# FWO DMP Template - Flemish Standard Data Management Plan

# Version KU Leuven

Project supervisors (from application round 2018 onwards) and fellows (from application round 2020 onwards) will, upon being awarded their project or fellowship, be invited to develop their answers to the data management related questions into a DMP. The FWO expects a **completed DMP no later than 6 months after the official start date** of the project or fellowship. The DMP should not be submitted to FWO but to the research co-ordination office of the host institute; FWO may request the DMP in a random check.

At the end of the project, the **final version of the DMP** has to be added to the final report of the project; this should be submitted to FWO by the supervisor-spokesperson through FWO’s e-portal. This DMP may of course have been updated since its first version. The DMP is an element in the final evaluation of the project by the relevant expert panel. Both the DMP submitted within the first 6 months after the start date and the final DMP may use this template.

The DMP template used by the Research Foundation Flanders (FWO) corresponds with the Flemish Standard Data Management Plan. This Flemish Standard DMP was developed by the Flemish Research Data Network (FRDN) Task Force DMP which comprises representatives of all Flemish funders and research institutions. This is a standardized DMP template based on the previous FWO template that contains the core requirements for data management planning. To increase understanding and facilitate completion of the DMP, a standardized **glossary** of definitions and abbreviations is available via the following [link](https://www.fwo.be/media/1024841/glossary-flemish-standard-data-management-plan.pdf).

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| 1. **General Project Information** | |
| Name Grant Holder & ORCID | Femke Konings (https://orcid.org/0000-0002-2034-6557) |
| Contributor name(s) (+ ORCID) & roles | Laura Vandenbosch (https://orcid.org/0000-0001-6834-8386), supervisor |
| Project number [[1]](#footnote-1) & title | PDMT2/24/012, Conceptualizing Mobile Dating Literacy: Unravelling Audience Group-and (Social-)Contextual Variances, Socializing Processes, and Well-being Outcomes through a Multimethod Approach |
| Funder(s) GrantID [[2]](#footnote-2) | KU Leuven Research Council - PDM Type 2 |
| Affiliation(s) | ☐ KU Leuven  ☐ Universiteit Antwerpen  ☐ Universiteit Gent  ☐ Universiteit Hasselt  ☐ Vrije Universiteit Brussel  ☐ Other:  ROR identifier KU Leuven: 05f950310 |
| Please provide a short project description | Mobile Dating Apps (MDAs) are popular among youth and facilitate the search for potential suitors. My Ph.D. revealed varying experiences among users, with some flourishing and others struggling. Currently users lack guidance in their MDA use, hence, this project proposes to introduce Mobile Dating Literacy (MDL) to empower users in navigating MDAs to enhance their relational (e.g., positive sexuality) and psychological wellbeing (e.g., positive affect). Special attention will be paid to disparities in MDL levels across different audience groups (i.e., age, gender, SES) and (social) contexts (i.e., sexual arousal, peer copresence), as well as to the socialization processes influencing MDL development. Multimethod research will involve MDL scale development research, cross-sectional survey research (*n* = 720) to examine group differences in MDL levels (trait), and daily diary (ESM) research (*n* = 350) to understand MDL activation across different contexts and interactions (state). |

