Protocol Intervention

(1) Will you be following a FERB-approved research programme of your supervisor?

○ Yes
◉ No

New data collection

(1) Will you be using:

☐ Online collection of questionnaire data
☐ Offline collection of questionnaire data
☐ Open or semi-open interview (incl. focusgroups)
☐ Regular observations
☐ Observations recorded on audio, video
☐ Naturalistic observation
☐ Participant observation (anthropology)
☑ Intervention study (not in lab)
☐ Behavioural tasks

(2) Where will the study (data collection) be conducted? If this is abroad, please note that you have to be sure of the local ethical codes of conducts and permissions

◉ the Netherlands
○ Abroad, outside the Netherlands. Please name the country

Intervention

(1) The intervention does not involve any invasive procedures

◉ Correct
○ Incorrect, please explain

(2) The intervention does not target clinical problems, sensitive issues and or content (e.g., anti-bullying, social security).

👁 Correct
O Incorrect, please explain

(3) The intervention does not preclude any group from treatment as usual

👁 Correct
O Incorrect, please explain

(4) There is no reason to expect any adverse effect, based on available literature, prior research findings, taking into account the duration and or intensity of the intervention.

👁 Correct
O Incorrect, please explain

(5) The intervention is aimed at improving the targeted processes, procedures or skills.

👁 Correct
O Incorrect, please explain

(6) I make sure that any participant that wished to stop can stop without providing a reason.

👁 Correct
O Incorrect, please explain

(7) I will do everything possible, within the limits of the prescribed procedures for data collection, to minimize the burden on participants.

👁 Correct
O Incorrect, please explain

(8) Prior to the data collection I have signed a non-disclosure agreement.

👁 Correct
O Incorrect, please explain

(9) I will strictly follow the protocol that has previously been approved by the FERB or METC

👁 Correct
O Incorrect
O Not applicable, there is no previous approval

Participants 1

(1) I will approach…

☑ Adults
☐ Children up to 12 years old
☐ Children from 13 to 15 years old
☐ Adolescents from 16 to 17 years old

(2) I will approach a vulnerable group

◉ No
○ Yes, please explain which vulnerable group you are working with and why this study cannot be carried out without this population.

(3) I am going to approach the participants using

☑ Recruitment websites, where participants register themselves
☐ Friends, familie, acquaintances
☐ Recruitment agencies that approached the participants
☐ Other, please explain

(4) I use the following resources

◉ Letter/e-mail
○ SONA systems
○ Other, please explain

(5) Participants receive PPU or financial compensation in proportion to their efforts

○ Yes, please explain
◉ No, please explain

(6) I'm going to ask consent to proceed using…

◉ A written procedure prior to data collection
○ Online, by ticking a box
○ An oral procedure prior to data collection (c.f. this description)

(7) This investigation cannot lead to coincidental findings

◉ Correct
○ Incorrect, please explain which coincidental findings may appear and how you will deal with this situation

(8) This study does not use deception

◉ Correct
○ Incorrect, it does use deception. Please explain

(9) If deception is used, I will ask permission from the participants again immediately after the data collection

O Not applicable
◉ Yes
O No, please explain

In accordance with the instruction document the information letter contains these elements:

(11) Aim of the study

◉ Yes
O No, please explain

(12) It is emphasized that it concerns student research

◉ Yes
O No, please explain

(13) Type of tasks, duration, load

◉ Yes
O No, please explain

(14) How the data are handled

◉ Yes
O No, please explain

(15) Right of removal (unless completely anonymous)

◉ Yes
O No, please explain

(16) Being able to stop voluntarily at any time without adverse consequences

◉ Yes
O No, please explain

(17) Contact person for questions

◉ Yes
O No, please explain

(18) Information letter in understandable language tailored to the target group

◉ Yes
○ No, please explain

(19) Participants will be given the opportunity to remove their personal data

◉ Yes
○ No, please explain

Informed consent

(1) For adults, I ask informed consent

◉ Actively (via 'wet' signature)
○ Online
○ Passively (explain why not actively)
○ No informed consent asked (explain)

Data management 1

(1) Data are or will be stored on faculty servers (YODA and/or FSBS research storage) in accordance with faculty protocol

◉ Yes
○ No, please explain

(2) Access to data is limited to student and supervisor(s).

◉ Yes
○ No, please explain

(3) Storage period is in accordance with faculty protocol and/or additional statutory provisions.

◉ Yes
○ No, please explain

(4) Data are not shared with external organization

◉ Correct

○ No, please explain

Attachments

(1) Questionnaire

(2) Information letter(s)

(3) Informed consent form(s)

(9) Miscellaneous documents e.g. data set description (optional)

No files have been uploaded yet

Signature

(1) Corona protocols

☑ I declare that I have read the relevant UU Corona protocols, and will follow them as I perform the study

(2) I declare that I have completed the above truthfully, my Solis ID is: