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.

	1. General Project Information
Name Grant Holder & ORCID	Dieter Stiers, 0000-0001-7242-8477
Contributor name(s) (+ ORCID) & roles	
Project number ¹ & title	Resurrecting Turnout, Saving Democracy?
Funder(s) GrantID ²	12A1Z24N
Affiliation(s)	X KU Leuven
	☐ Universiteit Antwerpen
	☐ Universiteit Gent
	☐ Universiteit Hasselt
	□ Vrije Universiteit Brussel
	□ Other:
	ROR identifier KU Leuven: 05f950310
Please provide a short project description	While there is high public support for electoral democracy worldwide, electoral turnout has been decreasing strongly in the past decades. This apparent contradiction leads to inequality between who turns out and who does not, and it also likely lowers people's perceptions of the legitimacy of elections. It is therefore unsurprising that political researchers and practitioners have looked for ways to increase turnout, but so far we know very little about the broader effects of these interventions. This project has as its main aim to investigate the effects of interventions to increase turnout on people's political attitudes and behaviour, and on their perception of the legitimacy of the electoral result. I rely on a unique electoral situation that presents itself in 2024 in Belgium in which two Election Days are held in the same year, each time presenting a new electoral rule, to examine the effects of lowering the voting age to 16 years old and compulsory voting on people's political attitudes using a more compelling design than previous research. I also include two novel conjoint experiments to investigate the effect of (interventions to increase) turnout on perceptions of the legitimacy of elections. Taken together, this project will reveal whether declining levels of turnout endanger the stability of representative democratic systems, and the extent to which interventions to increase turnout affect the legitimacy of elections.

¹ "Project number" refers to the institutional project number. This question is optional. Applicants can only provide one project number.

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.

2. 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.

				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,	Physical Volume
Electoral quality and legitimacy	Combination of existing survey datasets that measure satisfaction with democracy, complemented with data from objective indicators	☐ Generate new data ☑ Reuse existing data	⊠ Digital □ Physical	☐ Audiovisual ☐ Images ☐ Sound ☑ Numerical ☐ Textual ☐ Model ☐ Software ☐ Other:	. dta	TB)	
Voting at 16 Leuven- Ghent	Original dataset gathered among adolescents in Leuven and Ghent	☑ Generate new data☐ Reuse existing data	⊠ Digital □ Physical	☐ Audiovisual ☐ Images ☐ Sound ☑ Numerical ☐ Textual ☐ Model ☐ Software ☐ Other:	. dta		
Compulsory voting	Original dataset of a	□ Generate new data	☑ Digital☐ Physical	☐ Audiovisual ☐ Images	. dta	⊠ < 1 GB □ < 100 GB	

³ Add rows for each dataset you want to describe.

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		representative	☐ Reuse exist	ing		\square Sound		□ < 1 TB		
		sample in	data			Numerical		□ < 5 TB		
		Belgium				☐ Textual		□ > 5 TB		
						☐ Model		\square NA		
						☐ Software				
						\square Other:				
	Conjoint	Experimental	⊠ Generate n	ew	⊠ Digital	☐ Audiovisual	. dta	⊠ < 1 GB		
	experiments	data of a	data		□ Physical	☐ Images		□ < 100 GB		
		convenience	☐ Reuse exist	ing		\square Sound		□ < 1 TB		
		sample	data			Numerical		□ < 5 TB		
		gathered				☐ Textual		□ > 5 TB		
		through a				☐ Model		\square NA		
		survey company				\square Software				
						\square Other:				
_	GUIDANCE:									
		on forms the basis of	your entire DMI	, so make s	sure it is detail	led and complete. It in	cludes digital and phys	ical data and encompas	ses the whole spectrum	
			•			·		ıt need proper managen	•	
								nclude your own manusc		
	presentations; dod	cumentation is an int	egral part of you	r datasets a	and should des	scribed under documei	ntation/metadata.			
	<u>RDM Guidance on</u>	<u>data</u>								
										_
	•	ting data, please sp	•	I will reuse	e existing sur	vey data, but it is no	ot yet determined wh	nich data source I will u	use.	
	source, preferab	ly by using a persis	tent							
	identifier (e.g. D	OI, Handle, URL etc	.) per							
	dataset or data t	уре.								

Are there any ethical issues concerning the	☑ Yes, human subject data; provide SMEC or EC approval number:
creation and/or use of the data	☐ Yes, animal data; provide ECD reference number:
(e.g. experiments on humans or animals, dual	☐ Yes, dual use; provide approval number:
use)? If so, refer to specific datasets or data	□ No
types when appropriate and provide the	Additional information:
relevant ethical approval number.	Approval number for study voting at 16 Leuven-Ghent: G-2023-7066-R2(MAR)
	Approval numbers for other studies are not yet available.
Will you process personal data ⁴ ? If so, please	☑ Yes (provide PRET G-number or EC S-number below)
refer to specific datasets or data types when	□ No
appropriate and provide the KU Leuven or UZ	Additional information:
Leuven privacy register number (G or S number).	Approval number for study voting at 16 Leuven-Ghent: G-2023-7066-R2(MAR)
	Approval numbers for other studies are not yet available.
Does your work have potential for commercial	□ Yes
valorization (e.g. tech transfer, for example spin-	⊠ No
offs, commercial exploitation,)?	If yes, please comment:
If so, please comment per dataset or data type	
where appropriate.	
Do existing 3rd party agreements restrict	☐ Yes
exploitation or dissemination of the data you	⊠ No
(re)use (e.g. Material/Data transfer agreements,	If yes, please explain:
research collaboration agreements)?	
If so, please explain to what data they relate and	
what restrictions are in place.	
Are there any other legal issues, such as	☐ Yes
intellectual property rights and ownership, to be	⊠ No
managed related to the data you (re)use?	If yes, please explain:
If so, please explain to what data they relate and	
which restrictions will be asserted.	

