Erasmus School of Health Policy & Management

# Research Data Management Awareness Campaign



# How to write a DMP?

ESHPM RDM Awareness Campaign

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October 21, 2021



#### Outline

- Why a Data Management Plan?
  - FAIR
- What's a Data Management Plan?
  - Templates & requirements
- How?
  - EUR DMP Template v4.4
  - DMPOnline



# Why, oh why?

### Maximizing the impact of scientific data

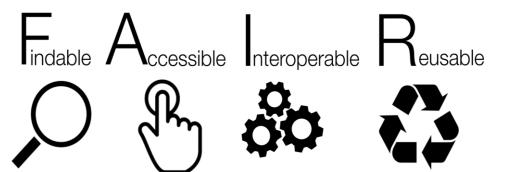




Research: making claims backed up by data

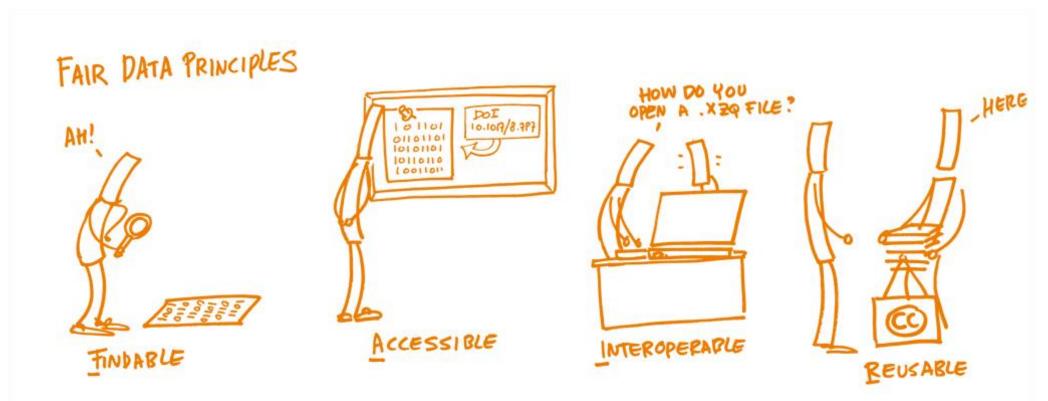
- ○Data = one of your main ingredients
- ○Data = one of your main outputs







#### Be FAIR: Findable, Accessible, Interoperable, Reusable



Picture: FAIR principles / Patrick Hochstenbach / CC0 1.0

- FAIR paper: Wilkinson et al., 2016. <a href="https://doi.org/10.1038/sdata.2016.18">https://doi.org/10.1038/sdata.2016.18</a>
- More resources at <a href="https://www.go-fair.org/">https://www.go-fair.org/</a>



#### Researcher's code



The European Code of Conduct for Research Integrity REVISED EDITION Research derives its status from the fact that it is a process governed by standards, which are expressed by a number of guiding principles. Researchers who are not guided by these principles risk harming quality and trustworthiness of research, undermine public trust and mutual trust between individual researchers.

- Honesty
- Scrupulousness
- Transparency
- Independence
- Responsibility



#### **Ethical research**

Research involving one of the following aspects:

- Human beings;
- Potential misuse of research results;
- (Special categories of) personal data;
- Potential conflict of interest;
- Non-EU countries;
- External stakeholders, among which funding organizations;
- Environment, health & safety issues, including potential harm to researchers.
- + if planning to publish in a (medical) journal.

#### **Ethical review**

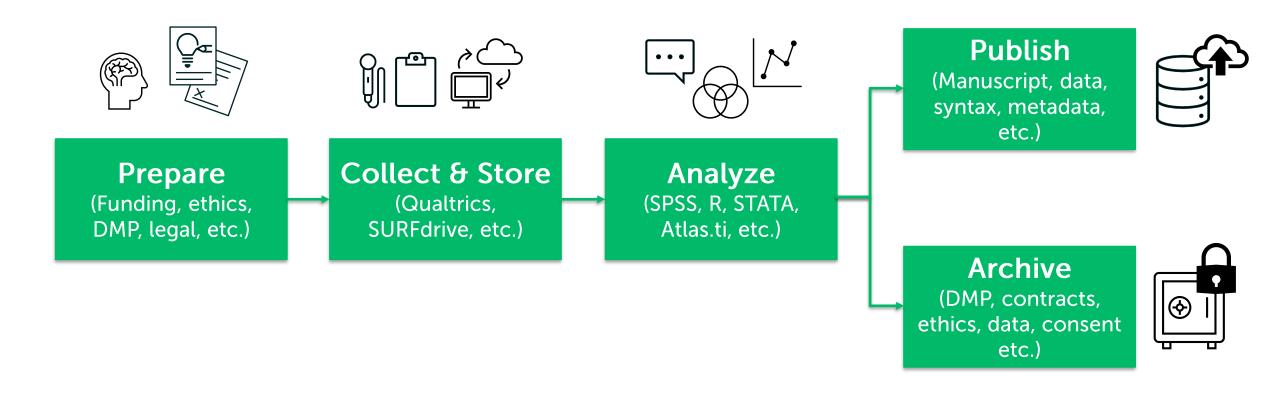
The Association of Universities in the Netherlands (VSNU) and the Royal Netherlands Academy of Arts and Sciences (KNAW) adopted the <u>Netherlands Code of Conduct for Research Integrity (NCCRI)</u> in 2018. In line with this document, Erasmus University Rotterdam underpins the importance of ethics review of research projects prior to data collection, and as of January 1st, 2021 a number of <u>principles and requirements</u> pertaining to research ethics review at Erasmus University Rotterdam has taken effect.

