Supplier Management for Device makers and Drugmakers: Qualification, Contracts and Audits

2 Day Virtual Seminar

Overview

This 2-day course will cover managing a supplier for the entire lifecycle of the relationship, beginning with identification and qualification of a supplier and continuing through building a relationship, risk management, ongoing assessment (including auditing) and finally planning for an exit. The course will show attendees how to use risk assessment for ranking suppliers and reducing the number of audits that are necessary to effectively manage suppliers. Strategies for determining whether a supplier will be sole source will be included

Who Will Benefit

- Quality Assurance Managers,
- Quality Control Supervisors,
- Quality Engineers,
- Procurement Professionals,
- Drug Development Scientists,
- Medical Device R&D Engineers
- Supply Chain Managers

8.0 RAC CREDITS

Agenda

- The benefits and components of a supplier management program
- Regulatory requirements for managing suppliers and contract manufacturers
- Strategic decision making for good supplier management
- How to manage risk and reduce the costs associated with having suppliers
- The steps involved in selecting and onboarding a supplier
- Developing good supplier relationships including managing improvement and nonconforming events
- Writing effective and useful quality agreements
- Reviewing supplier performance and making performance-based decisions
- How to perform a desktop assessment and a supplier audit (and when to use each)
- Managing supplier transitions

Faculty José Mora

Jose Mora is a Principal Consultant specializing in Manufacturing Engineering and Quality Systems. For over 30 years he has worked in the medical device industry specializing in manufacturing, process development, tooling, and quality systems. Prior to working full time as a consulting partner for Atzari Consulting, José served as Director of Manufacturing Engineering at Boston Scientific and as Quality Systems Manager at Stryker Orthopedics, where he introduced process performance, problem solving, and quality system methodologies

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REGISTRATION FORM

Name **Email**

Organization **Department** **Position**

Phone **Mobile**

Register Online At www.worldcomplianceseminars.com

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Terms And Condition

- Registration shall be reconfirmed only once payment has been made prior to the course.
- Cancellation of participant/s shall be submitted in writing to WCS ten days (10 days) before the course. There is a 50% liability on all bookings once made.
- WCS reserves the right to make alterations to the conference/executive briefing content, timing, speakers without notice. The event may be postponed or cancelled due to unforeseen events beyond the control of WCS. If such a situation arises, we will refund your registration fee or we will try to reschedule the event.