928	HVAC and GMP Environmental Control for Pharmaceutical Clean Rooms	Roger Cowan	LIVE RECORDED	2018-06-12	$150.000 $390.000
501953	Death By CAPA - Does your CAPA Program Need a CAPA?	Susanne Manz	LIVE RECORDED	2018-06-12	$150.000 $390.000
501843	21 CFR Part 11 Guidance for Electronic Records and Electronic Signatures in FDA-Regulated Industries	Carolyn Troiano	LIVE RECORDED	2018-06-12	$150.000 $390.000
501983	Laboratory Safety: Don't be Caught Unaware of Laboratory Hazards	Sheldon Primus	LIVE RECORDED	2018-06-12	$150.000 $390.000
501966	GDPR - European Privacy Regulation what and how	Derk Yntema	LIVE RECORDED	2018-06-12	$150.000 $390.000
501746	How to Comply with 21 CFR 11 Requirements for Electronic Medical Records	Angela Bazigos	LIVE RECORDED	2018-06-13	$150.000 $390.000
501945	Through the Eyes of an Auditor	Jose Mora	LIVE RECORDED	2018-06-13	$150.000 $390.000
501994	Drug Master Files: Understanding and Meeting your Regulatory and Processing Responsibilities	Robert J. Russell	LIVE RECORDED	2018-06-13	$150.000 $390.000
501864	FDA Issues Final Rule on Symbols in Labeling	David Dills	LIVE RECORDED	2018-06-13	$150.000 $390.000
501817	Medical Device Software Validation Meeting FDA Regulations	Edwin Waldbusser	LIVE RECORDED	2018-06-14	$150.000 $390.000
501919	Responding to FDA 483s and FDA Warning Letters	Angela Bazigos	LIVE RECORDED	2018-06-14	$150.000 $390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501991	The Cost of Ineffective Leadership: Maximizing Customer Satisfaction and Profit	Michael Abitz	LIVE RECORDED	2018-06-15	$150.000 $390.000
501964	FDA Regulations for Environmental Monitoring(EM) Program	Joy McElroy	LIVE RECORDED	2018-06-18	$150.000 $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	LIVE RECORDED	2018-06-18	$150.000 $390.000
501932	FDA's New Import Program for 2018	Casper Uldriks	LIVE RECORDED	2018-06-19	$150.000 $390.000
501997	What do the FDA, EMA and PMDA Look for When Conducting cGMP Regulatory Inspections	Angela Bazigos	LIVE RECORDED	2018-06-20	$150.000 $390.000
501999	3-Hour Virtual Seminar on Master Validation Plan - The Unwritten Requirements	John E Lincoln	LIVE RECORDED	2018-06-20	$290.000 $540.000
501973	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	LIVE RECORDED	2018-06-20	$150.000 $390.000
501958	Medical Devices: Hazard Identification and Risk Assignment Using Public Data Sources	Christina Bernstein	LIVE RECORDED	2018-06-20	$150.000 $390.000
501946	Chipping Away at Constraints - Practical Tips to Improve your Manufacturing Operations	Jose Mora	LIVE RECORDED	2018-06-20	$150.000 $390.000
501987	The New EU Medical Device Regulation	Salma Michor	LIVE RECORDED	2018-06-21	$150.000 $390.000
502008	2011 FDA Guideline on Process Validation	Louis Angelucci	LIVE RECORDED	2018-06-21	$150.000 $390.000
501989	3 hour Virtual Seminar On Medical Device Recalls: How to Properly, Compliantly, and Promptly Deal with a Recall	Angela Bazigos	LIVE RECORDED	2018-06-21	$290.000 $540.000
501970	The New ISO 13485: 2016 and Comparison with 21CFR820 - how to Comply with Both in the same Organization	Eyal Lerner	LIVE RECORDED	2018-06-22	$150.000 $390.000
501967	Software Verification & Validation	Lena Cordie	LIVE RECORDED	2018-06-22	$150.000 $390.000
502010	Project Management for Non-Project Managers	Charles H. Paul	LIVE RECORDED	2018-06-25	$150.000 $390.000
501986	3-Hour Virtual Seminar on Complaint Handling and Management: From Receipt to Trending	David Dills	LIVE RECORDED	2018-06-26	$290.000 $540.000
501874	Food Fraud in the Organic Industry	John Ryan	LIVE RECORDED	2018-06-26	$150.000 $390.000
501990	How to Audit Against ICH GCP 2 Addendum to Ensure Compliance	Laura Brown	LIVE RECORDED	2018-06-27	$150.000 $390.000
501975	Validation of Analytical Methods and Procedures	Dr. Ludwig Huber	LIVE RECORDED	2018-06-27	$150.000 $390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501976	How to Implement Good Clinical Practices for Successful FDA and International Regulatory Authority Inspections	Angela Bazigos	LIVE RECORDED	2018-06-27	$150.000 $390.000
501995	3-Hour Virtual Seminar on Japan - Regulatory Filing Requirements and Compliance Processes for Life Sciences	Robert J. Russell	LIVE RECORDED	2018-06-27	$290.000 $540.000
501938	CAPA, Failure Investigation and Root Cause Analysis to Meet FDA Expectations	John E Lincoln	LIVE RECORDED	2018-06-27	$150.000 $390.000
501954	Secrets to Writing Effective SOPs for Medical Device QMS	Susanne Manz	LIVE RECORDED	2018-06-28	$150.000 $390.000
501996	Chemistry 101 for Medical Device Regulatory & Quality Professionals: Essential knowledge needed to Manage Drug&Device Combination Product Project	Robert Michalik	LIVE RECORDED	2018-06-28	$150.000 $390.000
501969	FDA Compliance and Mobile Applications	Carolyn Troiano	LIVE RECORDED	2018-06-29	$150.000 $390.000
501992	Edward Deming's Cure for a Sick System: Improving the Reliability of Organizations Processes	Michael Abitz	LIVE RECORDED	2018-06-29	$150.000 $390.000
501977	How to Implement Lean Documents and Lean Document Control to Cut Costs, while Maintaining Compliance with Regulatory Authority Requirements	Angela Bazigos	LIVE RECORDED	2018-06-29	$150.000 $390.000
501985	Beginning with the End in Mind: Developing a Regulatory Strategy for Product Development	Lauren Neighbours	LIVE RECORDED	2018-07-02	$150.000 $390.000
502002	How to Correctly Submit Pharmacogenomics Data for Faster Approval by the Agency, and What to Expect from the Regulatory Agencies in the Near Future	Angela Bazigos	LIVE RECORDED	2018-07-05	$150.000 $390.000
501998	Auditing Analytical Laboratories for FDA Compliance	Steven S. Kuwahara	LIVE RECORDED	2018-07-10	$150.000 $390.000
501993	Emergency Communications for Medical Facilities and First Responders	Michael Abitz	LIVE RECORDED	2018-07-11	$150.000 $390.000
501978	How to Deal with the Contradictions and Challenges of Using Cloud in a Regulated Environment	Angela Bazigos	LIVE RECORDED	2018-07-11	$150.000 $390.000
502006	3 hour Virtual seminar on FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	LIVE RECORDED	2018-07-12	$290.000 $540.000
501988	Combination Products Registration	Salma Michor	LIVE RECORDED	2018-07-12	$150.000 $390.000
502009	Similarities and Differences Between an FDA and MHRA Audit	Louis Angelucci	LIVE RECORDED	2018-07-12	$150.000 $390.000
502003	21 CFR Part 11 Compliance for Computer Systems Regulated by FDA	Carolyn Troiano	LIVE RECORDED	2018-07-13	$150.000 $390.000
501941	Japan - Regulatory Filing Requirements and Compliance Processes for Life Sciences	Robert J. Russell	LIVE RECORDED	2018-07-13	$150.000 $390.000