The aim of the ethics review is to ensure that all research that is conducted by faculty affiliated researchers, complies with good practice in terms of protecting the privacy and security of research participants in the collection, analysis, and dissemination of research data, and with the five principles of proper academic practice as set out in the NCCRI: honesty, scrupulousness, transparency, independence, and responsibility. This will advance the quality and reliability of the research.



# Wait, but what?

## The lifecycle of data in a research project





# 10 practices from Briney et al.

- Practice 1: Keep sufficient documentation
- Practice 2: Organize files and name them consistently
- Practice 3: Version the Files
- Practice 4: Create a security plan, when applicable
- Practice 5: Define roles and responsibilities
- Practice 6: Back up the data
- Practice 7: Identify tool constraints
- Practice 8: Close out the project
- Practice 9: Put the data in a repository
- Practice 10: Write these conventions down [in a DMP]

→ Document & organize

→ Stay safe

→ Plan

- → Prevent technical issues
- → Closing & archiving

(Zafus

## Prepare by writing a data management plan (DMP)



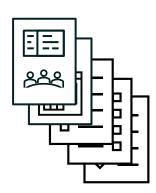
#### What a DMP is:

- OA way to formulate and record your plans and decisions about your data
- OA "living document", to be revisited every few months
- Saving you time
- OA tool for you to help with responsible RDM



#### What a DMP is not:

- A matter of "one-time approval"
- Written in stone
- Wasting your time
- A box to tick "and then RDM is taken care of"





### In a Data Management Plan you describe ...

- What type of data you are going to collect
- Where you are going to store your data
- How you are going to share your data
- How you address security & privacy matters

A DMP forces you to think through each step of your project making sure you have all you need



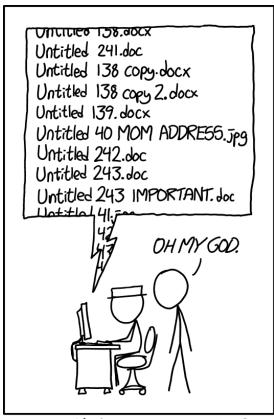
#### Write a Data Management Plan if ...

- You care deeply about research integrity and ...
- You want to prevent issues and set-backs
- You want to comply with funder and university policy



# Document and organize your research thoroughly



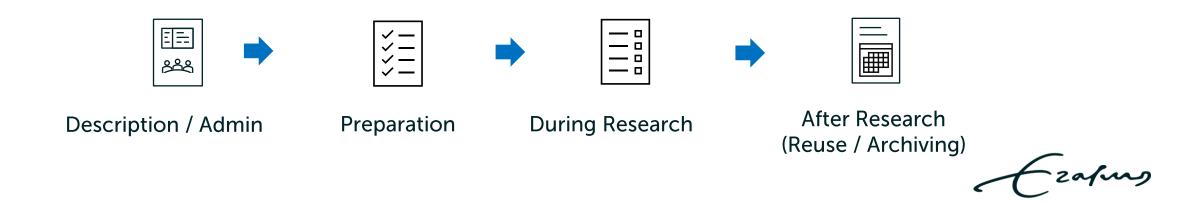


PROTIP: NEVER LOOK IN SOMEONE. ELSE'S DOCUMENTS FOLDER.



#### **DMP Templates**

- What does a DMP look like?
  - Many different DMP templates exist
  - I recommend to either use EUR template or your funder's template (if applicable)
    - EUR template is approved by NWO and ZonMw
- EUR DMP Template 4.4 (some outdated versions still in circulation!)
  - You can choose between offline version (Word document) or online tool (<u>DMPonline</u>)
  - Loosely structured in chronological order:



# How?

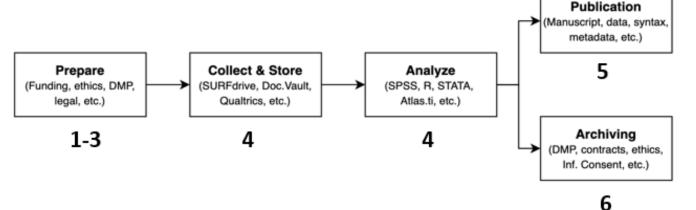
- EUR DMP Template
- Using DMPOnline

# EUR DMP Template

#### **EUR DMP Template**

#### **EUR Data Management Plan**

Version 4.4



The template was created by the RDM specialists at UL/EDSC and the data stewards. It has been approved by NWO and ZonMw, and it is also the default DMP format supported by the EUR.

