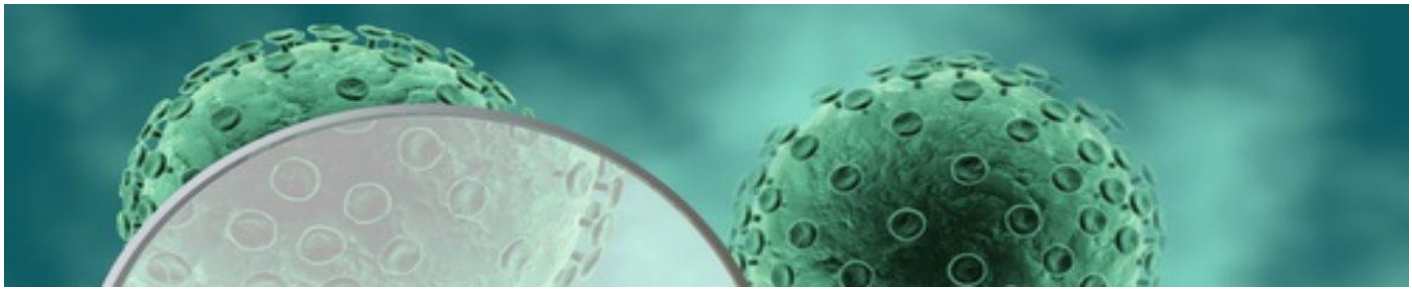
CAS in Clinical Research



Writing a study protocol

Description

Researchers are required to write various types of protocols. Many researchers consider this a difficult, or even the most difficult part of the research process. Understanding the differences between various types of protocols is important. This course concentrates on the processes of writing different protocols (not on the study design), using researchers' actual studies.

Objectives

The goal of this course is to enable researchers to know the different types of protocols (study protocol, grant proposal, submission to ethics committees) and to get them written and sent.

Dates

August Mon 17- Wed 19, 2020

Equipment

Please bring along your laptop

Course Structure

Participants each bring an idea for, or draft of a study protocol, a grant proposal or a submission to an ethics committee. We will help you to develop this. By the end of the course, participants will have a clear idea of how to complete their protocol – and have a timetable for doing so.

Day 1	Day 2	Day 3
Introduction: Differences between the various types and purposes of protocols/ pitfalls/ target audience/ expectations from reviewers **Exercises on structure and writing:** -Setting the brief -Organizing the material -Planning the protocol -Writing the protocol	Participants work on their own protocols, the teachers are available for personal support	**Morning:** Discussion and revision of drafts/ From a reviewer's perspective **Afternoon:** An afternoon with funders from EU/ SNF/ / Ethics Committee
Nicola Low/ Sven Trelle/ Dianne Egli	Nicola Low/ Sven Trelle	Sven Trelle/ Sasha Hugentobler/ Claudia Kühni/ Matthias Briel/ Peter Kleist

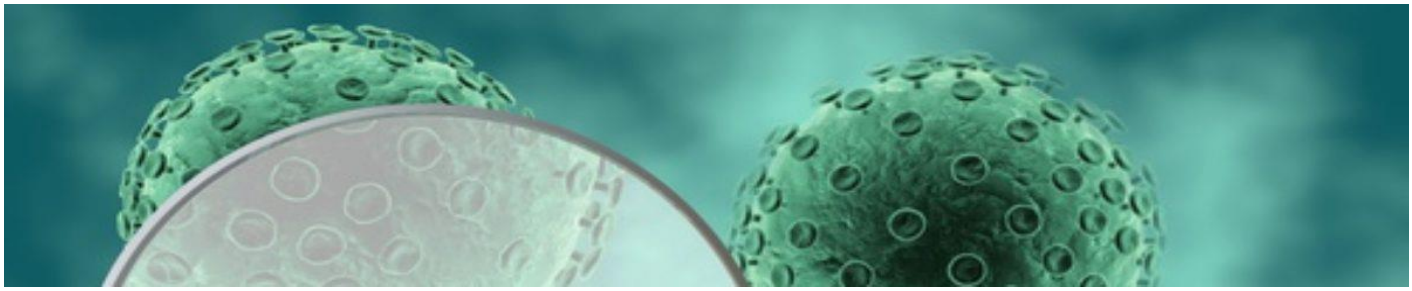
Contact:

Course coordination: Ann Walser

Address: ISPM Bern
Mittelstrasse 43, CH-3012 Bern

Telephone: +41 31 631 58 94
Email: casclinresearch@ispm.unibe.ch
Website: www.cas-clinicalresearch.ch

CAS in Clinical Research



Assessment

Attainment of learning objectives measured by assessing progress of participants' own proposed protocol

Credits

2 ECTS

Preliminary Work: 20+ h, Contact: 12 h, In-course work: 4-6 h, Wrap-up Work: 15-20 h

(1 ECTS corresponds to appr. 25/30 hours' work)

Facilitators

Prof. Nicola Low, ISPM, University Bern

PD Dr. Sven Trelle, CTU, University Bern

Dianne Egli, MSc, ISPM, University Bern

Prof. Claudia Kühni, ISPM, University Bern, SNF

PD Dr. Matthias Briel, Department of Clinical Research, University of Basel

Sasha Hugentobler, PhD, Euresearch

Peter Kleist, Cantonal Ethics Committee, Zurich

Location

University of Bern, Room: tba

Contact:

Course coordination: Ann Walser

Address: ISPM Bern
Mittelstrasse 43, CH-3012 Bern

Telephone: +41 31 631 58 94
Email: casclinresearch@ispm.unibe.ch
Website: www.cas-clinicalresearch.ch