ACUTA announces the release of ACUTA Regulatory Information Management (ARIM™) v2.3.3 and ACUTA Validator v2.3.3

Marlborough MA – May 5, 2016 – ACUTA announced the release of ACUTA Regulatory Information Management (ARIM) version 2.3.3, a next-generation solution that redefines how life science companies manage regulatory information, and ACUTA Validator v2.3.3, a desktop based application to support stand-alone validation of eCTD and NeeS submissions.

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 in a timely manner 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. "This release is focused on providing support for the upcoming EU regional DTD v3.0 requirements, that go into effect from July 1, 2016 and will become a requirement from October 1, 2016. "

ARIM is designed to assist Life Science companies in managing their regulatory information more efficiently and cost effectively. This v2.3.3 release includes a significant list of new capabilities along with improvements to existing features and functionalities.

ACUTA Validator v2.3.3 is designed to support validation of eCTD and NeeS submissions covering multiple agency requirements around the globe.

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 science and related companies with their regulatory information management, which ultimately benefits everyone and specifically the needs of patients around the world.

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

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