#### Table of contents

- General
- 2. Administration and Project Description
- 3. Preparation: Legal Issues, Policy and Guidelines
- 4. During Research: Collecting and Analyzing
- 5. After Research: Data Sharing and Re-use
- After Research: Archiving



#### General

Please tick the following boxes if you agree	to act according to the following terms:
☐ I will answer all questions truthf	fully and to the best of my knowledge
$\ \square$ I will discuss the data manageme	ent plan with my research team
$\square$ I will check and, if necessary, up	date my data management plan a minimum of once a year
Support <sup>i</sup> in writing a data management plan	is available through the faculty Data Stewards (their contact
details can be found on the web page of Eras	smus Research Services).
If applicable, please provide the name of th	e support staff consulted and the date of consultation:
Name of support staff	
Date of consultation	
	ata Management Plans to be completed in consultation with data nof the grant holder in order to be eligible for consideration.
management support stant at the nome institution	To the grant house in order to be engine for consideration.
Scientific research must be conducted in line	e with existing guidelines on good research practices and
integrity. Please tick the boxes if you have r	read and understand these guidelines and will act accordingly.
☐ The Nath colored Code of Coods	art for Donorando Intermite (VCNIII 2019)
	ct for Research Integrity (VSNU, 2018)
	OR
☐ The European Code of Conduct f	for Research Integrity (ALLEA, 2017)



List the names and affiliations of all members of the research team. List the researcher responsible for research data management first. For PhD projects, please indicate the Promotor(s) and/or Daily Supervisor(s) with a (!)

Name	Email	ORCID*	Research Institution

## Administration & Project description

Project title	
Project start date as intended	
Project duration in months as intended	
Funding body (if applicable)*	
Grant number (if applicable)	
Date of DMP version 1	
Date of update DMP and version*	Date: Version:

#### Also questions on:

- Project background & research questions
- Research type & methodology
  - Be as detailed as possible!
- Resources needed to implement the DMP
  - 'Standard' services are provided by EUR



#### Legal & policy

		Who	
4.	Do you need to obtain ethical approval for your research project?*		
	□ No, my project does not require ethical approval	Example: Research Pai	rticipants
	Yes, I am preparing to submit my application	Example: Multiple res	
	☐ Yes, I have submitted my application		
	☐ Yes, I have obtained ethical approval	<ol><li>List the agreen you are reusing</li></ol>	
	☐ I do not know	Who	Туре
5.	If you have obtained ethical approval, list the reference number:		
		Example: EUR	RDM p
		Property of the Property	

<ul> <li>□ With nobody / No reason → Go to Q3</li> <li>□ With research participants</li> </ul>				
	king with multiple resea	arcn partners		
☐ I do not	know → Go to Q3			
2. List the agreem		<u>te</u> and with whom you wi	ll make them.	
Who	Туре	of agreement		
Example: Research Part	icipants Informe	ed Consent		
Example: Multiple resea		Consortium Agreement		
3. List the agreeme	ents or other data man	agement policies that you	ı need to uphold but <u>did not initia</u> t	<u>te</u> . If
you are reusing	existing data, list the to	erms of use under which y	you may re-use them.	
Who	Туре		Version and date	
Example: EUR		us University Rotterdam (EUR)	Version 1.0 [August 14, 2020]	
Example: Zoom	Terms of Service		Version [August 20, 2020]	
Example: NWO	Funding Agreement		Version 11.4 [July 11, 2018]	
Example: EUR	Internet and ICT facil	ities policy	Version 1.0 [2015]	

1. With whom will you need to make legal arrangements?\*

#### **During research: Collecting & Analyzing**

6. Specify what data you will be collecting and indicate format, estimated size, and whether this is data that you will be generating or existing data that you will be re-using.\*

Туре	Format	Estimated size	Generate or Reuse
Example: Digital survey data	.CSV	1-5 GB	Generate
Example: Audio-recorded interviews + Transcripts	.mp3; .docx/.odt	5-10 GB	Generate
Example: Dutch Central Bureau of Statistics (CBS) data	.csv	10-50 GB	Re-use



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re	rso	nal	LO	ata

- Data that can [in]directly identify a natural person
- Name, address, IP-address, student number

#### Sensitive personal data

- Personal data that can potentially harm an individual if it were to become public
- Special category data: health, political preferences, sexual preferences, religion, race.

□ No

#### Sensitive non-personal data

Data that is otherwise sensitive such a proprietary data, patents, company data

7.	Will you be collecting [sensitive] personal data?*
	□ No – My research does not include [human] participants → Go to Q9
	$\ \square$ No – My research involves human participants, but I will collect fully anonymous data $ o$ Go to
	☐ Yes – Personal data (non-sensitive) → Consult your faculty's Privacy Officer
	□ Yes – Personal data (sensitive) → Consult your faculty's Privacy Officer
	□ I do not know → Consult your faculty's Privacy Officer
8.	If you collect [sensitive] personal data, how will you protect the privacy of participants when sharing your data?*
	☐ I will fully anonymize the data
	☐ I will pseudonymize the data
	□ I do not know → Consult your faculty's Privacy Officer
9.	Will you be collecting non-personal sensitive data?
9.	will you be collecting non-personal sensitive data?
	☐ Yes [e.g. confidential company data, data related to national security]



10. Where will you store your data during the project?\* You can select multiple options.

All types of files – EUR
employees & students
All types – Researchers N
All types — Researchers Non-NL & Third parties
Audio & Video
All types of data – EUR
employees