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| 1. **Research Data Summary** | |
| List and describe all datasets or research materials that you plan to generate/collect or reuse during your research project. For each dataset or data type (observational, experimental etc.), provide a short name & description (sufficient for yourself to know what data it is about), indicate whether the data are newly generated/collected or reused, digital or physical, also indicate the type of the data (the kind of content), its technical format (file extension), and an estimate of the upper limit of the volume of the data [[3]](#footnote-3).   |  |  |  |  |  |  |  |  | | --- | --- | --- | --- | --- | --- | --- | --- | |  | | | | *Only for digital data* | *Only for digital data* | *Only for digital data* | *Only for physical data* | | Dataset Name | Description | New or Reused | Digital or Physical | Digital Data Type | Digital Data Format | Digital Data Volume (MB, GB, TB) | Physical Volume | | In-depth interviews | Focus group interviews will be conducted among middle-late adolescents (14–17 years old), and emerging adults (18–30 years old). I will conduct in-depth interviews until the point of saturation is reached. This is expected to be around 15 participants. | Generate new data  Reuse existing data | Digital  Physical | Audiovisual  Images  Sound  Numerical  Textual  Model  Software  Other:  **Sound**: The in-depth interviews will be recorded in order to make data transcription possible. These recordings will be deleted after the interviews have been transcribed and no identifying data will be included in the interview transcripts. | **Sound**: Interview recordings will consist of audio recordings in MP3 format. **Textual**: The interviews will be transcribed in Word format. **Software:** an Nvivo project will be created (.nvp). | < 1 GB  < 100 GB  < 1 TB  < 5 TB  > 5 TB  NA  **Sound**: We expect to collect max 15 interviews of 1 hour. This will result in a volume of audio recordings below 1GB  **Textual**: The Word forms of the transcribed interviews are expected to be below 1GB. **Software**: The Nvivo project is expected to be below 1GB. | / | | Cross-sectional survey | A cross-sectional online survey will be conducted among a representative sample of Flemish middle-late adolescents (14-17 years old), and emerging adults (18-30 years old). We expect to recruit 1000 participants, proportionally distributed according to ages (14-30), genders, and socio-economic statuses. | Generate new data  Reuse existing data | Digital  Physical | Audiovisual  Images  Sound  Numerical  Textual  Model  Software  Other:  **Textual**: Participants will receive 1 survey. This survey will initially be formulated in Word but will be sent out using specific software (see below). At the beginning of the survey, participants will have to fill in an online informed consent form. The parents of the participants will receive the (passive/active) informed consent form via mail (PDF).  **Software**: The surveys will be sent out using Limesurvey, as software that follows the GDPR protocols. After data collection, data will be downloaded from Limesurvey and analysed using R. | **Textual**: The surveys and informed consents will be in Word/PDF format.  **Software**: Data will be downloaded and implemented in R in .csv format. | < 1 GB  < 100 GB  < 1 TB  < 5 TB  > 5 TB  NA  **Textual**: The survey and informed consents are expected to be below 1GB. **Software**: The data file of the cross-sectional survey study (.csv) is expected to be between 1 and 5 GB. | / | | Experience sampling method (ESM) study | A 14-day ESM study will be conducted among a representative sample of Flemish middle-late adolescents (14–17 years old), and emerging adults (18–30 years old). We expect to recruit 350 participants. Participants will receive 1 background survey at the beginning of the study, followed by one daily survey during the subsequent 14-day study. | Generate new data | Digital  Physical | Audiovisual  Images  Sound  Numerical  Textual  Model  Software  Other:  **Textual**: Participants will receive 1 background survey and one daily survey each day. These surveys will initially be formulated in Word but will be sent out using specific software (see below). At the beginning of the background survey, participants will have to fill in an online informed consent form. The parents of the participants will receive the (passive/active) informed consent form via mail (PDF).  **Software**: The surveys will be sent out using Avicenna, an app that allows real-time monitoring of participants and follows the GDPR protocols. After data collection, data will be downloaded from the Avicenna app and analysed using R. | **Textual**: The surveys and informed consents will be in Word/PDF format.  **Software**: Data will be downloaded and implemented in R in .csv format. | < 1 GB  < 100 GB  < 1 TB  < 5 TB  > 5 TB  NA  **Textual**: The surveys and informed consents are expected to be below 1GB. **Software**: The data file of the ESM study (.csv) is expected to be between 1 and 5 GB. | / | | |
| *Guidance:*  *The data description forms the basis of your entire DMP, so make sure it is detailed and complete. It includes digital and physical data and encompasses the whole spectrum ranging from raw data to processed and analysed data including analysis scripts and code. Physical data are all materials that need proper management because they are valuable, difficult to replace and/or ethical issues are associated.* *Materials that are not considered data in an RDM context include your own manuscripts, theses and presentations; documentation is an integral part of your datasets and should described under documentation/metadata.*  [*RDM Guidance on data*](https://www.kuleuven.be/rdm/en/guidance/data-standards) | |
| If you reuse existing data, please specify the source, preferably by using a persistent identifier (e.g. DOI, Handle, URL etc.) per dataset or data type. | Recruitment for adolescents (14-17 years old) will rely on a secondary dataset of adolescent MDA users previously collected by the Media Psychology Lab. Parents of adolescents who participated in the prior research will be asked to provide consent for their children’s participation. These parents will not be informed if their child was identified as a mobile dater in the previous research. After obtaining consent, adolescents who self-identified as mobile daters will be invited to join the new study to explore their mobile dating experiences. |
| Are there any ethical issues concerning the creation and/or use of the data  (e.g. experiments on humans or animals, dual use)? If so, refer to specific datasets or data types when appropriate and provide the relevant ethical approval number. | Yes, human subject data; provide SMEC or EC approval number: currently being processed  Yes, animal data; provide ECD reference number:  Yes, dual use; provide approval number:  No  Additional information: Data will be collected among young individuals aged 12 to 25. No special categories of personal data will be collected and special attention will be paid to the privacy/anonymity of participants. Ethical approval has been obtained. |
| Will you process personaldata*[[4]](#footnote-4)*? If so, please refer to specific datasets or data types when appropriate and provide the KU Leuven or UZ Leuven privacy register number (G or S number). | Yes (provide PRET G-number or EC S-number below)  No  Additional information: We will work with pseudonymised/anonymized data and have received ethical approval for this via the PRET platform (currently being processed). In this approval application, we thoroughly explain how we will pseudonymise the data and which particular safety measures will be taken.  **Focus group interviews:** The in-depth interviews will be recorded in order to make transcription and data analysis possible. Whilst transcribing these recordings, we will anonymize them right away (only fictive name, gender, and age will be kept). To create the fictional name, we will store participants' names in a separate, password protected file. This file can only be accessed by the researchers working on this project and will not be distributed to third parties. When the interviews are transcribed, this file will be deleted.  **Cross-sectional survey:** we will provide each participant with a unique identification code. The identification codes will only be used by the researchers. All information that allows identification of the participants (e.g., email address) will be kept in a separate data file that will be encrypted and can only be accessed by a password known by the primary researcher. Furthermore, this file will be deleted as soon as the research project is completed.  **ESM study**: We will provide each participant with a unique identification code to link the background questionnaire to the daily dairy checklists. The identification codes will only be used by the researchers and solely for the purpose of linking participants’ data over different data collection points. All information that allows identification of the participants (e.g., email address) will be kept in a separate data file that will be encrypted and can only be accessed by a password known by the primary researcher. Furthermore, this file will be deleted as soon as the research project is completed. |
| Does your work have potential for commercial valorization (e.g. tech transfer, for example spin-offs, commercial exploitation, …)?  If so, please comment per dataset or data type where appropriate. | Yes  No  If yes, please comment: |
| Do existing 3rd party agreements restrict exploitation or dissemination of the data you (re)use (e.g. Material/Data transfer agreements, research collaboration agreements)?  If so, please explain to what data they relate and what restrictions are in place. | Yes  No  If yes, please explain: |
| Are there any other legal issues, such as intellectual property rights and ownership, to be managed related to the data you (re)use?  If so, please explain to what data they relate and which restrictions will be asserted. | Yes  No  If yes, please explain: |

