**PRIDE QUESTIONNAIRE FOR APPROVAL RESEARCH PROGRAM**

(1.1) **Title**

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(1.2) **Study type**

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| Research program |

(1.3) **Division**

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(1.4) **Start date**

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(1.5) **End date**

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# [Basic study information](#_bookmark0)

1. Name(s), position(s) and division(s) of the responsible researcher(s):

Name Position Division E-mail

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1. Name(s), position(s) and division(s) of the executive researcher(s): Name Position Division E-mail

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1. Research area/discipline:

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1. What is the study's main objective (add hypothesis if applicable)?

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1. Primary funder of the study:

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1. Does the study concern a multi-center project, e.g. in collaboration with other universities, a GGZ mental health care institution, or a university medical center?
	* Yes
* No
1. 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

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1. Has an ethical committee already conducted a review? If so, please provide a registration ID. When reviewed by another institution, attach the review under 'Miscellaneous documents' at the attachments tab
	* Yes
* No

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# [Study details (I)](#_bookmark1)

1. The research will involve only the use of anonymous survey procedures, anonymous interview procedures or the observation of anonymous behavior. There is no way to identify participants in the dataset.
	* Yes
* No
1. If the study uses pseudonimized data, do the researchers have access to the key for re-identification?
	* Yes
* No
* Not applicable
1. Are any participants confined in a correctional or detention facility?
	* Yes
* No

(4) Will the research project use online data collection methods?

* + Yes
* No
1. Does this study involve medical research or research with clear implications for medical science?
	* Yes
* No
1. Will participants be subjected to procedures or be required to follow rules of behaviour, or does the study involve psychologically invasive questionnaires, including interviews?
	* Yes
* No

# [Study details (II)](#_bookmark1)

1. Will participants that are recruited be >16 years?
	* Yes
* No
1. Will participants that are recruited be mentally competent (wilsbekwaam in Dutch)?
	* Yes
* No
1. When applicable, will a legal representative (parent or care taker) sign the consent?
* Yes
* No
* Not applicable
1. Which legal basis is used for processing personal data in this study? (see [Legal bases for processing](https://privacy-fss.sites.uu.nl/working-with-personal-data/#legal-bases-for-processing) )
	* Not applicable (data is processed anonymously)
* Public interest (increasing knowledge for society as a whole)
* Legitimate interest (increasing knowledge with regard to own (educational) processes or commissioned

research)

* Informed consent (other types of studies)
1. Will participants that are recruited provide active (either written, digitally by ticking a box online or orally and recorded) informed consent for participating? If no, please clarify.
	* Yes
* No

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1. Will the probability and magnitude of possible harm or discomfort anticipated in the research be greater than ordinarily encountered in daily life or during the performance of routine physical or psychological examinations or tests?
	* Yes
* No
1. Does the participant population contain vulnerable persons?

# Yes No

Incapacitated people ○ ○

Children ○ ○

Mentally challenged people ○ ○

People who experienced trauma ○ ○

People who have suicidal thoughts ○ ○

1. Will you process special categories of personal information? (see [Special categories of personal data](https://privacy-fss.sites.uu.nl/working-with-personal-data/#special-categories))
	* No
* Yes, and I will ask participants for consent for this processing
* Yes, but I am unable to ask participants for consent. Please elaborate

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1. Will participants be subjected to:

# Yes No

Inquiries into their sexual behavior or orientation ○ ○

Inquiries into drug use (also alcohol, smoking, soft drugs) ○ ○

Assessment of delinquency ○ ○

Inquiries relating to religious or philosophical belief ○ ○

Inquiries relating to political opinions ○ ○

Inquiries into ethnic origin ○ ○

Inquiries into trade Union membership ○ ○

Inquiries into violent experiences ○ ○

Inquiries into the physical or mental health of an individual ○ ○

Inquiries into criminal convictions and offences ○ ○

Shocking images/videos ○ ○

Deception (information letter does not state real study objective) Are you purposefully withholding information from participants before the beginning of your study?

