

NEWS RELEASE

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The CR Toolkit and SAS Drug Development are an excellent match
Philadelphia, Pa - DataCeutics, Inc.

DataCeutics is set to meet a major milestone in an ongoing clinical reporting development project fusing SAS® Drug Development with its own CR Toolkit™. By utilizing the CR Toolkit as the underlying technology for developing a library of reusable SAS macros, and SAS Drug Development for an enterprise-wide collaboration tool, DataCeutics is positioned to deliver a fully configurable solution to a global user base spanning Japan, Europe and the United States.

The reusable macro development project entails the implementation of global standards in statistical computing, including the underlying data and output standards. With client-supplied CDISC safety data domains, DataCeutics is providing the expertise for development of standard, reusable macros for generating tables listing and figures; and for performing the data transformations from SDTM to ADaM for regulatory submissions.

DataCeutics' CR Toolkit has been in use by the client for over 10 years to facilitate in generating clinical reports, tables, listings and figures. Coupling the CR Toolkit with the industry-leading data management and analytic capabilities of SAS Drug Development, makes the data compliant, as well as more accessible to users across the enterprise. Partnering with an existing client to incorporate innovative repository technologies, and to harmonize data standards enables DataCeutics to focus on its core competencies, and qualifies the CR Toolkit as the tool of choice for automating and standardizing clinical report generation.

DataCeutics, Inc.

The Leading Functional Service Provider for the Life Sciences Industry

DataCeutics, Inc. is the leader in reporting, data management and compliance for life science companies. Our solutions include better processes, services and software products. 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 services, please contact us.