Evamela: ELIR Veda: Cenela C Sriba

Capus

	□ EUR supported
	□ Private
	□ I do not know → Contact your faculty's Data Steward
12. Wh	at hardware and software do you use?* Select all applicable options.
	☐ EUR supported hardware (e.g. @wEURk laptop, @wEURk workstation)
	☐ Private hardware (e.g. personal laptop, private external hard drive)
	☐ EUR supported software as found in the <u>software catalog</u>
	☐ Private software or freeware (e.g. private Dropbox)
13. If y	ou use private hardware, software, or freeware, please specify what and for what reason:
	During fieldwork I have no access to internet, so I temporarily store my data on an encrypted external drive. I do not have an EUR laptop, so I am using my own device.
14. Are	regular backups made of your data?
	☐ I do not know
	☐ I do not know ☐ No
	□ No

Note: Include who makes the backups and how often backups are made

#### Use EUR supported hardware and software

- University & Faculty provide necessary conditions, such as IT infrastructure
- Your responsibility to use the tools being provided
  - Use EUR supported hardware and software (see <u>software catalog</u>)
  - Use <u>MyApps</u> (Remote Desktop) when working on own device
  - Use <u>Teams/Sharepoint</u>, <u>SurfDrive</u>, <u>ResearchDrive</u>, <u>Document Vault</u>, EUR Network shares to store and process research data











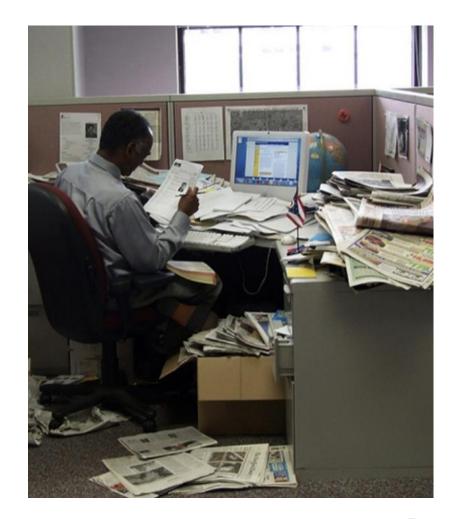








- Access to folder and files
  - Who manages access; who has access; how is access managed in case of staff changes or illness
- Structuring and naming folders and files
  - o In X time, would you [or someone else] be able to quickly find the right file and understand the content?
- Versioning of files
  - How many 'definitive' versions do you have of one document





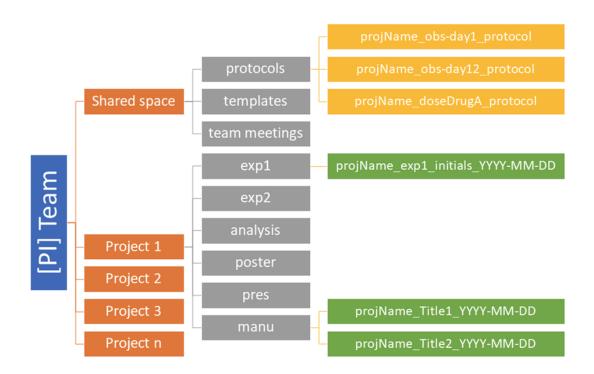
## Organize your data

- 1. Save your project in a single folder
- 2. Devise a logical system of subfolders
  - Make sure to have separate folders for data (raw vs. processed), documentation, figures
- 3. Introduce the project (or subfolder) in a README
- 4. Use interoperable file types
- 5. Use descriptive and logical file names
  - Machine readable
  - Human readable
  - Logically sortable (YYYYMMDD)
- 6. Make your data tidy (each variable is a column, each observation is a row)

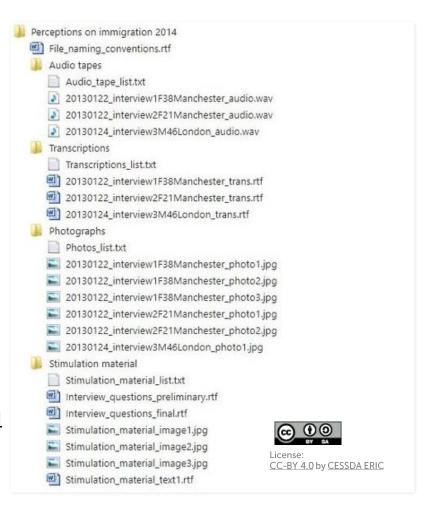


### Organize your data and use consistent filenames

- Organize files and name them consistently
  - Choose strategy + convention, and write these down



Refer to guidelines when designing your system: <u>CESSDA training</u>



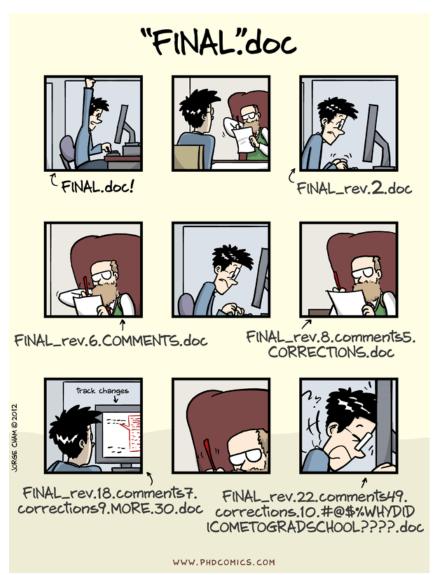
## Document your research thoroughly with a README

- Keep sufficient documentation
  - README.txt 'with enough detail so that the research can be picked up after a yearlong hiatus or by a collaborator'
  - Do not trust your memory!
- O What is the project about?
- Who are involved?
- O How was the data collected?
- When & where is the research conducted?
- Abbreviations, folder structure, file explanations, references & links





#### Use consistent filenames



- Version the Files
  - At minimum: separate raw + processed data
  - Use version numbers or dates (manually or automate using Git)
- Resist the temptation to ever use "FINAL" in a file name!



