Clinical Trials Center

Clinical Data Management

USZ Universitäts Spital Zürich

Services
for Clinical Research

Head

Dirk Smolinski

Consulting

- Requirements for Clinical Data Management regarding applicable Swiss rules
- Electronic Data Capture of study data
- Import of external data in study data base
- Simplify analysis of study data by using medical dictionaries
- Conduct of randomized and blinded trials
- Direct data capture by patient
- Secure archiving of electronic study data
- Process descriptions for quality management documents

Clinical Trials Software

- Professional software fulfilling the strict demands for clinical trials
- GCP-conform features
- Centralized data base on dedicated servers run by the University Hospital Zurich
- Web based data capture using standard browsers
- Secure pseudonymization of study data
- Comprehensive data validation tools
- Personalized access, role based user management
- Multi center trials in an international environment by multilingual user interface

Study software provided:
- secuTrial (ias GmbH, Berlin)
- REDCap (Vanderbilt University)

User friendly development tool - most of our customers build their eCRF by themselves. Development of the eCRF according to your draft by the CTC DM.
Please contact us for an offer to implement the software to your study

Training / Further Education

CAS Clinical Data Management (12 ECTS):
Theory and use in practice
Workshops and colloquia
CAS CDM can be completed within 1 year part-time

How to find us

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