
Researching the attitude on health, healthcare and eHealth in a low SES population

A Data Management Plan created using DMPonline

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Project abstract:

We are developing design principles that aim to support the design of eHealth interventions for people with a low socioeconomic status (SES). For the development of these guidelines, our first step is to understand how the target population perceives health, healthcare and eHealth. We will follow a Community Based Participatory Research (CBPR) approach, which entails the involvement of a community in all phases of the research. We will approach people with a low SES through a neighborhood community center in Rotterdam (Charlois). In total, we will be working with the community during the entire project (lasting 4 years). This application is for the first research phase, which entails the assessment of the community through observations and interviews. Our aim is to collect insights, stories and experiences on our target research question, i.e. the attitude on health, healthcare and eHealth in a low SES population. We will use these results to inform the design and setup of the next research phase within the CBPR, which will entail co-creation and participatory design activities. We aim to conduct this second study after 4 months and will apply for it at the HREC when our set-up is clear.

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General TU Delft data management questions

1. Is TU Delft the lead institution for this project?

- Yes, leading the collaboration

2. If you leave TU Delft (or are unavailable), who is going to be responsible for the data resulting from this project?

Valentijn Visch (V.T.Visch@tudelft.nl)

3. Where will the data (and code, if applicable) be stored and backed-up during the project lifetime?

- SURFdrive
- Project Storage at TU Delft

4. How much data storage will you require during the project lifetime?

- < 250 GB

5. What data will be shared in a research data repository?

- All data (and code) produced in the project

6. How much of your data will be shared in a research data repository?

- < 100 GB

7. How will you share your research data (and code)?

- I will upload the data to another data repository (please provide details below)
The data of this project will be uploaded for sharing on the DANS EASY database. This because the data that we are producing will mainly fall in the psychology/sociology/anthropology realm, which makes it applicable to the Social Science archive.

8. Does your research involve human subjects?

- Yes

9. Will you process any personal data? Tick all that apply (leave all unchecked if you do not process any personal data)

- Gender, date of birth and/or age

TU Delft questions about management of personal research data

1. Please detail what type of personal data you will collect, for what purpose, how you will store and protect that data, and who has access to the data.

Please provide your answer in the table below. Add an extra row for every new type of data processed:

Type of data	How will the data be collected?	Purpose of processing	Storage location	Who will have access to the data
Age	Interview Recording	To determine the age of the respondents to make appropriate correlations with the respondent's beliefs which is part of the research question	Project Drive	The project team (Promotor: Valentijn Visch, daily supervisor: Jos Kraal, and external partner: Rita van den Berg Emmons)
Gender	Interview Recording	To determine the gender of the respondents to make appropriate correlations with the respondent's beliefs which is part of the research question	Same as above	Same as above
Stories, experiences, insights	Interview Recording / field notes	To learn about the attitude of the community member towards health, healthcare, and eHealth.	same as above	same as above
Signed consent forms	Scanning and uploading the form	To record the consent of participants who agreed for their data processing	Same as above	Same as above

2. Will you be sharing personal data with individuals/organisations outside of the EEA (European Economic Area)?

- No

3. What is the legal ground for personal data processing?

- Informed consent - please describe the informed consent procedures you will follow
Before conducting the interviews, the participants will be taken through the project explanation and the informed consent form. Subsequently, they are asked to tick the boxes that they agree with / understand.
The forms are scanned and uploaded to the TU Delft project drive. The physical forms will be destroyed.

4. Will the personal data be shared with others after the end of the research project, and if so, how and for what purpose?

No

5. Does the processing of the personal data results in a high risk to the data subjects?

If the processing of the personal data results in a high risk to the data subjects, it is required to perform a Data Protection Impact Assessment (DPIA). In order to determine if there is a high risk for the data subjects, please check if any of the options below that are applicable to the processing of the personal data during your research (check all that apply).

If two or more of the options listed below apply, you will have to [complete the DPIA](#). Please get in touch with the privacy team: privacy-tud@tudelft.nl to receive support with DPIA. If only one of the options listed below applies, your project might need a DPIA. Please get in touch with the privacy team: privacy-tud@tudelft.nl to get advice as to whether DPIA is necessary.

If you have any additional comments, please add them in the box below.

- None of the above apply