



EtQ  
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EtQ is the leading Enterprise Quality and Compliance Management Software for identifying, mitigating and preventing high-risk events through integration, automation and collaboration.

EtQ's mission is to make compliance management and business communication easy and efficient. Founded in 1992, EtQ has always had a unique knowledge of quality and compliance processes, and has always strived to focus on making overall quality operations and quality systems better for businesses.

EtQ Pharmaceuticals/Biotechnology is an integrated FDA Compliance Management System that has been pre-configured to specifically address the needs of the Pharmaceuticals/Biotechnology industry, and exceed 21 CFR part 11 requirements. EtQ's unique modular approach provides unparalleled flexibility and automation.

As required by 21 CFR Part 11, a secure environment must be maintained, and the system must be validated to maintain electronic signature integrity. As a result, validation services can double, or even triple the cost of a software system. EtQ's web-based system is designed specifically for the Pharmaceutical industries, and our staff has developed the test scripts necessary to ease system validation. Because of the EtQ software is usable right out of the box and configurations are executed at the server level, the need for extensive validation across the enterprise is less than comparative systems in the market.

EtQ uses best of breed-integrated modules and enterprise applications to manage the process, provide business intelligence visibility and execute organizational change. Key modules include: Corrective Action/Preventive Action, Complaint Handling, Nonconformance, Document Control, Risk Assessment, Training, Reporting and more.

EtQ's unique FDA Compliance Software System is designed to minimize the number of CAPAs using an advanced filtering model; which features automatic segregation and categorization of events at the source, automatic identification and display of related events, built-in risk assessment software module, initial assessment to allow early closure, risk assessment throughout the process to guide decision making, full investigation with step-by-step root cause analysis, automatic lookup and display of related investigations and CAPAs, comprehensive CAPA action and effectiveness check plan with risk mitigation history.

Large companies with multiple divisions typically have independent, hybrid systems in place. Bridging the gap between these systems is inefficient in the long-term and ultimately will require expensive customizations to link the enterprise. Re-routing documents, meeting to discuss integration strategy, and IT man hours in customizations result in lost time and money. EtQ's FDA Compliance Software is designed to incorporate the whole enterprise in collaborating at the enterprise level, while maintaining each division's independent business workflow. The result is a unified Quality Management system that allows for each division to collaborate with the enterprise.

As more companies rely on Contract Manufacturers to help bring their products to the market, it is

important that the systems implemented allow for increased visibility downstream. EtQ provides integration at multiple levels of the process, allowing contract manufacturers to take part in the overall business process, without direct access to the main system. This allows the Contract Manufacturers to fill in their pieces of the process, and provide a seamless quality system, from start to finish.