



**Press release – 20th September 2006**

**XClinical presents and exhibits at the CDISC Interchange Conference in Bethesda, MD**

At the upcoming CDISC Interchange Conference in Bethesda, MD, September 25 through 29, XClinical will present a first-in-class real-life implementation of a full end-to-end CDISC-based clinical trial data flow. The process includes automated CRF design and e-CRF set-up, online data capture and cleaning, and CDISC ODM export all the way down to the automatic generation of CDISC SDTM datasets for submission to the FDA and storage in a clinical data warehouse.

Our team would like to invite you to attend our plenary presentation and to visit us at our exhibition booth. MARVIN, our cost-effective online EDC-CDM system stands for very fast e-CRF set-up and is fully compliant with CDISC data standards as well as all applicable legal requirements (21 CFR 11, GCP system validation, data privacy protection, etc.). If you would like to make an appointment for a personal discussion and an in-depth demonstration of MARVIN, please send an email to [info@xclinical.com](mailto:info@xclinical.com) or call us at +1 / 301 841-7784 or toll free at +1 / 866 871-5889.

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XClinical GmbH ([www.xclinical.com](http://www.xclinical.com)) is an EDC-CDM system vendor based in Munich, Paris and Bethesda, USA. XClinical was founded in 2002 by a team of highly experienced software developers, clinicians and managers. Since 4 years XClinical has sustained profitable growth without external investors. More and more biopharmaceutical corporations and medical devices manufacturers prefer MARVIN over competitive offerings, since MARVIN is designed to be an attractive working tool especially for investigators, the key people that drive the success of a clinical trial; moreover, MARVIN is fully compliant with the CDISC data standards and offers fast and flexible e-CRF set-up. Till today, MARVIN was used to collect and process data of more than 43,000 patients in 26 studies.

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