COMPLIANCE associates

VALIDATOR

Validator is the only automated compliance solution available. Validator significantly reduces validation time. **Results guaranteed.**

Book a demo

Greetings from Compliance Associates! We have exciting new validation and compliance resources to share including recently conducted industry research, and computer system validation training.

And if you haven't already, be sure to check out our video for a quick overview of our business.

Vaughan Chamber of Commerce Award Nomination

We've been nominated for the Vaughan Chamber of Commerce Innovation Award, by Deloitte. We are among the top 3 out of 11 candidates - stay tuned for the results!

Learn more about the award

Compliance and Validation Research

We recently conducted research that focused on understanding pain points, trends and purchasing behaviours in computer system validation and compliance in the pharmaceutical industry in North America.

Contact us for an exclusive opportunity to review the research and gain insights and learnings that will help drive successful validation strategy and planning for your business.

Computer System Validation: How Much to Validate?

How much to validate is the biggest challenge facing validation professionals and their organizations. Deciding how much to validate is a delicate balance between satisfying compliance requirements and the time it takes to

Training Course: Computer System Validation Fundamentals

During this training we will teach basic validation principals that will help companies establish a solid foundation for delivering technology solutions in FDA and Health Canada regulated environments.

Date: April 11, 2014
Time: 9am-4pm
Location: 8000 Jane Street, Concord ON L4K 5B8
Duration: 1 day
Cost: $800 per student
Group Pricing: Enroll two students at full price and receive a third seat FREE.

implement critical technologies.

Read more

In order to help our customers get their questions answered quickly, we've put together a list of frequently asked questions.

Visit our FAQ page

Read more and register

Quality by Design and Process Validation

The FDA has made significant progress in implementing the concepts of "Quality by Design" (QbD) into pre-market processes. The QbD concept is that quality should be built into a product through a deep understanding of the product, both how it's developed and how it's manufactured.

The QbD initiative forms the basis for process validation activities and attempts to provide guidance on pharmaceutical development to facilitate design of products and processes with maximum efficacy, safety profile and product manufacturability.

Contact us to speak with a process validation expert

Customer Success Story: Updating Existing Software to Optimize Regulatory Infrastructure and Automate the Validation Process

Manufacturers like KIK Custom Products use many types of software throughout their organizations, all of which must be validated in order to meet regulations set forth by the FDA and Health Canada. The IT Team and Quality Group at KIK were looking for a software package capable of automating the validation life-cycle.

Read more

Your partner for regulatory compliance with over 25 years of experience.

Consult with a validation expert today.

Stay connected!