Medvacon Promulgates its Laboratory Information Management Systems (LIMS) Services

December 23, 2020 - (SPARTA, NJ) - When electronic records need to be validated, you can turn to Medvacon Life Sciences—a Quality, Compliance and Validation consulting company. And when it comes to Laboratory Information Management Systems (LIMS) in the biotech, pharmaceutical or medical device industries, Medvacon offers validation support in every phase of development from pre-clinical to manufacturing.

LIMS is s a software-based solution with features including sample management, workflow and data tracking support, flexible architecture, and data exchange interfaces in a regulated environment. The features and uses of a LIMS have evolved over the years from simple sample tracking to a fully integrated enterprise tool that manages multiple aspects of laboratory data from multiple instrument sources. LIMSs are dynamic because the laboratory's requirements are rapidly evolving and different labs often have different needs.

“Much of the software out there requires significant configuration for each customer’s needs,” explains Joe Toscano, a founding member of Medvacon Life Sciences, “and it can be specialized to communicate with several instruments within a life sciences organization. That’s where Medvacon comes in. Even if your software has specific user requirements and configurations,” he continued, “we can work with you to customize your LIMS validation plan and execution.”

According to Toscano, most LIMS are cloud-based today, and Medvacon has the experience to validate your laboratory information management system and other electronic record systems, such as document control, SOPs, and Quality Management Systems. The consulting firm offers experts in optimizing and streamlining validation master plans and protocols for LIMS.

Medvacon is a full-service Quality and Compliance consulting firm offering a suite of quality services such as: Quality Systems Implementation and Remediation; Computer System Validation; Process and Cleaning Validation; Equipment, Facility and Utility Qualification; Mock FDA Inspections and Audits; Project Management; and cGMP Training. These services help clients to ensure FDA Compliance and reduce compliance cost in the Life Sciences’ industry. The company provides experienced project administration, trouble-shooting, and/or training services, on-site, with local resources and senior management involvement to optimize flow and collaboration. Headquartered in Sparta, NJ, Medvacon Life Sciences also has offices in San Diego California. More information can be found at www.Medvacon.com.

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