Clinical Trials Center

ISO 9001 CERTIFIED 2015

Monitoring

USZ Universitäts Spital Zürich

Services for Clinical Research

Head	**Karin Huber, MSc**
Consulting	Monitoring by the Clinical Trials Center staff Process optimization Standard Operating Procedures and Working Instructions
Support	Compilation of Notification Files Setup and maintenance of Trial Master Files Support with the organization of Investigator Meetings
Site Qualification Visits	Facilities, equipment, staff Responsibilities of Investigator/Sponsor and Investigator Discussion of Study Protocol, recruitment, study conduct
Site Initiation Visits	Discussion of study conduct and documentation Safety Reporting, use of case report forms, cooperation with laboratories
Routine Monitoring Visits	Verification of source data and study documentation, e.g. Trial Master File, Safety Reporting
Close-out Visits	Debriefing regarding responsibilities, reporting and archiving Close-out verification of study documentation and disposal of study medications
Tariff	Please ask for a quotation

How to find us

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