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| 1. **Documentation and Metadata** | |
| Clearly describe what approach will be followed to capture the accompanying information necessary to keep **data understandable and usable**, for yourself and others, now and in the future (e.g. in terms of documentation levels and types required, procedures used, Electronic Lab Notebooks, README.txt files, Codebook.tsv etc. where this information is recorded).  [*RDM guidance on documentation and metadata*](https://www.kuleuven.be/rdm/en/guidance/documentation-metadata)*.* | **In-depth interviews**:   * We will provide Word documents with the meaning of the codes. * We will make use of an Nvivo project to generate documentation and metadata. * We will follow the guidelines as described in the following link 🡪 <https://dam.ukdataservice.ac.uk/media/622387/ukda-datamanagement-nvivo.pdf>   **Cross-sectional survey:**   * We will provide a codebook with the naming of the variables, the meaning of the values, and the labels in order to interpret the dataset. This will be generated using SPSS. * R code to analyse our data will be stored in specific R scripts.   **ESM study:**   * We will provide a codebook with the naming of the variables, the meaning of the values, and the labels in order to interpret the dataset. This will be generated using SPSS. * R code to analyse our data will be stored in specific R scripts.   For all studies, we will publish our materials within our organisation via the shared drive and outside our organization by making use of the Open Science Framework (OSF). We will develop our documents with a special eye for transparency and reproducibility. |
| Will a metadata standard be used to make it easier to **find and reuse the data**?  If so, please specify which metadata standard will be used. If not, please specify which metadata will be created to make the data easier to find and reuse.  *Repositories could ask to deliver metadata in a certain format, with specified ontologies and vocabularies, i.e. standard lists with unique identifiers.* | Yes  No  If yes, please specify (where appropriate per dataset or data type) which metadata standard will be used:  If no, please specify (where appropriate per dataset or data type) which metadata will be created:  We will make use of the personal (I-drive), shared KU Leuven drive (J-drive) and KU Leuven Onedrive to make our research project visible within our organization. In addition, the Open Science Framework (OSF) will be used as a repository for all materials that can be publicly shared, including code books, anonymized data, and code for analyses. |

