| Seminar Topic | Registration Type / Location / Price | Product ID |
|---|---|---|
| Managing Your Complaints and Obstacles in Post-Market Requirements- Results from Top Medical Device Observations During an Inspection: 2-day In-Person Seminar by Ex-FDA Official | Seminar One Registration<br>September 12-13, 2019, San Diego, CA<br>(Registrations till August 3, 2019 - $1499)<br>(Registrations after August 3, 2019 - $1699) , Discounted Price: $1,499 | 80068SEM |
| Risk Management in Medical Devices Industry: 2-Day In-Person Seminar | Seminar One Registration<br>September 26-27, 2019, Tampa, FL<br>(For Registrations till August 27, 2019 - $1499)<br>(For Registrations after August 27, 2019 - $1899) , Discounted Price: $1,499 | 80490SEM |
| Navigating through Maze of In-vitro Diagnostics (IVD) Regulations: A systematic approach from Regulatory Strategy to Regulatory Approvals in U.S./Europe/Canada: 2-day In-Person Seminar | Seminar One Registration<br>October 14-15, 2019, Boston, MA<br>(For Registrations till August 30, 2019 - $1699)<br>(For Registrations after August 30, 2019 - $1899) , Discounted Price: $1,699 | 80148SEM |
| FDA`s Medical Device Software Regulation Strategy: 2-Day In-Person Seminar by Ex-FDA Official | Seminar One Registration<br>November 7-8, 2019, Philadelphia, PA<br>(Registrations till August 21, 2019 - $1299)<br>(Registrations after August 21, 2019 - $1499) , Discounted Price: $1,299 | 80242SEM |
| Medical Device - QSR Compliant Product Development Process: One and half day In-person Seminar | Get the Invitation<br>Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar.<br>Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com | 80136SEM |
| Medical Device Software Risk Management, Cybersecurity and Assurance Case: One and a Half Day In-Person Seminar | Get the Invitation<br>Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar. | 80169SEM |

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| | Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com | |
| Understanding FDA QSR for Med Devices and Laser Product Performance Standards for Medical and Industrial Radiation Emitting Devices: 2-day In-Person Seminar | Get the Invitation<br>Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar.<br>Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com | 80188SEM |
| CAPA and its Strategy: Root Cause Analysis: 2-day In-person Seminar | Get the Invitation<br>Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar.<br>Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com | 80209SEM |
| Effective Complaint Handling, Medical Device Reporting and Recalls and Avoiding Costly Errors: One and a Half-day In-Person Seminar | Get the Invitation<br>Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar.<br>Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com | 80187SEM |
| Medical Device Risk Management A to Z - Best Practices for Effectiveness and Efficiency: 2-day In-person Seminar | Get the Invitation<br>Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar.<br>Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com | 80213SEM |

  

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| Effective and Efficient Internal and Supplier Quality System Auditing for Medical Devices: 2-day In-person Seminar | Get the Invitation<br>Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar.<br>Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com | 80235SEM |
| FDA`s Regulation of Medical Devices and Strategies for their Successful Import, Marketing and Sale in the United States: 2-day In-Person Seminar | Get the Invitation<br>Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar.<br>Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com | 80179SEM |
| Human Factors and Design Controls for Medical Devices and Combination Products: 2-day In-person Seminar | Get the Invitation<br>Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar.<br>Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com | 80301SEM |
| The CE Mark: Understanding the Medical Device Directives: One and a Half Day In-Person Seminar | Get the Invitation<br>Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar.<br>Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com | 80214SEM |

  

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| 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<br>Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar.<br>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<br>Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar.<br>Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com | 80222SEM |
| Process validation and process defect prevention to increase device reliability: 2-Day In-Person Seminar | Get the Invitation<br>Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar.<br>Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com | 80276SEM |
| Understanding and Implementing the Medical Device Directive: 2-Day In-person Seminar | Get the Invitation<br>Pre-Register yourself and get the official Invite when venue and dates are announced for this seminar.<br>Call here to register +1-1-888-771-6965 or email at editor@grcseminars.com | 80119SEM |

  