

NEWS RELEASE

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DataCeutics Chosen for Phase I of Multi-Phase, International Report Standardization Project.

Philadelphia, Pa - DataCeutics, Inc.

DataCeutics announced today that it has been chosen to lead the development and implementation of the first phase of a multi-phase, international clinical reporting and standards project. This project will include the sponsor's Japanese, US and European offices but will be managed from its offices in Chicago. The first phase of the effort will require development of requirements and design specifications. SAS has been chosen as the software tool for new macros and report standards that will be adopted across all sponsor sites and clinical reporting departments. CDISC Adam was identified as a possible standard that the company will embrace.

Paul Gilbert, President and System Architect for DataCeutics said, "DataCeutics high quality standards, solid software development life cycle, and excellent documentation practices will guarantee that the system developed for our sponsor will meet all of their requirements and FDA regulations." He continued, "We have over 100 cumulative years' experience developing and delivering software solutions for our customers in this market and we will meet their timelines and budget."

This project represents just the first phase of a much larger effort that will require overhauling all of the company's reports, their reporting environment and code management. Phase I is expected to run until mid-May with the follow-on work to commence soon after and lasting until 2009.

DataCeutics, Inc.

The Leader in Human Research Information Technology

DataCeutics, Inc. is the leader in human research IT and compliance services. Our solutions include both software products and services. We were founded in 1993 and have over 50 clients in the pharmaceutical, biotechnology, medical device and CRO industries. Primary services include Human Research Business Process Consulting, Project Management, EDC, CDM, Safety and Clinical Reporting Systems Selection, Development, Implementation & Support, Compliance Assessments and Remediation, Systems Validation, Auditing, System and Data Migration, CDM and SAS Programming, Software Verification, Training, Regulations (GxPs, 21 CFR Part 11), Standards (IEEE, CDISC, etc.), e-Submissions.

DataCeutics' leading software products are the Clinical Reporting Solution™, CR Toolkit™ and DataCeutics Report Portal™.

To obtain more information about DataCeutics, our products and services, please contact us.

