Seminar Topic	Registration Type / Location / Price	Product ID
Ensuring Compliance with Advertising and Promotional Requirements for Drugs and Medical Devices: 2-day In-person Seminar	Seminar One Registration February 7-8, 2019, San Francisco, CA (Registrations till January 20, 2019 - $1299) (Registrations after January 20, 2019 - $1699) , Discounted Price: $1,299	80066SEM
REACH and RoHS Compliance: Gain a Deeper Understanding: 2-Day In-Person Seminar	Seminar One Registration February 7-8, 2019, San Diego, CA , Discounted Price: $1,699	80428SEM
Medical Device Single Audit Program [MDSAP] Implementation- Participating Country Regulatory Processes: U.S., Canada, Brazil, Australia and Japan: 2-Day In-Person Seminar	Seminar One Registration February 7-8, 2019, New Jersey, NJ , Discounted Price: $1,599	80493SEM
FDA`s Medical Device Software Regulation Strategy: 2-Day In-Person Seminar by Ex-FDA Official	Seminar One Registration February 28-March 01, 2019, San Francisco, CA (Registrations till January 28, 2019 - $1299) (Registrations After January 28, 2019 - $1499) , Discounted Price: $1,299	80242SEM
eCTD Submissions of IND and NDA/BLA to the US FDA, EU and Canada: 2-Day In-Person Seminar by Ex-FDA Official	Seminar One Registration March 12-13, 2019, San Francisco, CA (Registrations till February 8, 2019 - $1699) (Registrations after February 8, 2019 - $1899) , Discounted Price: $1,699	80480SEM
FDA`s New Import Program for 2019 - Strict Precision: 2-Day In-Person Seminar by Ex-FDA Official	Seminar One Registration March 14-15, 2019 , Orlando, FL (Registrations till February 5, 2019 - $1299) (Registrations After February 5, 2019 - $1499) , Discounted Price: $1,299	80133SEM
Latin America: Regulatory Compliance Requirements for Life Science Products (Focus: Brazil, Mexico, Argentina): 2-day In-person Seminar	Seminar One Registration March 21-22, 2019, Tampa, FL (Registrations till February 10, 2019 - $1299) (Registrations after February 10, 2019 - $1699)	80016SEM

Seminar Topic	Registration Type / Location / Price	Product ID
	, Discounted Price: $1,299	
HIPAA Omnibus Rule Compliance- Understanding Roles and Responsibilities of Privacy and Security Officers: One and Half day In-Person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80184SEM
Statistical Considerations for ICH Guidelines: 2-day In-person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80018SEM
Fundamentals of Antitrust and Competition Law: What you need to know to identify issues, mitigate risk and maximize flexibility: 2-day In-person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80230SEM
Schedule M-1/M-3 and Schedule UTP Reporting and Companies Requirements: 2-day In-person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80221SEM

Seminar Topic	Registration Type / Location / Price	Product ID
Key GMP Systems in Pharmaceutical and Biotech Labs: 2-Day In-person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80006SEM
Conducting Effective Investigation of Out-of-Specs and Atypical Laboratory Data: One and a Half day In-Person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80115SEM
Auditing the Human Resource Function-Human Resource Metrics: 2-day In-person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80035SEM
Lifecycle Approach to Analytical Methods for Drug Products: Incorporating QbD Concepts into Method Development, Validation, Verification and Transfer: 2-day In-person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80236SEM

Seminar Topic	Registration Type / Location / Price	Product ID
GMP Compliance for Quality Control and Contract Laboratories: 2-day In-person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80112SEM
Detecting and Preventing Embezzlement in Your Organization: One-Day In-Person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80292SEM
Ready, Aim, Hire!! Employee Screening and Background Checks for Newly Hired Employees: 2-Day In-Person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80261SEM
The Microbiological Element of Cleaning Validation in Oral Solid Dosage Manufacturing: 2- day In-person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80215SEM
Implementing UDI (Unique Device Identification) - Plan Now for	Get the Invitation	80303SEM

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Success: One-day In-person Seminar	Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	
OSHA 10 Hour Course: 2-Day In-Person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80282SEM
Human Factors and Design Controls for Medical Devices and Combination Products: 2-day In-person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80301SEM
Regulatory Compliance Auditing for pharma manufacturers: 2-Day In-person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80309SEM
APT Audit Process â€“ How Agile is Your Audit Process? Risk Based Auditing 2020: 2-Day In-Person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this	80254SEM

Seminar Topic	Registration Type / Location / Price	Product ID
	seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	
Clinical Research Project management for Life science industry - 2 day Workshop	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80306SEM
Implication of Global Med Device Regulatory Requirements to Total Product Lifecycle: Practical Approach - from Concept to Market and Beyond: 2-Day In-Person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80177SEM
Software Risk Management - Ways to FDA and MDD Compliance: 2-day In-person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80222SEM
How to Prepare for, Manage, and Follow-up to an FDA Inspection: Risk-driven Approach: One and a Half day In-Person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or	80159SEM

Seminar Topic	Registration Type / Location / Price	Product ID
	email at editor@grcseminars.com	
Regulatory Crisis Management: Best Practices for Dealing with the Common Crisis Events for the FDA-Regulated Industry: 2-Day In-Person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80166SEM
Process validation and process defect prevention to increase device reliability: 2-Day In-Person Seminar	Get the Invitation Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com	80276SEM

