

OMNICIA ENRICHES ITS OMNIEFILE™ PRODUCT FAMILY BY ADDING NEW PRODUCTS AND FEATURES

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At the RAPS 2004 Annual Conference and Exhibition in Washington DC, Omnicia showcased the pre-launch of version 2 of the OmnieFile™ Suite of authoring and publishing products for the Life Sciences Industry. The OmnieFile Suite includes the OmnieTemplate Authoring Package, the OmnieFile Toolkit Productivity Tools, the OmnieFile Publishing System Software, and the complete bundled package – OmnieFile Suite.

“We are very proud of what we’ve accomplished for less than a year in product development and feature-enhancements. Feedback from our clients played a major role.” said Ms. Peachy Dimanlig, Corporate Operating Officer.

By implementing new features and enhancing the existing ones, Omnicia again offers a fully integrated, feature rich, and easy-to-use solution. Customers now have access to a wide gamut of integrated tools for compiling electronic, paper or hybrid submissions in a timely manner and at a down to earth cost.

“The Life Sciences Industry finally has a robust, fully integrated, intuitive, easy-to-use publishing alternative to compile submissions. Our goal was to produce a product line that would work in concert with the drug development process and reduce FTE costs associated with submission activities. We have successfully achieved our goal.” said Mr. Reuben Jenkins, President and Chief Executive Officer.

“At Omnicia our clients are the highest priority and their needs drive our product development process. Industry regulations and technology are constantly changing, so we continue to enhance our products to be fully compliant with the FDA and ICH guidance documents. We always strive for excellence.” proudly added Ms. Dimanlig.

Dedication, diligence and client interaction has yielded a feature rich product family to compile regulatory submissions to the worldwide agencies. Customers can begin submission activities at any and every stage of the drug development process, no longer being forced to wait until documents are final. Omnicia’s technology can accommodate all the aspects of the electronic, hybrid, or paper submission’s life cycle, and are designed by industry practitioners to automate and streamline the author’s and publisher’s work. The results are increased productivity, reduced FTE costs and faster regulatory approvals.

Pharmaceutical, Biotechnology, and Medical Device companies can now leverage any or all of Omnicia's technology solutions including:

- **OmnieTemplates** - a set of powerful templates specifically designed to ease the generation of submission-ready documents: eCTD, eNDA, eBLA, IND, 510K, PMA, Amendments, Supplements, Reports, Protocols, SOPs.
- **OmnieFile Toolkit** - integrates feature rich tools to automatically create and manage bookmarks and hyperlinks, headers and footers, and Microsoft Word and Adobe PDF tables of contents.
- **OmnieFile Publishing System** - an intuitive solution designed to publish compliant electronic drug development reports, marketing applications and view eCTD, M1 and Study Tagging File indexes.
- **OmnieFile Suite** - a robust and complete solution for authoring and publishing compliant electronic submissions that comes with powerful submission templates; advanced bookmark, cross-reference, and cross-document TOC generation tools; an optimized Adobe PDF rendering engine; and a fully integrated eCTD builder and viewer.

About Omnicia, Inc.

Omnicia, a privately held corporation founded in 2001, provides software solutions and professional services to compile electronic submissions for the Life Sciences Industry. Omnicia's worldwide headquarters are located in the heart of Biotech Bay, South San Francisco. For more information, call 650-588-2188, email info@omniciainc.com, or visit www.omniciainc.com.