ACUTA announce the release of ACUTA Regulatory Information Management (ARIM™) version 2.2.0

Marlborough MA – January 30, 2015 – ACUTA announced the release of ACUTA Regulatory Information Management (ARIM) version 2.2.0 – a next-generation solution that redefines how life sciences companies manage regulatory information. ARIM brings together the data, content, and interactions with agencies needed to deliver a fully informed Regulatory Information Management (RIM) solution. Designed to work in the cloud or on premise, ARIM provides a centrally coordinated user experience while ensuring compliance. It supports Regulatory Affairs and Operations processes regardless of their complexity or geographic location, and it offers global access.

"ACUTA's goal is to offer this next generation RIM solution to enable companies of all sizes to adhere to strict standards set by regulatory authorities around the world and help them collect and manage regulatory information with reliability" said Shy Kumar, ACUTA's Founder, President and CEO. "With this release we are very proud to announce that support for the upcoming FDA v3.3 DTD is available to ARIM users now! With this release, ARIM users have the opportunity to plan and prepare to deliver the submissions using the new DTD. Also, this release offers the capability to publish Non-eCTD Electronic Submissions (NeES), which is another critical requirement to support rest of the world (ROW) submissions."

ARIM is designed to assist Life Science companies in managing their regulatory information more efficiently and cost effectively. This v2.2.0 release includes a significant list of new capabilities along with improvements to existing features and functionalities.

The ARIM Cloud offering is powered by Microsoft Azure and advances the life science industry's urgent drive for an affordable solution using technology tailored to small and medium size life science company's needs, with increased efficiencies and cost savings.

For more information on ARIM, please visit www.acutallc.com/arim.html

About ACUTA, LLC

ACUTA was founded in November 2012 to assist life science companies in complying with ever changing regulatory requirements that guide the product lifecycle through approval and maintenance. ACUTA's founder Shy Kumar, and his team members are well known in the industry, with over 20 years of experience. They have successfully deployed state-of-the-art solutions that were well received by industry and users.

ACUTA's vision is to develop innovative solutions to assist life sciences and related companies with their regulatory information management, which ultimately benefit everyone and specifically the needs of patients around the world.

ACUTA is headquartered in the Marlborough, MA, with offices in Bangalore India. For more information, visit www.acutallc.com.

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