Picture: "Piled Higher and Deeper" by Jorge Cham / www.phdcomics.com

### After research: Data Sharing & Reuse

20. What data [and code] will be shared in a research data repository?

☐ I do not know
☐ All data [and code] underlying published papers / reports → Go to Q22
☐ All data [and code] produced in the project → Go to Q22
☐ A selection of the data [and code] → Go to Q21
☐ I cannot share the data [and code]; I will share only the metadata → Go to Q21
21. Please specify why you are unable to share (all) data [and code].

- The EUR is an Open Science university
  - All publications should be published open access
  - All data underlying publications should be shared as openly as possible, as closed as necessary
- Three questions: Why, How, Where
- Why: for your future self, for a possible reuse & collaborations
- How: complete and interpretable metadata and documentation
- Where: EUR Data Repository [EDR] or domain-specific repository



#### Tension between what you need to share and what you can share

- From the Netherlands Code of Conduct for Research Integrity (2018)
  - https://doi.org/10.17026/dans-2cj-nvwu

Ensure that, in accordance with the FAIR principles, data is open and accessible to the extent possible and remains confidential to the extent necessary.





#### Data sharing & reuse

22. List the data [and code] that you plan to share in a research data repository. Also list the information / documentation / metadata that you will include to make the data package self-explanatory and re-usable in the future (for other researchers and yourself)\*

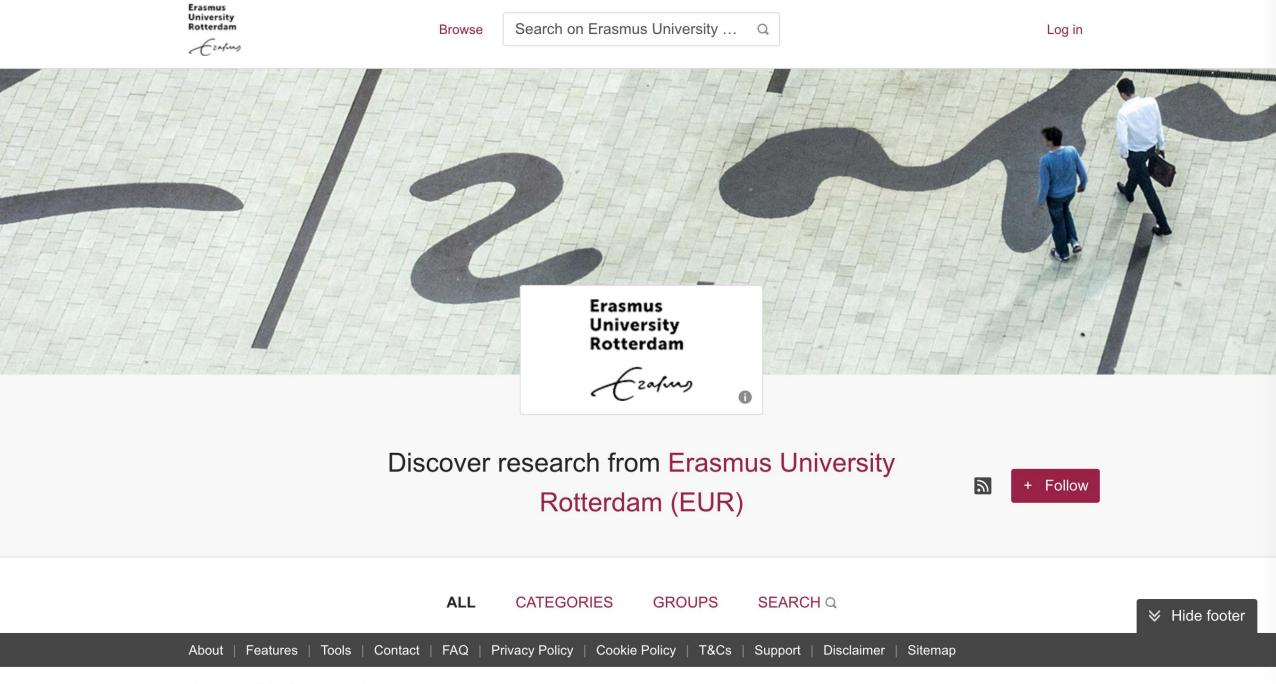
Data	Format	Size
Example: Anonymized survey data	.CSV	<1 GB
Example: Codebook, Blank questionnaire	.pdf	<1 GB
Example: readme text file (general description of the data, incl. date of collection, selection procedure of participants, tools used to collect the data, etc.)	.txt	<1 GB

- Information needed to interpret, understand, and reuse data
  - Codebook, methods, date of collection, definitions
  - Metadata standard: if we all used the same e.g. coding and definitions, life would be much easier!



