




2-day In-person Seminar:

Supplier Management for Medical Device Manufacturers

-  Washington, DC
-  March 23rd & 24th, 2017
-  9:00 AM to 6:00 PM



Betty Lane

Founder and President,
Be Quality Associates, LLC

Betty Lane has over 30 years' experience in

Medical Device quality assurance and regulatory affairs. She is the founder and President of Be Quality Associates, LLC, a consulting company helping small and medium sized medical device and diagnostic companies implement and improve their quality systems. Her work enables companies to manage their business in compliance with FDA and ISO 13485 requirements, as well for quality system requirements for other geographic area such as Europe and Canada. Her background in digital systems engineering enables her to facilitate quality system processes for design controls and software validation. Her areas of expertise include training, auditing, supplier management, document and records management, design controls, and software validation.

Overview :

Supplier selection and management is one of the critical issues for medical device manufacturers. Suppliers provide materials and services to the device manufacturer, which means that they can be critical to performance and delivery of your device. Neither the FDA nor your notified body regulates your suppliers (with a few exceptions). They expect you to have an effective process to ensure your suppliers perform in the regulatory environment.

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~~

ENROLL

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



Agenda:

Day One

Lecture 1: Introductions

Lecture 2: Fundamentals Regulatory Requirements

- FDA Requirements
 - ISO 13485 requirements
 - Understanding the role of the Global Harmonization Task Force Guideline
 - Understanding NBOC Guideline and why it should be used
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Lecture 3: Planning the Supplier Management Program

- Supplier Classification
 - Supplier QA agreements what are they and why are then
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Who will benefit:

- Quality Managers
- Quality Engineers
- Audit Managers
- Supplier Engineers
- Internal quality auditors
- Supplier auditors
- Quality associates
- Quality Specialists
- Regulatory Compliance Managers

Day Two

Lecture 1: Planning Supplier Selection

Lecture 2: Potential Suppliers

Lecture 3: Supplier Selection

Lecture 4: Implementing Supplier Controls

Lecture 5: Monitoring, Measuring, and Evaluation

- Periodic Monitoring
 - Re-evaluations
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Lecture 6: Supplier Audits - where do they add value

- Planning your supplier audit schedule
 - How Notified Body unannounced audits affect your contract manufacturer
 - What you should do to prepare yourself and your contract manufacturer for unannounced Notified body audits
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Lecture 7: Feedback and Communication

- Supplier meetings: Partnering with Key suppliers
 - Supplier Corrective Actions
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Lecture 8: Evaluating your current program to see how it measures up to regulatory Expectations

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
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What You will get

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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