

# **Pre-exam answers**

## **Research ethics in empirical studies**

**Lecture 6**

**Usability evaluation CS-E5210**

**Antti Salovaara**

# Guest lecture advertisement

# Next Tuesday: Esko Kurvinen



Service design lead at Elisa  
Doctor of Arts (industrial design)

Experimentation, A/B testing,  
theory, practices, examples...

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# Pre-exam answers

# Questions

1. Scenarios are important in both walkthroughs and empirical usability evaluations. Why? (10 p)
2. Using the concepts of reliability and/or validity, explain why it is a good idea that a heuristic evaluation is carried out by 3–5 evaluators. (10 p)
3. Consider a heart rate monitoring machine installed in a hospital. Briefly describe three critical aspects of usability from the perspective of nursing staff and patients. Explain why you identified these aspects in particular. (10 p)

# Why did we have an exam?

## Improved group communication:

Same knowledge basis helps you understand each other

Faster and more precise communication

## Quality control:

Students' different starting levels get closer to each other

## Professionalism:

You can now meet the customer companies with good confidence

## Rewarding you for personal effort:

You can affect your grade and prove your skills to us in more ways than just participating in group work

# Question 1

*Scenarios are important in both walkthroughs and empirical usability evaluations. Why? (10 p)*

- Scenario is a fictitious story and context that depicts a typical user persona, her motivations, needs, and activities
- How the user can accomplish an action or goal?
- Also exposes associated problems and limitations
- A typical scenario documents the process via which a user might use the design
- User scenario are often used in various stages of product design particularly ideation and usability evaluation

# Significance of scenarios

- Inexpensive, quick, and easy to create
- Reflect multiple tasks, use context, and user characteristics
- Can be tailored to test new features
- Evaluations become more realistic and natural
- Actual needs, motivations, and problems can be better comprehended



## Question 2...

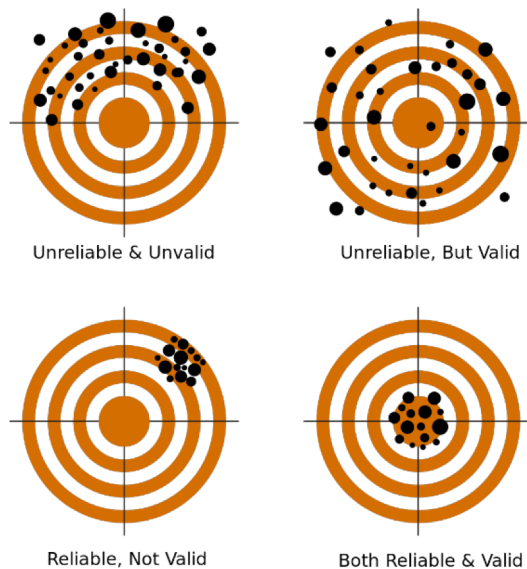
*Using the concepts of reliability and/or validity, explain why it is a good idea that a heuristic evaluation is carried out by 3–5 evaluators. (10 p)*

“How many evaluators are needed to make the analysis work? That depends on how knowledgeable the evaluators are. If the evaluators are experienced interface experts, then 3 to 5 evaluators can catch all of the "heuristically identifiable" major problems [...].”

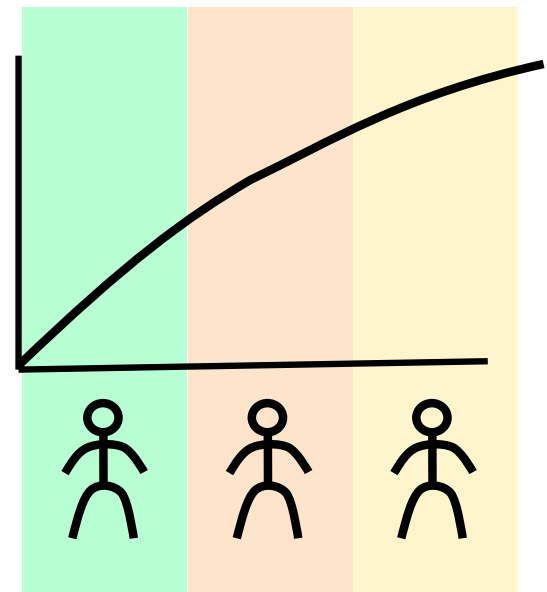
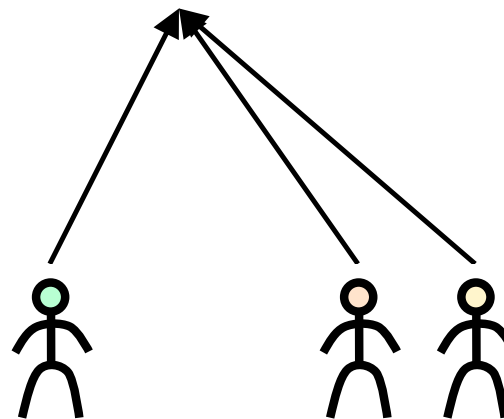
Lewis, C. & Rieman, J. (1994) Task-Centered User Interface Design, Heuristic Analysis.

## Question 2

*Using the concepts of reliability and/or validity, explain why it is a good idea that a heuristic evaluation is carried out by 3–5 evaluators. (10 p)*



### Usability



## Question 3

*Consider a heart rate monitoring machine installed in a hospital. Briefly describe **three critical aspects** of usability from the perspective of nursing staff and patients. Explain **why** you identified these aspects in particular. (10 p)*

This question did not have a simple correct answer

The purpose was to elicit analyses and prioritizations of different aspects

A possible answer structure:

General analysis of the context of use

Staff viewpoint: 3 x ( aspect + explanation )

Patient viewpoint: 3 x ( aspect + explanation )

# Research ethics in empirical studies

How human participants need to be treated  
Ownership and access to research data

# Research ethics

Informed consent

Ethics guidelines

Anonymity

Access to raw data vs only findings

Reporting the results to companies

Intellectual property rights



ing area of social science research that needs to be approached with sensitivity and with vigilance regarding personal privacy issues.