# Data sharing & reuse

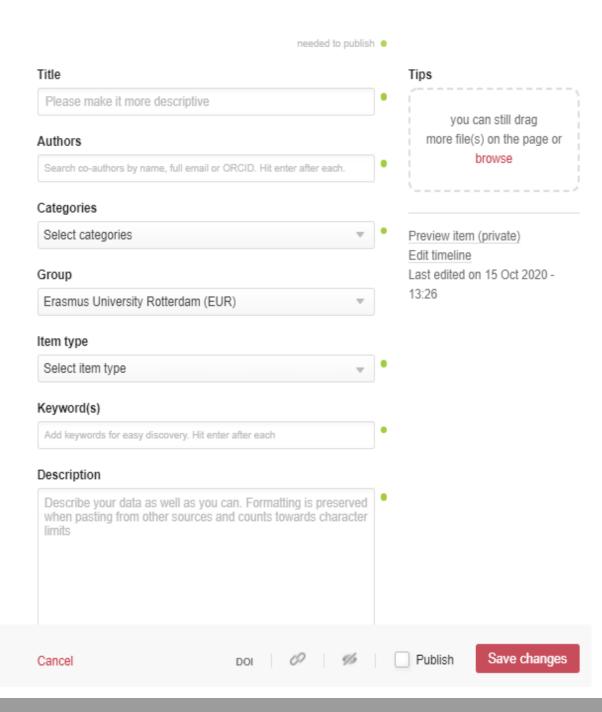
	23. In w	vhich repository will you place the [meta]data and/or code associated with your paper?*
		☐ I do not know
		☐ EUR Data Repository (EDR)
		☐ Other – please specify:
		Example: DANS EASY, Zenodo, Dryad, Dataverse, 4TU, Open Science Framework (OSF)
24. V	What metadata standard will you use to document your research?*	
	☐ I do not know	
	□ None	
	☐ <b>DCMI</b> (Dublin Core Metadata Initiative) Note: Default within the EUR Data Repository	
	☐ DDI (Data Documentation Initiative)	
	☐ SDMX (Statistics Data and Metadata Exchange)	
	☐ Other – please specify:	
		Fragues
	L	- Zajun)



# **EUR Data Repository**

### Metadata (Dublin Core):

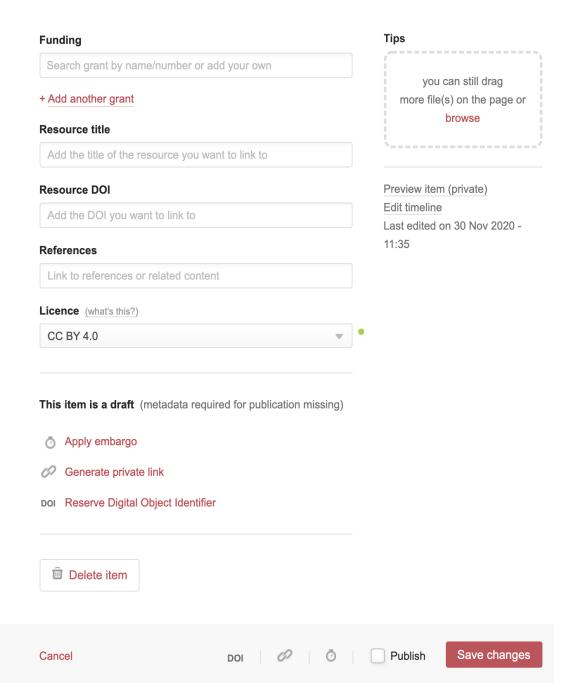
```
Title;
Authors;
Categories;
Group;
Item Type,
Keywords;
Description;
Funding;
Resource title;
Resource DOI;
References;
License
```



# **EUR Data Repository**

### Metadata (Dublin Core):

```
Title;
Authors;
Categories;
Group;
Item Type,
Keywords;
Description;
Funding;
Resource title;
Resource DOI;
References;
License
```



### Data sharing & reuse

After uploading data, you stay in control and can choose your data to be:

- Open to all
- Placed under embargo (temporary vs. permanent)
- Made restricted access:
  - Sharing restrictions: specified person(s), faculty, IP-range, etc.
  - o Confidential: accessible only upon request, user agreement can be set up

28. Und	28. Under which license will you make your data available for re-use?*		
	☐ I do not know		
	☐ Creative commons (e.g. CC-BY)		
	☐ License for specific types of data (e.g. software license)		
	☐ Other – please specify:		
29. Plea	29. Please specify which license:		

### After research: Archiving

- Closed archive 'cold storage', unlike e.g. EUR Data Repository
  - o Doubles as emergency backup; original files for auditing and verification
- Typical files for archiving include:
  - Informed Consent; Contracts; DMP; Ethics documents
  - Complete raw data; Code and Software; Codebook
- Per research domain the term may vary between 5 to 15+ years
  - The EUR recommends a minimum of 10 years



# **Archiving**

- Select what needs to be destroyed
  - Exemption for personal data under GDPR
- Determine what you want to archive
- Use durable formats

