# Clinical Trials Center

ISO 9001 CERTIFIED 2015

## Regulatory Affairs

USZ Universitäts Spital Zürich

# Services
# for Clinical Research

**Head**

**Annette Widmann, MD**

**Consulting**

Personal consulting on
- study planning, study preparation, study conduct
- project-specific study files
- submitting to IRB (Cantonal Ethics Committee) and authorities (Swissmedic and others)
- insurance certificates for clinical trials
- general regulatory questions

**Document Review**

Review of study files (study protocols, etc.)
Support with creation and revision of study documents

**Document Templates**

Document templates are made available for
- study protocol
- amendment
- investigator‘s brochure

etc.

**Tariff**

For projects of the UniversityHospital Zurich, the University of Zurich and associated hospitals services are free of charge. For external clients rates are according to contract

**How to find us**


**Office** Moussonstr. 2, via basement/terrace, 8044 Zurich

**Post** Clinical Trials Center
University Hospital Zurich
Raemistrasse 100 / MOU2
8091 Zurich

**Phone** +41 43 254 0186
**Mail** CTC-RA@usz.ch
**Web** www.ctc.usz.ch