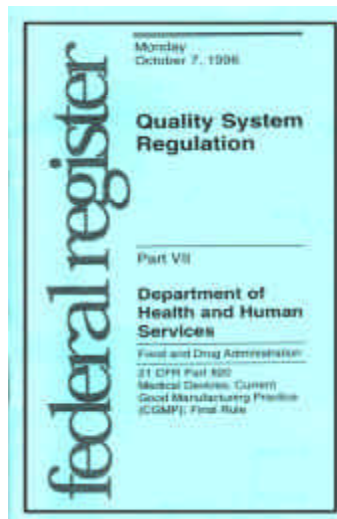


QARACC News

FDA QSR TRAINING



*An In-Depth 2-Day Seminar for:
Executives, Managers, Engineers,
Quality and Regulatory Professionals,
Auditors, Documentation Specialists,
and Compliance Specialists*

Date: February 23rd and 24th, 2004

Time: 8:00 AM to 4:30 PM

COURSE EXPECTATIONS

Manufacturers of medical devices are mandated to implement and comply with the FDA's Quality System Regulation (QSR) codified under 21 CFR Part 820. This two-day course will provide medical professionals with an in-depth knowledge and perspective of the FDA's QSR and enable personnel to implement an FDA-compliant quality management system. The course provides an in-depth review of the QSR subparts in easy to understand language. It is ideal for those individuals responsible for complying with the regulations as well as those individuals involved in the development, implementation and maintenance of the organizations quality management system.

Highlights

- ◆ **CGMP/QSR History**
- ◆ **Organizational Benefits of Compliance**
- ◆ **Key Terms & Definitions**
- ◆ **Outline FDA's Quality System Inspection Technique (QSIT)**
- ◆ **In-Depth QSR Subpart Review**
- ◆ **General Auditor Questions**
- ◆ **Interactive discussion**
- ◆ **Real World Examples**
- ◆ **Tips for Compliance**
- ◆ **Participant Manual**

REGISTER NOW!

SAVE \$100 off the \$895 registration fee by registering before 12/31/03

CONTACT

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Odessa, FL 33556

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Email: info@qaracc.com
Website: <http://www.qaracc.com>



The seminar will be held at the:
Wyndham Harbour Island Hotel
725 South Harbour Island Blvd
Tampa, FL 33602-5707
Tel: (813) 229-5000 Fax: (813) 229-5322
Web: www.wyndhamharbourisland.com

Call early for the discount rate!