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For immediate Release

Datafarm™ announces HollisterStier Contract Manufacturing Regulatory Affairs Deploys S-Cubed® eCTD

-HollisterStier expands electronic regulatory submission capabilities with Datafarm's S-Cubed eCTD-

Marlborough, MA, USA September 30, 2009 — Datafarm™, a world leading provider of high performance electronic regulatory submission solutions for the life sciences industry, today announced that HollisterStier Contract Manufacturing installed and validated S-Cubed® eCTD to manage and compile electronic regulatory submissions for the FDA. This would include drug product information supplied by HollisterStier, along with the clients' submission ready drug substance, clinical and non-clinical data.

HollisterStier's Regulatory Affairs department adds this software to their existing electronic capability which includes assisting clients in the creation of an FDA Electronic Submission Gateway (ESG) account at client sites for direct submission of electronic information to the FDA. Additionally, HollisterStier can perform electronic establishment registration and electronic drug listing using Structured Product Labeling (SPL) eXtensible Markup Language (XML) functions for clients or train clients' staff to perform these functions. HollisterStier's electronic submission offerings provide a comprehensive set of regulatory support options under the same roof as fill/finish operations.

About S-Cubed eCTD

Datafarm's S-Cubed® eCTD includes all of the latest global regulatory agency requirements so that users can quickly create consistent and compliant eCTD submissions. S-Cubed eCTD enables complete lifecycle management of the eCTD allowing users to publish, compile, review, validate and track all content throughout the entire drug development process. S-Cubed® eCTD conforms to the current ICH and Regional requirements for delivering eCTD submissions to the US, EU, Canada, Japan, and Taiwan regulatory agencies. For more information about S-Cubed eCTD visit

<http://www.datafarminc.com/Products-S-Cubed.aspx#eCTD>.

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

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world, helping them achieve quality, accuracy, and compliance to efficiently deliver regulatory reports and submissions. Headquartered in Marlborough, Massachusetts, US, Datafarm has regional offices in California in the US, UK, France, and India. Information about Datafarm's products and services can be found at <http://www.datafarminc.com>.

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