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501910	Bad SOPs, Bad Training: Garbage In, Garbage Out	Michael Esposito	LIVE RECORDED	2018-07-17	$150.000 $390.000
501965	Key Factors to Write an Effective Standard Operating Procedure (SOP) and Work Instructions (WIs)	Angela Bazigos	LIVE RECORDED	2018-07-17	$150.000 $390.000
501950	Process Validation Requirements & Compliance Strategies	Jose Mora	LIVE RECORDED	2018-07-17	$150.000 $390.000
501955	Non-Conforming Material and Failure Investigation	Susanne Manz	LIVE RECORDED	2018-07-18	$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
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
501951	Effective Design of Experiments (DOE) Strategies	Jose Mora	LIVE RECORDED	2018-07-26	$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
501959	Protocol Deviation Reporting and Management	Grace Morgan Holmes	LIVE RECORDED	2018-07-31	$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
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
501933	FDA's Revolutionary Change in Software Regulation	Casper Uldriks	LIVE RECORDED	2018-08-28	$150.000 $390.000
501980	Tools for Human Error Reduction	Ginette Collazo	LIVE RECORDED	2018-08-30	$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
501981	Controlling Human Error in the Manufacturing Floor	Ginette Collazo	LIVE RECORDED	2018-09-25	$150.000 $390.000
501982	Training in Human Error: Reducing Training Related Errors	Ginette Collazo	LIVE RECORDED	2018-10-24	$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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
500010	Foodservice Sanitation	Norman G. Marriott	RECORDED	2014-07-15	$149.000
500012	Foreign Corrupt Practices Act - Audits Role	Denise Cicchella	RECORDED	2014-05-27	$149.000
500013	Operational Risk - Beyond Regulatory Constraints	Fred Vacelet	RECORDED	2014-07-02	$149.000
500015	System Safety Engineering	Dev Raheja	RECORDED	2014-05-20	$149.000
500016	Fundamental ATC Organization, Structure, and Concepts	Stephen Alvania	RECORDED	2014-05-20	$149.000
500017	ATC Radar Approach Control (TRACON)	Stephen Alvania	RECORDED	2014-06-10	$149.000
500018	Mastering the Power of Sensitivity Tools in Financial Modeling	Miles Hutchinson	RECORDED	2014-08-28	$149.000
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
500038	The National Shipbuilding Procurement Strategy- Changing the playing field for shipbuilding in Canada	Anthony Goode	RECORDED	2014-07-17	$149.000
500039	The Business of Defence Procurement in Canada- A Market Over	Anthony Goode	RECORDED	2014-08-07	$149.000
500040	ISO Standard for Business Continuity (ISO 22301)	Ms. Michael Redmond	RECORDED	2014-09-16	$149.000
500041	Using Forensic Accountants Effectively in the Legal Process	Ray Kulzick	RECORDED	2014-08-07	$149.000
500042	Detecting Frauds in Purchasing and Payables	Ray Kulzick	RECORDED	2014-09-09	$149.000
500046	1099 and W-9 Update - Complying with IRS Information Reporting	Miles Hutchinson	RECORDED	2014-10-02	$149.000
500051	Controlled Goods in Canada- A primer	Anthony Goode	RECORDED	2014-08-27	$149.000
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	

