Global CompliancePanel

Netzealous
Knowledge, a Way Forward…

2-day In-person Seminar:

Regulatory Requirements and Principles for Cleaning Validation

San Diego, CA

December 7th & 8th, 2017

9:00 AM to 6:00 PM

Price

Price: **$1,495.00**

(Seminar for One Delegate)

Register for 5 attendees

Price: **$4,485.00** You Save: $2,990.0 (40%)*

~~$7,475.00~~

Register for 10 attendees

Price: **$8,222.00** You Save: $6,728.0 (45%)*

~~$14,950.00~~

ENROLL

***Please note the registration will be closed 2 days (48 Hours) prior to the date of the seminar.*

Joy McElroy

Principle Consultant, Maynard Consulting Company

Upon earning a degree in Zoology at North Carolina State University, Joy made her debut in the pharmaceutical industry in 1992 at Pharmacia & Up John performing Environmental Monitoring and Sterility Testing. Her hard work allowed her to move into a supervisory role at Abbott Laboratories where she oversaw their Quality Control Lab.

Now with 12 years experience as a consultant, and over 20 years total experience in the pharmaceutical and biotech industries, Joy has gained extensive knowledge of Quality Assurance, Process and Cleaning Validation, and Equipment Qualification. She has written and executed Equipment Qualification and Validation Protocols for numerous Companies such as Mallinckrodt, Wyeth Lederle, Merck, BioMerieux, Catalent, Phillips Medisize, Xcelience, and Novartis.

Overview :

This 2 day course will cover practical guidance on cleaning validation regulatory compliance, in conjunction with, risk-based, reasonable and informed decision making and activity planning. This two day interactive course will cover fundamental principles of a cleaning validation program, exploring such concepts as the determination of residues to be targeted, selection of analytical and sampling methods, determination of appropriate limits in various pharmaceutical and biotechnology processes, and establishment of scientific rationales acceptable to regulatory inspectors.

Agenda:

Day One

Lecture 1: FDA Requirements and Industry Standard Practices

Lecture 2: How to Develop/Review your Cleaning Procedures and the Adequate Selection of Cleaning Agents and Parameters

Lecture 3: How to Develop a Cleaning Validation Policy/Program

Lecture 4: How to Implement a Robust Cleaning Validation Plan

Why you should attend:

Attendance at 2 day seminar will be beneficial to personnel directly involved in the development of cleaning procedures, cleaning validation programs and plans. Additionally, those responsible for cleaning validation protocols and execution activities, including validation and laboratory personnel, as well as, beginning or seasoned operational personnel who will eventually participate in such efforts, will find this course particularly useful. This includes Analytical Method Development, Quality Control and Quality Assurance personnel.

Areas Covered in the Session:

- Understand the importance and underlying principles of cleaning validation and the requirements to have adequate cleaning procedures for manufacturing equipment in contact with the product
- Understand the FDA perspectives on cleaning validation and areas of concern during regulatory inspections
- Be able to set up cleaning validation procedures, protocols and reports that meet current FDA, WHO, PIC/S and EU regulations
- Prepare and defend your own cleaning validation approach/program and avoid costly delays and/or rejections by regulatory agencies

Day Two

Lecture 1: Laboratory Issues in Cleaning

Lecture 2: Microbiological aspects of a cleaning validation program for manufacturing equipment

Lecture 3: Keys to Cleaning Validation Maintenance - Remaining Compliant

Lecture 4: Current FDA concerns about validation of cleaning processes

Regulatory Requirements and Principles for Cleaning Validation

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2-day In-person Seminar:

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Group Participation

10%	2 Attendees to get offer
20%	3 to 6 Attendees to get offer
25%	7 to 10 Attendees to get offer
30%	10+ Attendees to get offer

Payment Option

1. Credit Card: Use the Link to make Payment by Visa/Master/American Express card click on the register now link
2. Check: Kindly make the check payable to NetZealous DBA GlobalCompliancePanel and mailed to 161 Mission Falls Lane, Suite 216, Fremont, CA 94539, USA
3. PO: Please drop an email to support@globalcompliancepanel.com or call the our toll free +1-800-447-9407 for the invoice and you may fax the PO to 302 288 6884
4. Wire Transfer: Please drop an email to support@globalcompliancepanel.com or call our toll free +1-800-447-9407 for the wire transfer information

What You will get

1. Learning Objectives
2. Participation certificates
3. Interactive sessions with the US expert
4. Post event email assistance to your queries.
5. Special price on future purchase of web based trainings.
6. Special price on future consulting or expertise services.
7. Special price on future seminars by GlobalCompliancePanel.
8. Seminar Kit – includes presentation handout, ID card, brochure, trainings catalog, notepad and pen.
9. Networking with industry's top notch professionals

Contact Information: Event Coordinator

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Kindly get in touch with us for any help or information.

Look forward to meeting you at the seminar

GlobalCompliancePanel