

How2GCP e-Learning Tool – User evaluation sheet

On behalf of the how2GCP.ch team, we would like to ask you to take a few minutes of your time in order to give us a feedback on your experience with the e-Learning Tool. With your help we may implement subsequent improvements of this tool.

Please check only one answer per question.

1. What type of organization do you belong to?

- Ethics Committee/Regulatory Authority..... 0
- Investigator/Study Coordinator/Study nurse..... 0
- Contract Research Organization (CRO)..... 0
- Industry- Headquarters/Biotech (headcount headquarter < 500)..... 0
- Industry- Headquarters (headcount headquarter > 500) 0
- Industry- Local Affiliates..... 0
- Other 0

2. How did you hear about the how2GCP.ch e-Learning Tool?

- My company recommended the participation..... 0
- From a colleague..... 0
- From someone who has worked with the tool 0
- I received a flyer 0
- From the SwAPP annual meeting..... 0
- From the SwAPP homepage 0
- The Sponsor of a clinical trial offered it 0
- Other: _____ 0

3. How many years of GCP experience do you have? _____ years.

4. Would you consider using how2GCP.ch as a reference tool? Yes 0 / No 0

5. Did you encounter any technical problems using the e-Learning GCP tool how2GCP.ch?

6. In general, how would you rate the following items?

(circle one per line; 4 very good; 3 good; 2: acceptable; 1 unacceptable)

- Overall quality of the e-Learning Tool	4	3	2	1
- Usefulness of the topics to your everyday work	4	3	2	1
- Level of teaching	4	3	2	1
- Amount learned	4	3	2	1
- User-friendly set-up of the tool	4	3	2	1
- Loading time in your browser	4	3	2	1
- User support	4	3	2	1

7. In general, how would you rate the quality of the content in the different modules?
(circle one per line; 4 very good; 3 good; 2: acceptable; 1 unacceptable)

- What is GCP?	4	3	2	1
- Roles and responsibilities	4	3	2	1
- Clinical trial processes for Investigators	4	3	2	1
- Clinical trial processes for Sponsors	4	3	2	1
- Clinical trial processes for Monitors	4	3	2	1
- Investigator selection and agreement	4	3	2	1
- Pre-trial processes	4	3	2	1
- During the trial processes	4	3	2	1
- Post-trial processes	4	3	2	1

8. Did you learn anything new? Yes / No

9. If SwAPP were to extend this course at a later point of time, what would your preferences for additional content be (e.g. additional languages, EU Directive, additional content for Regulatory, other topics)?

10. Would you recommend how2GCP.ch to a colleague? Yes / No

11. Any additional comments?

Thank you very much for taking the time to complete this form.

Please return:

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