Туре	<ul> <li>Preferred format(s)</li> </ul>	<ul> <li>Non-preferred format(s)</li> </ul>	
Text documents	<ul><li>PDF/A (.pdf)</li><li>ODT (.odt)</li></ul>	<ul> <li>Microsoft Word (.doc)</li> <li>Office Open XML (.docx)</li> <li>Rich Text File (.rtf)</li> <li>PDF other than PDF/A (.pdf)</li> </ul>	
Plain text	<ul> <li>Unicode text (.txt)</li> </ul>	<ul> <li>Non-Unicode text (.txt)</li> </ul>	
Markup language	<ul> <li>XML (.xml)</li> <li>HTML (.html)</li> <li>Related files: .css, .xslt, .js, .es</li> </ul>	<ul><li>SGML (.sgml)</li><li>Markdown (.md)</li></ul>	
Programming languages	<ul><li>MATLAB</li><li>NetCDF</li><li>TextFabric</li></ul>		
Spreadsheets	<ul><li>ODS (.ods)</li><li>CSV (.csv)</li></ul>	<ul><li>Microsoft Excel (.xls)</li><li>Office Open XML Workbook (.xlsx)</li><li>PDF/A (.pdf)</li></ul>	

30. You	30. You may be obliged to destroy some data before archiving. Do any of such obligations apply to you?		
	☐ I do not know		
	□ No		
	☐ Yes - Contractual obligation (e.g. licenses)		
	☐ Yes - Commercial objectives		
	☐ Yes - Privacy law (e.g. personal data of participants)		
	☐ Yes - Other - please specify:		

#### 31. List the data you will be archiving. These data constitute your archival package.

Data	Format	Size

Example: Informed Consent forms (signed)	.pdf	<1 GB
Example: Raw data	.csv, .docx, mp3, mp4	5-10 GB
Example: Processed data	.csv, .odt, .mka	1-5 GB
Example: Questionnaires	.pdf	<1 GB
Example: Contracts & Terms of use	.pdf	<1 GB
Example: Data Management Plan	.pdf	<1 GB
Example: Grant application & project description	.pdf	<1 GB
Example: Ethical review application & approval document	.pdf	<1 GB

# **DMPOnline**

### **DMPOnline**



Public DMPs DMP Templates



#### Welcome

DMPOnline-Erasmus helps you to create, review, and share data management plans that meet institutional and funder requirements.

DMPonline is developed by the UK's Digital curation centre (DCC)and the University of California Curation Center (UC3) as a shared resource for the research community.

You can download funder templates without logging in, but the tool provides tailored guidance and example answers from funders and your organisation.

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More information can be found at this url: https://www.eur.nl/en/campus/university-library/erasmusdata-service-centre/research-data-management-rdm/data-management

If you have questions don't hesitate and contact the Datateam of the Erasmus Data Service Centre (EDSC) <edsc@eur.nl>



17,622 Users



203 Organisations

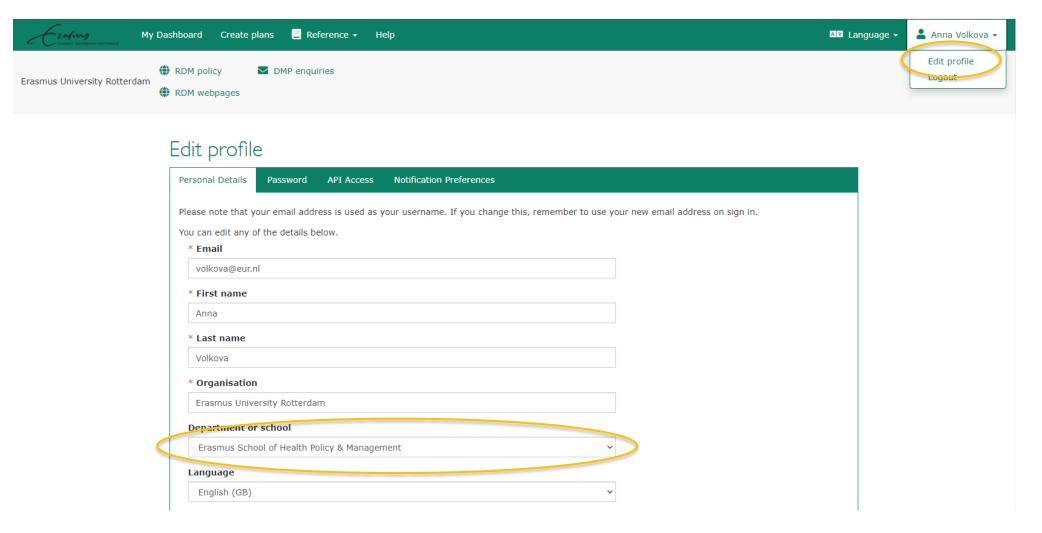


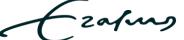
89 Countries

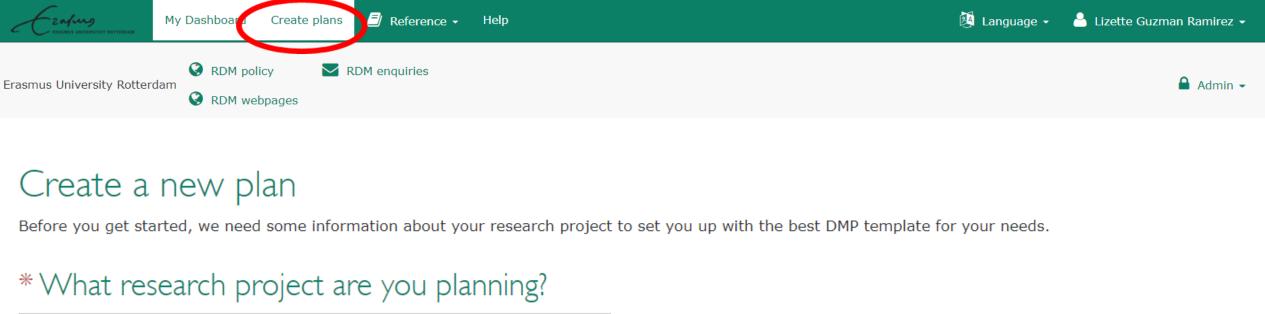
Sign in	Create account
* First N	ame
* Last N	ame
* Email	
Organisa	ation
Begin t	yping to see a filtered list
My organ	isation isn't listed.
* Passw	ord
	password cept the terms and conditions
Create	account