* ○

A stimulus causing the participant pain or discomfort (electrical/ thermal shocks, noise or other forms of inducing physical discomfort)

* ○

A new technique for data collection? ○ ○

Inquiries into financial information (such as income or purchasing behavior)? ○ ○

Inquiries into study results? ○ ○

1. Will data of the following categories be processed:

# Yes No

Photo data (containing people) ○ ○

Video data (containing people) ○ ○

Genetic data (buccal, blood, hair) ○ ○

Biometric data (fingerprint, iris or retinal scan, voice recognition and face scan) ○ ○

Directly identifying data (name, address, date of birth or a combination of data that makes it possible to identify a person)

* ○

Location data (GPS, IP-address ○ ○

Electronic communication data, e.g. text messages, phone metadata ○ ○

# [Study details (III)](#_bookmark2)

1. What is the study's theoretical and practical relevance? (500 words max.):

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1. What are the central questions?

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1. What is the study's design and procedure? (500 words max.):

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1. Optional attachments:
2. Describe the data collection instruments, stimuli and/or manipulations (If data are collected online, please specify the platform used, e.g. Qualtrics, Gorilla, LimeSurvey.

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1. Optional attachments:
2. Please explain the statistical or methodological or analytical approach you will use

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1. Will a method be used that may, by coincidence, lead to findings of which the participant should be informed? If so, what actions will be taken in the case of a coincidental finding?:
	* Yes
* No

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# [Participants](#_bookmark3)

1. What possible risks could participating in the study hold for participants?

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1. What measures are implemented to minimize risk for participants?

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1. How does the burden on the participants compare to the study's potential scientific contribution (theory formation, practical usability)?:

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1. What is the number of participants? Provide a power analysis and/or motivation for the number of participants. The current convention is a power of 0.80. If the study deviates from this power, the FERB would like you to justify why this is necessary:

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1. Age of the participants? Please specify a range in years: minimum - maximum

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1. How will the participants be recruited

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1. Please state any specific in- and exclusion criteria and how these are tested.

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1. How much time will the prospective participants have to decide as to whether they will indeed participate in the study?:

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1. Are the participants fully free to participate and terminate their participation whenever they want and without stating their grounds for doing so?
	* Yes
* No
1. Will the participants be in a dependent relationship with the researcher?
	* Yes
* No
1. Is there an independent contact person or a general email address of a complaint officer to whom the participant can contact?
	* Yes
* No
1. What time investment and effort will be requested from participants?

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1. Will the participants be compensated for their efforts? (Please consult the FAQ [Participant compensation](https://ferb.sites.uu.nl/relevant-documents/) [policy](https://ferb.sites.uu.nl/relevant-documents/) (under Guidelines > Local)
	* No, please explain why not
* Yes, standard recruitment. Please provide the amount according to the table in the FAQ Participant

compensation policy FSBS research and describe how payment is transferred.

* Yes, complex recruitment. Please provide the amount and provide a brief explanation of the chosen

amount. Please describe how payment is transferred.

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1. Will this compensation depend on certain conditions, such as the completion of the study?
	* Yes
* No
* Not applicable

# [Datamanagement](#_bookmark4)

1. Who will be responsible for managing access to the data?

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1. What type of data does the dataset contain? Please provide a brief description of the data, including the type, volume (if known), format and content.

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1. Will you be exchanging personal data with organizations/research partners outside the UU?
	* No
* Yes, with another research organization
* Yes, with an ICT supplier (for instance, software made by external party)
1. If personal data is shared, will a data processing agreement or data transfer agreement be made up?
	* Not applicable
* Yes
* No
1. Will standard minimum and maximum retention periods apply to the data?
	* Yes
* No, namely:

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1. Will data be collected and stored according to FSBS protocol? For the current version of the protocol, please see [this page](https://privacy-fss.sites.uu.nl/data-management/). Please explain how data collection and storage will be organized.
	* Yes
* No

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1. Is future use by yourself, other UU researchers or students foreseen?
	* Yes
* No
1. If so, where will you make your data available?

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1. If so, what access and usage conditions will apply? See: [FAIR data cheatsheets](https://www.uu.nl/en/research/research-data-management/tools/fair-cheatsheets-to-publish-your-research-data-and-software-fair)

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1. Will a possible transfer of personal data to a third country be part of this project?
	* No
* Yes, a country within the European Economic Area (EEA). Which country?
* Yes, a country outside the European Economic Area (EEA). Which country?

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# [Attachments](#_bookmark5)

1. Text (advert) for the recruitment of participants (optional)
2. Information letter for participant *(required)*
3. Consent form for participants *(required)*
4. Written or oral feedback information (debriefing text) (optional)
5. (Descriptions of) questionnaires (optional)
6. (Descriptions of) measurement instruments (optional)
7. Consent form for participants *(optional)*
8. Miscellaneous documents e.g. data set description (optional)