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For Immediate Release

Datafarm advances regulatory submission authoring with the release of S-Cubed[®] Templates version 2.0

-Microsoft[®] Word[®] 2007 support is one of the various major enhancements to the latest release of S-Cubed Templates-

Marlborough, MA (PRWEB) November 17, 2009 — Datafarm Inc., a world leading provider of electronic regulatory submission solutions for the Life Sciences industry, today announced that S-Cubed Templates version 2.0 offers users major feature enhancements as well as new features including support for Microsoft Word 2007. Regulatory submission document authors have relied on S-Cubed Templates since 2006 to reduce the time manual formatting requires while enabling compliant, multi-agency documents.

“Document authoring is a critical part of the drug development and regulatory approval process,” said Shy Kumar, Founder and CEO at Datafarm. “This latest version of S-Cubed Templates is the most comprehensive, yet simple solution for regulatory submission document authoring on the market. It is a significant step in our mission to simplify the entire electronic submissions process.”

S-Cubed Templates is an essential utility for the Regulatory and Clinical authoring community that offers ICH and regional agency compliant document templates. The package includes over 250 comprehensive shell documents to create CTD, eCTD and NeES documents as well as CTA documents required for EU regulatory submissions. S-Cubed Templates are pre-formatted with instructions and references to various guidelines, available in Arial and Times New Roman fonts, and US Letter and A4 sizes. S-Cubed Templates also supports both Microsoft Word 2003 and 2007 and offers some of the complex word functionalities with a single button click.

For more information about S-Cubed Templates version 2.0 please visit
<http://www.datafarminc.com/Products-Templates.aspx>.

About Datafarm Inc.

Established in 1997, Datafarm is a world leader in high performance electronic regulatory submission solutions for the life sciences industry. Datafarm's open, modular technologies and professional services experts enable life sciences companies to meet the strict standards of regulatory authorities across the world, helping them achieve quality, accuracy, and compliance to efficiently deliver regulatory reports and submissions. Headquartered in Marlborough, MA, Datafarm has offices in the US, UK, France, and India. For more information, visit www.datafarminc.com.

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