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| 1. **Data Storage & Back-up during the Research Project** | |
| Where will the data be stored?  *Consult the*[*interactive KU Leuven storage guide*](https://icts.kuleuven.be/storagewijzer/en)*to find the most suitable storage solution for your data.* | Shared network drive (J-drive)  Personal network drive (I-drive)  OneDrive (KU Leuven)  Sharepoint online  Sharepoint on-premis  Large Volume Storage  Digital Vault  Other: OSF Framework |
| How will the data be backed up?  *What storage and backup procedures will be in place to prevent data loss?* | Standard back-up provided by KU Leuven ICTS for my storage solution  Personal back-ups I make (specify)  Other (specify) |
| Is there currently sufficient storage & backup capacity during the project? If yes, specify concisely. If no or insufficient storage or backup capacities are available, then explain how this will be taken care of. | Yes  No  If no, please specify: |
| How will you ensure that the data are securely stored and not accessed or modified by unauthorized persons?  *clearly describe the measures (in terms of physical security, network security, and security of computer systems and files) that will be taken to ensure that stored and transferred data are safe.*  [*Guidance on security for research data*](https://icts.kuleuven.be/storagewijzer/en) | All data will be encrypted and can only be accessed by a password known by the researchers working on the project. Data will be stored on the networks provided by KU Leuven, which provide high security. |
| What are the expected costs for data storage and backup during the research project? How will these costs be covered? | No expected costs. |

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| **5. Data Preservation after the end of the Research Project** | |
| Which data will be retained for at least five years (or longer, in agreement with other retention policies that are applicable) after the end of the project? In case some data cannot be preserved, clearly state the reasons for this  (e.g. legal or contractual restrictions, storage/budget issues, institutional policies...).  [*Guidance on data preservation*](https://icts.kuleuven.be/storagewijzer/en) | ​​ All data will be preserved for 10 years according to KU Leuven RDM policy  All data will be preserved for 25 years according to CTC recommendations for clinical trials with medicinal products for human use and for clinical experiments on humans  Certain data cannot be kept for 10 years (explain)  The recordings from the in-depth interviews will be deleted as soon as all interviews are transcribed. |
| Where will these data be archived (stored and curated for the long-term)?  [*Dedicated data repositories*](https://www.kuleuven.be/rdm/en/policy)*are often the best place to preserve your data. Data not suitable for preservation in a repository can be stored using a KU Leuven storage solution, consult the*[*interactive KU Leuven storage guide*](https://www.kuleuven.be/rdm/en/guidance/data-sharing)*.* | KU Leuven RDR  Large Volume Storage (longterm for large volumes)  Shared network drive (J-drive)  Other (specifiy): |
| What are the expected costs for data preservation during the expected retention period? How will these costs be covered? | No expected costs. |

