PRIDE QUESTIONNAIRE FOR APPROVAL SINGLE STUDY

(1.1) **Title**

(1.2) **Study type**

Single Study/project

(1.3) **Division**

(1.4) **Start date**

(1.5) **End date**

# Basic study information

1. Name(s), position(s) and division(s) of the responsible researcher(s): Name Position Division E-mail
2. Name(s), position(s) and division(s) of the executive researcher(s): Name Position Division E-mail
3. Research area/discipline:
4. What is the study's main objective (hypothesis)?
5. Primary funder of the study:
6. 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

# Study details (I)

1. Does your study exclusively concern the analysis of existing data, document or records? Where can the data be found?

* Yes
* No

1. Are the sources of the existing data, documents or records publicly available?

* Yes
* No

1. Not applicable. Will the data be processed by the principal investigator in such a manner that participants can be identified either directly or indirectly (through identifiers (such as a code) linked to them)?

* Yes
* No

1. If the study uses de-identified (or pseudonymized) data, does the responsible or executive researchers have access to the key to the code permitting re-identification of the person whose data are being studied?

* Yes
* No
* Not applicable

1. The research will involve only the use of anonymous survey procedures, interview procedures or the observation of public behavior

* Yes
* No

1. Will participants be asked to report their own or others' sexual experiences, alcohol or drug use, or suicidal thoughts, and will their identities be known to you?

* Yes
* No

# Study details (II)

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. Will participants that are recruited provide active informed consent?

* Yes
* No

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? (e.g., incapacitated, children, mentally challenged, traumatized, pregnant), or a taboo subject (e.g., own or others' sexual activity, hard drug use, suicide thoughts, religious belief, political preference)

* Yes
* No

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 personal health ○ ◉

Inquiries into criminal convictions and offences ○ ◉

Shocking images/videos ○ ◉

Deception (information letter does not state real study objective) ○ ◉

Physical pain (electrical/ thermal shocks, noise) ○ ◉

Following orders behaviorally (by force, or outside the context of the lab with possible harmful consequences for the participant or his/her social environment?)

○ ◉

A new technique for data collection? ○ ◉

1. Will data of the following categories be processed:

# Yes No

Photo data ○ ◉

Video data ○ ◉

Biological material (buccal, blood, hair) ○ ◉

# Yes No

Genetic data ○ ◉

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

Directly identifying data (name, address, date of birth or a combination of those items) ○ ◉

# Study details (III)

1. What is the study's theoretical and practical relevance? (500 words max.):
2. What are the central hypotheses?
3. What is the study's design and procedure? (500 words max.):
4. Optional attachments:

1. What data collection instruments, stimuli and/or manipulations will be used?
2. Optional attachments:
3. Please state which statistical procedures will be used.
4. 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

# Participants

(5) 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 :

1. How will the participants be recruited

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

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

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. What time investment and effort will be requested from participants?

1. Will the participants be compensated for their efforts? How (financial reimbursement, travelling expenses, otherwise). What is the amount?

* Yes
* No

1. Will this compensation depend on certain conditions, such as the completion of the study?

* Yes
* No

# Datamanagement

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

1. What type of data will you collect or create? Please provide a brief description of the data, including the type, volume (if known), format and content.

1. Will you be exchanging (personal) data with organizations/research partners outside the UU?

* Yes
* No

1. If so, will a data processing agreement be made up?

* Yes
* No
* Not applicable

1. Will standard minimum and maximum retention periods apply to the data?

* Yes
* No, namely:

1. Will data be collected and stored according to FSBS protocol? For the current version of the protocol, please see this page: https://ferb.sites.uu.nl/relevant-documents. Please explain how data collection and storage will be organized.

* Yes
* No

1. Is secondary use of your data is intended or foreseeable?

* Yes
* No

1. If so, where will you make your data available?

1. If so, what access and usage conditions will apply?

# Attachments

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