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500079	FDA's Proposed Rule Regarding Device Establishment Registration and Listing and How to Register and List	David Dills	RECORDED	2014-11-24	$189.000
500080	FDA Compliant HPLC Qualification and Performance Testing	Dr. Ludwig Huber	RECORDED	2014-11-06	$189.000
500081	GMP Perspectives on Working with Contracting Laboratories	Steven S. Kuwahara	RECORDED	2014-11-11	$189.000
500082	Cleaning and Sanitation Training for Food Processors	Melinda Allen	RECORDED	2014-11-13	$189.000
500083	Auditing Analytical Laboratories for FDA Compliance	Steven S. Kuwahara	RECORDED	2014-11-18	$189.000
500084	Compiling the Design History File, and Technical File, Design Dossier	John E Lincoln	RECORDED	2014-11-18	$189.000
500086	FDA's 2011 Draft Guidance on Financial Disclosure by Clinical Investigators	David Dills	RECORDED	2014-11-19	$189.000
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

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500113	Device Corrections and Removals	Dan OLeary	RECORDED	2014-12-18	$189.000
500114	GMP Expectations for Products Used in Early Phase IND Studies	Steven S. Kuwahara	RECORDED	2015-01-06	$189.000
500115	IT Auditing - Principles and Practices (2nd Edition)	Robert E. Davis	RECORDED	2014-11-20	$149.000
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
500150	ATC Radar Approach Control (TRACON)	Stephen Alvania	RECORDED	2015-01-07	$149.000
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
500185	FDA Regulation of Dietary Supplements	Thomas E. Colonna	RECORDED	2015-02-19	$189.000
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 Dills	RECORDED	2015-02-24	$189.000
500207	Bullet Proof 510(k) – Latest FDA Changes to the Process	David 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 Dills	RECORDED	2015-03-24	$189.000
500209	Conducting Successful Product Complaint Investigations	David 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 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 Fruijtier	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 Fruijtier	RECORDED	2015-05-19	$189.000
500269	Scientific Advice in the EU	Adriaan Fruijtier	RECORDED	2015-05-27	$189.000
500270	Product Information in the EU	Adriaan Fruijtier	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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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 Dills	RECORDED	2015-07-23	$155.000
500394	Bullet Proof 510(k) – Latest FDA Changes to the Process	David Dills	RECORDED	2015-09-01	$158.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
500395	Draft Guidance for Device Industry and FDA - Postmarket Surveillance	David 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 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 Dills	RECORDED	2015-10-29	$190.000
500400	Conducting Successful Product Complaint Investigations	David 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 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 Fruijtier	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
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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
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500698	Introduction to FDA Regulation of Combination Products	Thomas E. Colonna	RECORDED	2016-03-31	$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
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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
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500914	Secrets to Writing Effective SOPs for Medical Device QMS	Susanne Manz	RECORDED	2016-07-06	$390.000
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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
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501026	Better Alternatives to AQL Sampling Plans for Risk Management in Incoming QC	John N. Zorich	RECORDED	2016-10-11	$390.000
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501049	Valid Statistical Rationales for Sample Sizes	John N. Zorich	RECORDED	2016-11-08	$390.000
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501053	Device Changes, FDA Changes, and the 510(k)	John E Lincoln	RECORDED	2017-01-25	$390.000
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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 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 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
501688	The New ISO 13485:2016 and Comparison with 21CFR820 - How	Eyal Lerner	RECORDED	2018-02-23	$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
	to Comply with both in the Same Organization				
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
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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
501797	Onboarding Employees in a GMP Environment: Best Practices for Foundational Employee Success	Michael Esposito	RECORDED	2018-04-30	$390.000

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
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
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
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501841	Computer System Validation (CSV) for FDA-Regulated Computers	Carolyn Troiano	RECORDED	2018-05-21	$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
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
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
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

Webinar Id	Webinar Name	Speaker Name	Webinar Type	Date	Price
501905	FDA's New Enforcement of 21 CFR Part 11	Dr. Ludwig Huber	RECORDED	2018-04-30	$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
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
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