**PRIDE QUESTIONNAIR FOR APPRORVAL EXISTING DATA**

(1.1) **Title**

(1.2) **Study type**

Existing Data

(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
* Not applicable

1. 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. What are the terms and conditions regarding the use of the data? Did participants give consent for their data to be re-used? If not, on what basis is re-use of the data justified?

1. Please state what (if any) conditions the data archive imposes (e.g. registration, signing of confidentiality agreement, specific training etc.).
2. Do you intend to link two or more datasets? If YES, please give details of which datasets will be linked and for what purposes.

# Datamanagement

1. Who will be responsible for managing access to the data?
2. 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.
3. 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

1. NoIf so, where will you make your data available?
2. If so, what access and usage conditions will apply?

# [Attachments](#_bookmark0)

* Consent form for participants *(optional)*
* Miscellaneous documents e.g. data set description (optional)