Questions have been raised about the principles of informed consent and opportunity to opt out in connection with the research in this paper. The authors noted in their paper, “[The work] was consistent with Facebook’s Data Use Policy, to which all users agree prior to creating an account on Facebook, constituting informed consent for this research.” When the authors prepared their paper for publication in PNAS, they stated that: “Because this experiment was conducted by Facebook, Inc. for internal purposes, the Cornell University IRB [Institutional Review Board] determined that the project did not fall under Cornell’s Human Research Protection Program.” This statement has since been [confirmed by Cornell University](#).

Obtaining informed consent and allowing participants to opt out are best practices in most instances under the US Department of Health and Human Services Policy for the Protection of Human Research Subjects (the “[Common Rule](#)”). Adherence to the Common Rule is [PNAS policy](#), but as a private company Facebook was under no obligation to conform to the provisions of the Common Rule when it collected the data used by the authors, and the

# Exercise

1. Make a list of issues that your group work will have to address (silent work 2 min)
2. When you listen to the following slides, make additions
3. Discuss these issues in your group mentor meeting

**Informed consent**

**Access to raw data vs only findings**

**Ethics guidelines**

**Reporting the results to companies**

**Anonymity**

**Intellectual property rights**



# Aspects of participant-facing ethics

Old University of Helsinki psychologists' guidelines:

Respect for the participant

Anonymity

Informed consent + voluntariness of participation

Possibility to opt out

Debrief after the study

Priority in participant's well-being

(+ and many other aspects)

Also other dimensions of ethics exist (e.g., lawfulness, fairness in academic publishing, independence and integrity, etc.)



# When you are in the planning stage

Your study needs to be approved by a Research Ethics Committee if it includes at least one of the following:

1. Intervention in the **physical integrity** of subject
2. The study deviates from the principle of **informed consent** (excluding archival data)
3. The subjects are **children** under the age of 15
4. **Exceptionally strong stimuli** whose harmfulness needs to be evaluated by an expert
5. Possible long-term **mental harm** (trauma, depression, sleeplessness)
6. Possible **security risk** to subjects

(see <http://www.tenk.fi/en/ethical-review-human-sciences/ethical-review>)

@ Aalto: see more information at:

<https://inside.aalto.fi/display/aallosta/Tutkimuseettinen+toimikunta>

<https://inside.aalto.fi/display/AboutAalto/Research+Ethics+Committee>

# Before starting an experiment with participant

Researcher describes the study, its requirements, participant's rights, researchers' responsibilities, etc:

- Who are the members of the research team that organize this study

- That the purpose is not to evaluate the participant, but to investigate a research question

- That the participant may opt out any time during the study

- That the **relevant** material created by participants may be used in reports and publications

  - material not relevant to the RQ will not be reported

- Confidentiality of the data: who will see it and in what form (see the next slide)

Get the participant sign an informed consent

# Levels of anonymity + access to data

(with some suggestions)

## 1. All (“raw”) data

Including the participant’s name

→ Access to research team only

## 2. Aggregated data

= Anonymized and not traceable to individuals in other ways either

→ Access to research team only or also to companies

## 3. Reportable data

= Data that addresses the research questions or is evidence of your emergent findings

→ Can be published and delivered to e.g., involved companies, preferably not as datasets but as reports

# If you have third parties in the project

## Possible third parties:

“**The client**”: one that is interested in the results

“**The case company**”: the context where the study is conducted

“**Instrument provider**”: a company that provides an instrument (e.g., collaboration platform) that is used in the research

Different access rights to data may be needed for each party

Never promise access to raw data!

You would have to tell this to participants and this would potentially bias your data 😞

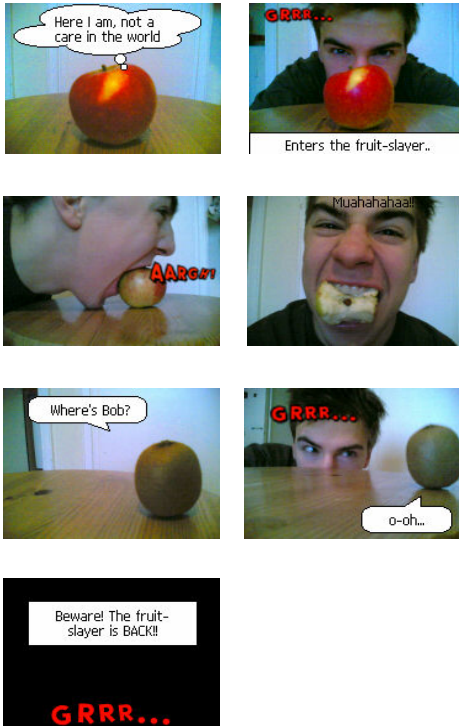
→ Provide only reports to 3<sup>rd</sup> parties

→ Report to each party only the findings that are relevant to them

→ Tell the participants which 3<sup>rd</sup> party will be reported what

# Intellectual property rights

© Unnamed participant in Salovaara (2007)



Participant-created works of art are participants' IPR

E.g., in communication studies

Publication of others' works of art requires participant's permission

Remember to mention the intention to publish participants' works already in the beginning of study

Usually participants are proud of having their works published!

If in doubt, ask your school's legal team

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