



**2010 Course Locations and Dates**

**Sept. 13-17, 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**

Cardholder's Name: \_\_\_\_\_

Address: \_\_\_\_\_

City/State/Zip: \_\_\_\_\_

Signature (Required) \_\_\_\_\_

**\*\*Five courses to choose from!\*\***

For the best price and value, choose the **EduTracks** or select individual courses.

**The EduTrack - \$3980.00 USD**

**Track A**

*FDA Auditing of Computerized Systems & Part 11, Introduction to Risk Management for FDA Compliance, and Process Risk Management Explained*

**Track B**

*FDA Auditing of Computerized Systems & Part 11, Introduction to Risk Management for FDA Compliance, and Effective and Compliant CAPA Systems*

**FDA Auditing of Computerized Systems and Part 11**

\$2795. USD

*Monday-Wednesday - 8:30 AM - 5:30 PM*

**Introduction to Risk Management for FDA Compliance**

\$495. USD

*Thursday, 8:30 AM - 12:30 PM*

**Effective and Compliant CAPA Systems**

\$990. USD

*Thursday -1:30 PM - 5:30 PM*

*Friday - 8:30 AM - 12:30 PM*

**Process Risk Management Explained**

\$990. USD

*Thursday, 1:30 PM - 5:30 PM*

*Friday, 8:30 AM - 12:30 pm*

**Dangerous Documents**

\$495. USD

*Thursday, 8:30 AM - 12:30 PM*

**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

## ***We Trained FDA and are Now Offering Our Courses to the Industry!***

Dear Industry Colleague,

FDA is continuing to update its approach to GMP, quality systems, 21 CFR Part 11, risk management, and the industry's use of new process analytical and control technologies. At the same time, the industry is increasing its reliance on software, computerized systems, and other advanced technologies to support and streamline manufacturing, clinical, laboratory, and other regulated operations. Because quality system, validation, and computerized system issues continue to be sources of controversy and frequently cited deviations during FDA inspections, it is critically important for today's technology-dependent companies to understand and prepare for FDA's shifting priorities and inspectional scrutiny in these areas.

Our most popular training courses focus specifically on these issues, and we are now offering more courses in parallel tracks to allow you to choose the training that best fits your company's needs. We invite you to take advantage of this unique opportunity to learn from the same expert team that trained FDA.

We have enclosed information regarding our 2009 training course dates and locations.

**Please note our upcoming courses in Las Vegas, Nevada (March 22-26, 2010) and Wilmington, Delaware (September 13-17, 2010).**

Please feel free to call us with questions at 301-874-6031 or e-mail us at [Info@EduQuest.net](mailto:Info@EduQuest.net).

To register, use Registration Form on reverse side or visit our website [www.EduQuest.net](http://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]*

1896 Urbana Pike, Suite 14, Hyattstown, MD 20871, (Ph) 301-874-6031, (Fax) 301-874-6033,  
[Info@EduQuest.net](mailto:Info@EduQuest.net) \* \* \* \* [www.EduQuest.net](http://www.EduQuest.net)