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| **6. Data Sharing and Reuse** | |
| Will the data (or part of the data) be made available for reuse after/during the project?  Please explain per dataset or data type which data will be made available.  *Note that ‘available’ does not necessarily mean that the data set becomes openly available, conditions for access and use may apply. Availability in this question thus entails both open & restricted access. For more information:* [*https://wiki.surfnet.nl/display/standards/info-eu-repo/#infoeurepo-AccessRights*](https://wiki.surfnet.nl/display/standards/info-eu-repo/#infoeurepo-AccessRights) | Yes, as open data  Yes, as embargoed data (temporary restriction)  Yes, as restricted data (upon approval, or institutional access only)  No (closed access)  Other, please specify: |
| If access is restricted, please specify who will be able to access the data and under what conditions. | Fully anonymized data will be provided on the Open Science Framework to encourage open science practices. |
| Are there any factors that restrict or prevent the sharing of (some of) the data (e.g. as defined in an agreement with a 3rd party, legal restrictions)? Please explain per dataset or data type where appropriate. | Yes, privacy aspects  Yes, intellectual property rights  Yes, ethical aspects  Yes, aspects of dual use  Yes, other  No  If yes, please specify: All data will be fully anonymized before making them available. No identifying information will be kept and the demographic variables (gender, age, socio-economic status) will not be sufficient to identify the respondents should anyone try to do so. |
| Where will the data be made available?  If already known, please provide a repository per dataset or data type. | KU Leuven RDR  Other data repository (specify)  Other (specify)  Open Science Framework (OSF) for both the transcripts of the focus group interviews and the data derived from the ESM study. |
| When will the data be made available? | Upon publication of research results  Specific date (specify)  Other (specify) |
| Which data usage licenses are you going to provide? If none, please explain why.  *A data usage license indicates whether the data can be reused or not and under what conditions. If no licence is granted, the data are in a grey zone and cannot be legally reused. Do note that you may only release data under a licence chosen by yourself if it does not already fall under another licence that might prohibit that.*  *Check the*[*RDR guidance on licences*](https://www.kuleuven.be/rdm/en/rdr/licenses)*for data and software sources code or consult the*[*License selector tool*](https://ufal.github.io/public-license-selector/)*to help you choose.* | CC-BY 4.0 (data)  Data Transfer Agreement (restricted data)  MIT licence (code)  GNU GPL-3.0 (code)  Other (specify)  We will make use of the following license (https://creativecommons.org/licenses/by-nc-nd/4.0/legalcode.en). CC BY-NC-ND 4.0 only allows people to download and share our work for no commercial gain and for no other purposes. |
| Do you intend to add a PID/DOI/accession number to your dataset(s)? If already available, please provide it here.  *Indicate whether you intend to add a persistent and unique identifier in order to identify and retrieve the data.* | Yes, a PID will be added upon deposit in a data repository  My dataset already has a PID  No  A DOI will be attached to our data once collected and made available. |
| What are the expected costs for data sharing? How will these costs be covered? | No costs are expected for sharing as both KU Leuven RDR and OSF are free. |

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| **7. Responsibilities** | |
| Who will manage data documentation and metadata during the research project? | Data preservation for this project will be managed by Postdoctoral Researcher, Dr. Femke Konings, under the supervision of supervisor Prof. Dr. Laura Vandenbosch. |
| Who will manage data storage and backup during the research project? | The data storage and backup will be managed by Femke Konings (postdoctoral researcher on this project). |
| Who will manage data preservation and sharing? | Data preservation and sharing will be managed by Femke Konings (postdoctoral researcher on this project). |
| Who will update and implement this DMP? | The postdoctoral researcher of this project, named Femke Konings, will update and implement this DMP. |

1. “Project number” refers to the institutional project number. This question is optional. Applicants can only provide one project number. [↑](#footnote-ref-1)
2. Funder(s) GrantID refers to the number of the DMP at the funder(s), here one can specify multiple GrantIDs if multiple funding sources were used. [↑](#footnote-ref-2)
3. Add rows for each dataset you want to describe. [↑](#footnote-ref-3)
4. See Glossary Flemish Standard Data Management Plan [↑](#footnote-ref-4)