



2010 Course Locations and Dates

Sept. 20-22, 2010
Wilmington, Delaware
Hotel duPont
11th & Market Streets
Wilmington, DE 19801

Registration Form:

Name: _____
Please print First LAST

Title: _____

Organization: _____

Mailing Address: _____

City: _____

State: _____ **Zip/Postal Code:** _____

Country: _____

Telephone: () _____

Fax: () _____

E-mail: _____

How did you hear about EduQuest? _____

Method of Payment:

_____ Check Payable to EduQuest, Inc.

_____ Company Purchase Order

Please bill company PO# _____

_____ Credit Card (**Circle One – MC / VISA / AmEx**)

Card# _____

Exp. Date _____ Security Code _____

Name on credit card & billing address

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Signature (Required) _____

****Two courses to choose from!****
For the best price and value, choose the
EduTrack
or select individual courses.

The EduTrack - \$2795.00 USD

All 3 Days of Training:
*QSR Compliance Fundamentals, and
Design Control for Medical Devices*

QSR Compliance Fundamentals:

Complying with FDA's Medical Device 21 CFR 820
Quality System Regulation
\$990. USD
8:30 AM - 5:30 PM

_____ **Wilmington, Delaware September 20, 2010**

Design Control for Medical Devices:

Meeting FDA's 21 CFR 820.30 Rules
for Quality Design and Manufacturing
\$1980. USD
8:30 AM - 5:30 PM

_____ **Wilmington, Delaware September 21-22, 2010**

Multiple Registrations:

Group rate for attendees from same company at same location and courses. Call for price, 301-874-6031

Method of Payment:

Please note that payment is required in advance prior to course date. Please make checks (in US funds drawn on US bank) payable to EduQuest, Inc. Confirmation of your registration will be sent. Full payment must accompany registration form.

Cancellations/Substitutions:

Company substitutions are not permitted without prior notification to EduQuest. Cancellations received before the beginning of a course will be subject to a refund according to the following schedule and rates. Up to 6 PM EST, 10 business days in advance of the course, a 95% refund will be provided. If less than 10 business days advance notice is provided, the refund amount will be 50%. Individuals requesting to change course location will be charged a \$500 course transfer fee if less than 10 business days before course. No-shows will be charged the full amount. EduQuest reserves the right to cancel the courses and is not responsible for any airfare, hotel, or other costs incurred by registrants.



For registration or additional information,
please contact us at:

Mail:
EduQuest, Inc.
1896 Urbana Pike, Suite 14
Hyattstown, MD 20871 USA

Call: 301.874.6031
Fax: 301.874.6033
E-mail: Info@EduQuest.net
Website: www.EduQuest.net



EDUcation: QUality Engineering, Science and Technology

**Learn straight from the source – former FDA inspectors, rulemakers, and trainers
from the global consulting team of EduQuest**

Dear Industry Colleague,

The scope of FDA's Quality Systems Regulation (QSR) is huge – more than 500 pages of rules and guidance. Keeping up with all these regulations is quite a challenge. Just within the past 12 months, 27 medical device companies received Warning Letters for deficient quality management programs, and several were threatened with removal of their products from the market.

That's why I'm inviting you to attend these two new courses – offered separately or as an integrated three-day learning package:

QSR Compliance Fundamentals:

Complying with FDA's Medical Device 21 CFR 820 Quality System Regulation

March 17, 2010 • Las Vegas, NV
September 20, 2010 • Wilmington, DE

Design Control for Medical Devices:

Meeting FDA's 21 CFR 820.30 Rules for Quality Design and Manufacturing

March 18-19, 2010 • Las Vegas, NV
September 21-22, 2010 • Wilmington, DE

The goal of both training programs is to give you and your staff real-world, step-by-step compliance information. Through plain-English instruction, detailed course materials, and interactive exercises, **you learn to cost-effectively comply with FDA's QSR rules and related international standards.**

Please feel free to call us with questions at 301-874-6031 or email us at Info@EduQuest.net. To register, use Registration Form on reverse side or visit our website www.EduQuest.net. We look forward to seeing you at our training courses soon!

Sincerely,

Martin Browning

Martin Browning
President, EduQuest

[Former FDA Investigator, National Computer and Medical Device Expert, Special Assistant to the Associate Commissioner for Regulatory Affairs, Vice Chair of FDA's Electronic Record and Signature Working Group, and Chair of the U.S. Government ISO-9000 Committee]

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