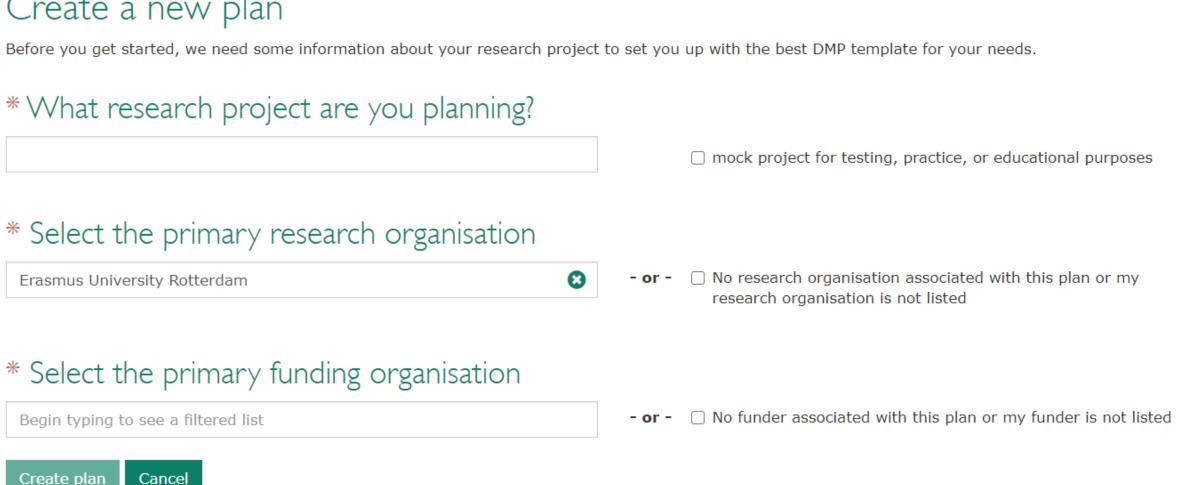


### **DMPOnline: Edit profile**









Support in writing a data management plan is available through the faculty Data Stewards. If your faculty has no Data Steward, support is available through the Research Data Management Specialist of the Erasmus Data Service Centre (EDSC is part of the University Library). Which research support professional is available for you? Data Steward of my own faculty - ERIM / RSM Data Steward of my own faculty - ESE Data Steward of my own faculty - ESSB Data Steward of my own faculty - ESHCC Research Data Management Specialist of the EDSC Save Scientific research must be conducted in line with existing guidelines on good research practices and integrity. Please select which guidelines you are familiar with. The European Code of Conduct for Research Integrity (ALLEA, 2017) The Netherlands Code of Conduct for Research Integrity (VSNU, 2018) Save

Guidance

Comments

**EUR** 

Support at the faculty RSM / ERIM & Faculty ESE

Support faculty ESSB

Guidance

Support faculty ESHCC

Support by the University Library / EDSC

h

Comments

**EUR** 

You can find The European Code of Conduct for Research Integrity (ALLEA, 2017) here.

You can find The Netherlands Code of Conduct for Research Integrity (VSNU, 2018) here.

## If you need anything ...

- If you have questions or need support
  - Let me know!
- If you have written a DMP...
  - Send it to me for review!
- If you used DMPOnline to write your DMP
  - Click 'Request Feedback' and it will be sent to me for review

volkova@eur.nl datasteward@eshpm.eur.nl



### **Questions**



Ezafus,

#### Useful resources

- Erasmus University Rotterdam:
  - Research Data Management pages
  - Legal Support pages
  - MyEUR Privacy pages
  - MyEUR Software pages & Software catalog
  - MyEUR Cybersecurity pages
  - <u>EUR DMPonline</u> (for drafting data management plans)
  - EUR Data Repository (for sharing data)
  - Infographics on Privacy: Why, What and How
  - EUR Privacy & Security app: Apple App Store



#### **Useful resources**

- Research Data Management information:
  - 23 things for RDM
  - Research Data Management the Turing Way
  - Harvard Data Management pages
  - <u>UK Data Service research data management</u>
  - Go FAIR FAIR principles
  - National Coordination Point Research Data Management



# Making your data FAIR(er)

#### Findable

 Assign persistent IDs, provide rich metadata, register in a searchable resource

#### Accessible

 Retrievable by their ID using a standard protocol, metadata remain accessible even if data aren't

#### Interoperable

 Use formal broadly applicable languages, use standard vocabularies, qualified references

#### Reusable

 Rich, accurate metadata, clear licenses, provenance, use of community standards



### **Enters RDM...**

Research data management (RDM) is a term that describes the

- organization,
- storage,
- preservation, and
- sharing of data collected and used in a research project.

