



LIQUENT Announces XEVMPD Educational Series and Data Assessment Tool

Horsham, PA – January 13, 2012: LIQUENT, Inc. the global leader in regulatory information management solutions and services, is pleased to announce LIQUENT's XEVMPD Educational Series and Data Assessment tool. Both are designed to assist in complying with the July 2012 deadline for submitting product information in extended EudraVigilance Medicinal Product Dictionary (XEVMPD) format.

LIQUENT's XEVMPD Educational Series is designed to educate internal personnel about the requirements for XEVMPD. The sessions consist of five recorded modules, for a total of four hours of training, focused on areas such as the understanding of why EMA is requiring XEVMPD, recent announcements and timelines, and requirements for authorized products, approved substances, and development substances and products.

The XEVMPD Data Assessment tool will assist organizations in determining what data is needed for compliance and where the data is currently managed or tracked within the organization.

"Companies are facing an unprecedented timeline for compliance," said Sarah Powell, Executive Director of Regulatory Affairs and Writing Services. "The LIQUENT XEVMPD Educational Series and XEVMPD Data Assessment tool provide an opportunity for companies to get a jump start on preparing their organizations, by ensuring employees are educated about the requirements and providing a clear understanding of the data needs."

Successful transition to comply with the new XEVMPD requirements will mandate companies to evaluate standards, processes, technology, and organizational needs. LIQUENT's educational and solution offerings will support companies as they make necessary assessments.

For more information, Please contact info@liquent.com.

About LIQUENT, Inc.

LIQUENT provides technology and outsourcing solutions focused around regulatory submission preparation and tasks, dossier planning, eCTD to CTD publishing and registration tracking capabilities. LIQUENT is the premier provider of a scalable, regulatory information management platform and associated regulatory & clinical services that can be leveraged throughout a client organization.