⁴ See Glossary Flemish Standard Data Management Plan

3. Documentation and Metadata Clearly describe what approach will be followed Each original dataset that I will gather, will be accompanied with the original questionnaire that serves as to capture the accompanying information codebook. This will be uploaded together with the datafile to Harvard's Dataverse. This upload includes keywords making it easier to find the dataset. The upload is also linked to an academic publication that necessary to keep data understandable and **usable**, for yourself and others, now and in the details all the data gathering procedures. future (e.g. in terms of documentation levels and types required, procedures used, Electronic Lab The dataset consists of variables that are clearly named and the values are labelled to indicate their Notebooks, README.txt files, Codebook.tsv etc. content. where this information is recorded). RDM guidance on documentation and metadata. Will a metadata standard be used to make it ☐ Yes \bowtie No easier to find and reuse the data? If yes, please specify (where appropriate per dataset or data type) which metadata standard will be used: If so, please specify which metadata standard will be used. If not, please specify which metadata will be created to make the data If no, please specify (where appropriate per dataset or data type) which metadata will be created: easier to find and reuse. All variables in the dataset will be named, and their values labelled. Besides the data, the original questionnaires will be stored. The data always include information on when exactly they were gathered. REPOSITORIES COULD ASK TO DELIVER METADATA IN A CERTAIN FORMAT, WITH SPECIFIED ONTOLOGIES AND VOCABULARIES, I.E. STANDARD LISTS WITH UNIQUE IDENTIFIERS.

4. Data Storage & Back-up during the Research Project

Where will the data be stored?	Shared network drive (J-drive)
	☐ Personal network drive (I-drive)
Consult the <u>interactive KU Leuven storage guide</u> to	☐ OneDrive (KU Leuven)
find the most suitable storage solution for your data.	☐ Sharepoint online
	☐ Sharepoint on-premis
	☐ Large Volume Storage
	☐ Digital Vault
	☐ Other:
How will the data be backed up?	☑ Standard back-up provided by KU Leuven ICTS for my storage solution
	☐ Personal back-ups I make (specify)
WHAT STORAGE AND BACKUP PROCEDURES WILL BE IN PLACE TO	☐ Other (specify)
PREVENT DATA LOSS?	
Is there currently sufficient storage & backup	⊠ Yes
capacity during the project? If yes, specify	□ No
concisely. If no or insufficient storage or backup	
capacities are available, then explain how this	If no, please specify:
will be taken care of.	
How will you ensure that the data are securely	During the fieldword, when sensitive data are only pseudonymized, the data are stored offline (on an external hard drive) following the requirements of the ethical board.
stored and not accessed or modified by	Once the data are anonymized, they will be stored on the J-drive of the university and
unauthorized persons?	on Box.
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	

What are the expected costs for data storage	
and backup during the research project? How	No additional costs are expected.
will these costs be covered?	

	5. Data Preservation after the end of the Research Project
Which data will be retained for at least five	☐ All data will be preserved for 10 years according to KU Leuven RDM policy ☐ All the control of the control
years (or longer, in agreement with other retention policies that are applicable) after the	☐ 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
end of the project? In case some data cannot be	☐ Certain data cannot be kept for 10 years (explain)
preserved, clearly state the reasons for this	
(e.g. legal or contractual restrictions,	
storage/budget issues, institutional policies).	
Guidance on data preservation	
Where will these data be archived (stored and	☐ KU Leuven RDR
curated for the long-term)?	☐ Large Volume Storage (longterm for large volumes)
	☐ Shared network drive (J-drive)
<u>Dedicated data repositories</u> are often the best place to preserve your data. Data not suitable for	☑ Other (specifiy): Harvard's Dataverse
preservation in a repository can be stored using a KU	
Leuven storage solution, consult the interactive KU	
<u>Leuven storage guide</u> .	
What are the expected costs for data	No costs are expected.
preservation during the expected retention period? How will these costs be covered?	
period: flow will these costs be covered?	

	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	 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: All original data will be shared on Harvard's Dataverse once the envisioned academic paper has been finished − i.e., it will be published alongside the research paper. The reused data is already openly available online.
If access is restricted, please specify who will be able to access the data and under what conditions.	
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: In the process of gathering the original data, data will be pseudonymized. Once the data collection is complete, all information that would allow identifying individual respondents will be deleted. The anonymized datasets will be shared openly.

Where will the data be made available? If already known, please provide a repository per dataset or data type. When will the data be made available?	 □ KU Leuven RDR □ Other data repository (specify) ☑ Other (specify) Harvard's Dataverse ☑ 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 quidance on licences for data and software sources code or consult the License selector tool to help you choose.	 □ CC-BY 4.0 (data) □ Data Transfer Agreement (restricted data) □ MIT licence (code) □ GNU GPL-3.0 (code) □ Other (specify)
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
What are the expected costs for data sharing? How will these costs be covered?	No additional costs are expected

	7. Responsibilities
Who will manage data documentation and metadata during the research project?	I will manage the documentation and metadata
Who will manage data storage and backup during the research project?	I will manage the data storage and backup during the research project
Who will manage data preservation and sharing?	I will manage the data preservation and sharing
Who will update and implement this DMP?	I will update and implement this DMP