SAS® PROGRAMMING

clinovo
Clinical Data Solutions

LEVERAGE SAS TECHNOLOGY TO ENHANCE YOUR EFFICIENCY

SAS® is the indisputable leading programming platform in the clinical trial industry. We leverage SAS® technologies to monitor key study metrics and prepare your statistical submission to the FDA. We also use SAS® to create user-friendly, safe and compliant web applications to streamline data management and biostatistics processes.

PROGRAM

CLINOVO SAS® EXPERTS

- Our local team members have **15 years of experience** in the clinical trial industry.
  - ✓ Pharmaceutical and Medical Device
  - ✓ 100% staff SAS® certified
  - ✓ Experts in complex programming
  - ✓ Speakers at all major SAS® events
- To supplement our local SAS® experts, we have an offshore team of SAS® programmers to meet your highest expectations on time and on budget.

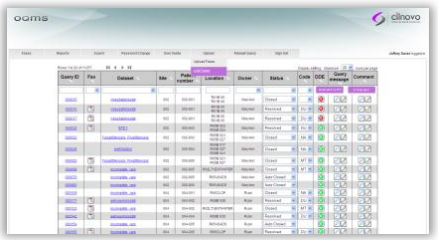
SAS®-BASED SYSTEMS

CDISC® Express: Open source CDISC conversion tool

NEAT: Faster data cleaning and query resolution

OQMS: Query Management System

Patient Profile: Comprehensive, on demand and easy-to-read patient data



CLINOVO SAS® PROGRAMMING SERVICES

We offer a comprehensive range of SAS® Programming Services tailored to your needs.

Clinical Programming

- ✓ Documentation (Data Transfer Plan)
- ✓ Ad hoc clinical data reports
- ✓ Edit check programming
- ✓ Dashboard reports
- ✓ Report automation
- ✓ Data transfers

Statistical Programming

- ✓ Analysis dataset
- ✓ Tables, listings and figures
- ✓ Patient profiles
- ✓ CDISC SDTM and ADaM mapping
- ✓ System validation

SAS GLOBAL FORUM 2011

SAS GLOBAL FORUM BEST PAPER

FDA

2 FDA PANEL SUPPORT

sas

SAS BUSINESS PARTNER

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