



FOR IMMEDIATE RELEASE

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SAGE SUBMISSIONS ANNOUNCES RELEASE OF SAGE TEMPLATES VERSION 1.1.3

New eCTD Templates support EMEA Module 1, multiple languages and options for additional user flexibility

San Diego, CA – October 6, 2009 – Sage Submissions ([Sage](#)), a leading supplier of templates for eCTD submissions, announces the release of Sage Templates Version 1.1.3. Sage Templates are immediately available for purchase on the [Sage](#) web site and through Sage Submissions reseller partners.

Sage Templates were first launched in 2007 and delivered to customers in North America, Western Europe and Asia. Sage Templates operate within Microsoft Word versions 2000, 2003, and 2007. Extensive customer input was gathered to enhance the design of Sage Templates and extend the number of templates available with Version 1.1.3. Version 1.1.3 has been enhanced in many areas including:

- Redesign of header and footer content to take into account regulatory authority feedback about ease-of-review features desired by submission reviewers
- Redesign and extension of the Styles, Tables and Symbols toolbars
- Addition of the Pharmacokinetics toolbars
- New content templates for:
 - European Medicines Agency (EMA) Centralised Procedure Module 1 for eCTD

- EMEA Clinical Trial Authorisation-Investigational Medicinal Product (CTA-IMP)
- US FDA Office of Generic Drugs Question based Review (QbR) for Module 2 Quality Overall Summary
- Enhancements to User Documentation and Online Training.
- Acrobat Job Options in US Letter and A4 page size to assure conversion of Sage Templates to comply with ICH, EMEA, Health Canada, and FDA guidance for PDF files for regulatory review.
- Full set of templates are available with Toolbars and Styles in Spanish, German and Polish for users operating in multilingual versions of Microsoft Word. Other Western European languages are available upon request
- Subsets of the Sage Templates and Training are now available. This provides a low cost acquisition point for companies that prepare Drug Master Files, CROs that conduct nonclinical or clinical studies, or companies that submit only IND or CTA-IMP submissions.

Information about this and other News and Events may be found at http://www.sagesubmissions.com/content_main/news_and_events.php.

Representatives will also be available at the upcoming DIA Electronic Submissions conference in San Diego, CA, November 17-19, 2009.

ABOUT SAGE SUBMISSIONS

Founded in 2007, Sage Submissions is a proven market leader in providing regulatory submission templates, training and consulting to life science companies. Sage Submissions serves life science companies of all stages of development and business models in North American, Western Europe, Asia and all regions that produce drug or biologic submissions in the electronic Common Technical Document (eCTD) format.

Sage Submissions is a privately held company with headquarters in Colorado and business and product development operations in California. For more information visit <http://www.SageSubmissions.com>.

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