TrialStat Portal

A New Paradigm in Data Management Technology

TrialStat Portal is a paradigm shift in how you manage your studies. Providing insight across all of your studies, coupled with integration into 3rd party systems, Portal is your ultimate study management platform. Capabilities such as real-time visualization of your study data allows you to make informed decisions quickly and easily and implement those new best practices in future studies.

Portal is seamlessly integrated with all TrialStat products providing advanced user dashboards to retrieve and drill down into the data. Portal enables users to make critical decisions quickly and efficiently allowing you to continuously maintain control of the study while staying on time and on budget

TrialStat Portal – Real-Time Data & Decisions

- ✓ Real-Time Data Visualization
- ✓ Cross Study Reporting
- ✓ 40+ Standard Reports
- ✓ Unlimited Custom Reports & Dashboards
- ✓ Dynamically Drill Down Into Live Data
- ✓ Make Critical Decisions Sooner
- ✓ Browser Based & Mobile Responsive
- ✓ Integrate With 3rd Party Systems

Empower Your Sites

TrialStat Portal is your central location for managing the performance and conduct or your sites, including all site documentation and communication. Keep sites engaged and informed with timely study updates, site-specific documents, information and real-time comparison metrics with other sites . Use real-time site comparison data to monitor empower troubled sites and encourage top performers to effectively manage enrollment and compliance.

Empowering Sponsors

Sponsors can finally make study decisions in real-time, no longer having to wait for reporting on key study metrics. TrialStat's configurable dashboards with real-time data and instant notifications ensure that Sponsors have the data they need when it's needed most! TrialStat's real time dashboards, remove the unnecessary limitations of manual reports with stale data ensuring trials progress efficiently.

Features CROs Have Always Wanted

CRO's finally have a Data Management platform that meets all their specific study requirements. Completely configurable eCRFs, User Roles, Workflows and multi-lingual capabilities, to support for all study phases including pilot / proof of concept studies, Phase I, II, III and IV studies and the unique requirements of Medical Device studies and the necessary pricing models to support this diverse set of study requirements. Coupled with the extensibility of our API to connect with 3rd party systems such as EDC, and Safety Systems and our ability to create custom validated solutions, CROs no longer have to compromise on requirements or use inferior technology.

Jubilant Biosys

FDA 21 CFR Part 11 Compliant

HIPAA Compliance Solutions

FISMA Compliance Solutions

rackspace

Data Without Limitations

Book Your Demo Today

1-888-488-0312

Corporate Overview

- ✓ Delivering Successful Studies since 2003
- ✓ Global studies in more than 50 countries & 3000 users
- ✓ Subsidiary of Jubilant Life Sciences
- ✓ Over 500 completed studies

Professional Services

- ✓ Study Development - Case Report Form Design
- ✓ Project Management
- ✓ Complete eCRF Validation
- ✓ Custom Programming & Systems Integration

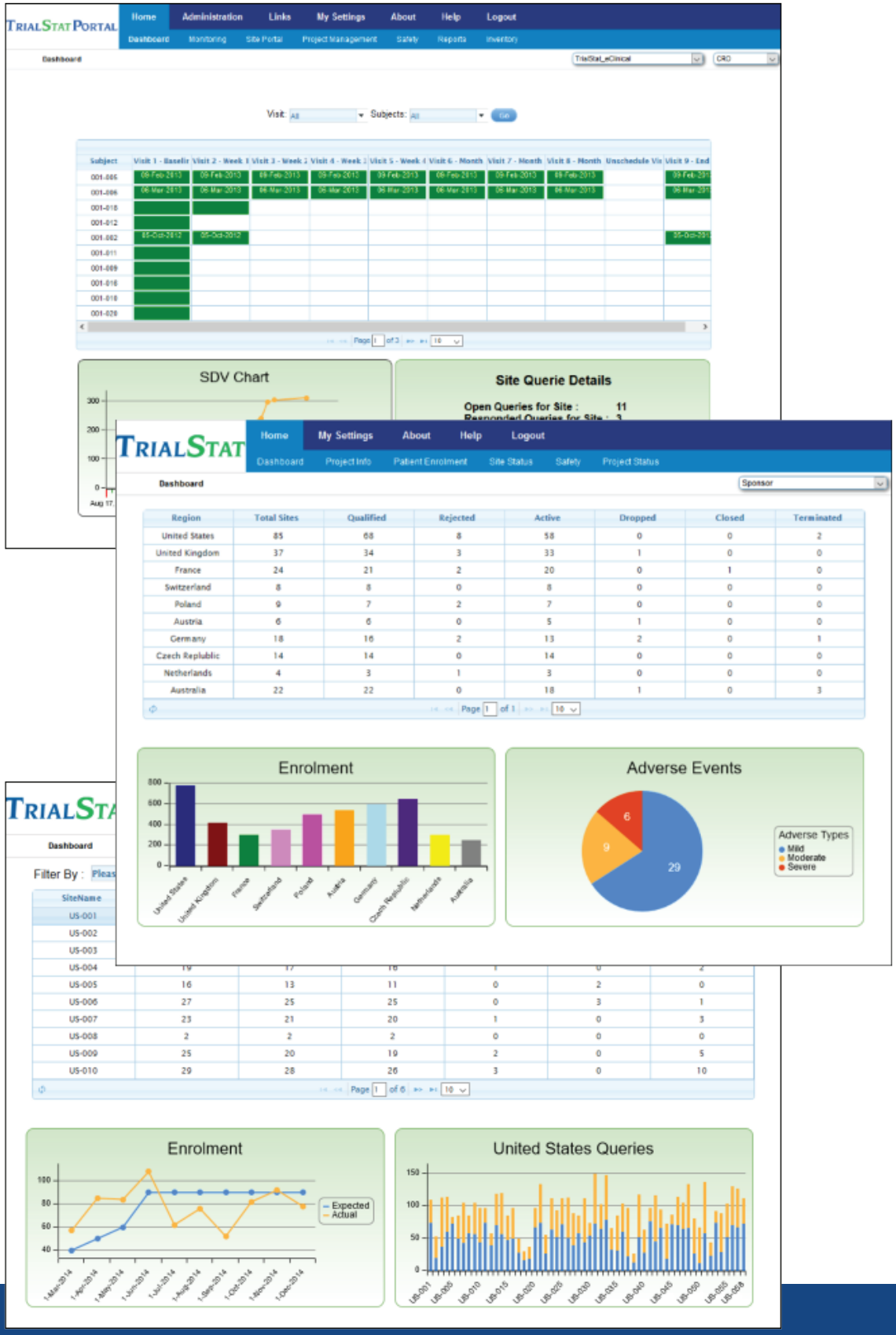
Regulatory Compliance

- ✓ 21 CFR Part 11 Compliant
- ✓ HIPAA Compliant
- ✓ SSAE 16 SOC 1 data centers
- ✓ SAS 70 Type II

TrialStat

TrialStat Solutions Inc.
www.trialstat.com | 888-488-0312

A Jubilant Life Sciences Company



TRIALSTAT

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