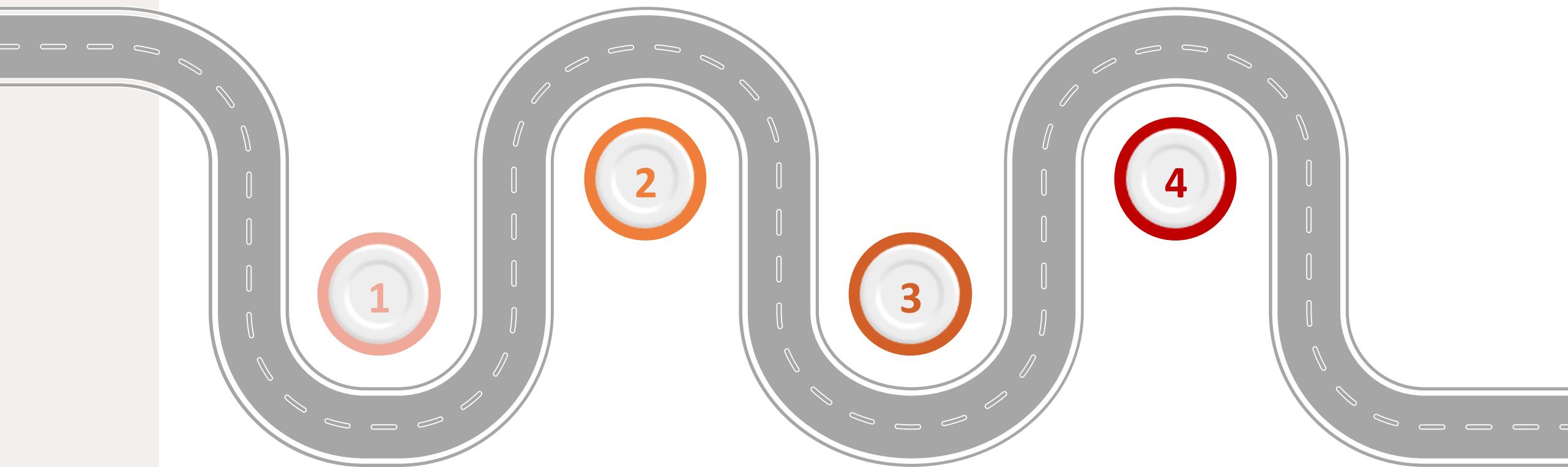


Research Data Management Onboarding



Your RDM Roadmap



Explore Funding & Data



Funding

VU IXA-GO

- Identify and secure **external funding**
- Provides **workshops and trainings**
- Personalised **funding scans** from VU grants office
- **CV scan** to increase competitiveness in securing related grants

Research Connect

- Broad platform to find **external funding**
- Offers complementary services to **customise funding search**

Centre for International Cooperation (CIS)

- **Specialised grant advising** for projects with an international focus in:
 - Food and Nutrition and Sustainable Land Management
 - ICT for Development
 - Conflict Resolution and Mediation and Governance for Society (incl. Security & Rule of Law)

Pre-existing Datasets



VU Resources

[LibSearch](#)
[VU Research Portal](#)
[PURE](#)
[Other data sources](#)



Repositories

[DataVerseNL](#)
[DANS: Data Stations](#)
[4TU.Research Data](#)
[B2SHARE](#)
[Zenodo](#)
[Re3data](#)



Journals

[Data Journals](#)
Check supplements
and references of
research articles



Search Engine

[DataCite](#)
[Google DataSet Search](#)

2



Investigate your
legal
responsibilities

Knowledge Security

What is it

The undesirable transfer of **sensitive information** impacting **Netherlands' national security**.

Covert influencing of education and research **by dangerous state actors**

Collaboration with individuals and institutions **from countries** that **do not respect fundamental rights**

How does it impact you*

Primary responsible researcher **answers 6 framework** questions

One **yes = risk assessment needed** (contact faculty knowledge security officer for assistance)

If needed, they **contact the VU knowledge security advisory group** for advice

*If you're conducting international collaborations (formal, informal, hiring, visitors)

See [Knowledge Security](#) for more information

VU Knowledge Security Framework

Legal framework – what is not permitted, or under conditions

1. Does the collaboration concern a sanctioned person, entity or body? Or is the combination of the technology and country sanctioned? - See [EU sanctions list](#).
2. Is the Dual Use regime applicable to the research? - Contact the [KS advisory team](#) for advice.

Risk management – what else should be considered

3. Is, or has the partner been affiliated to a military organization outside the EU? - *collaboration with military organisations outside EU is undesirable.*
4. Does the collaboration involve sensitive research?
5. Do ethical dilemmas play a role? - *check if the partner is based in county which scores 0.4 or less on the Academic Freedom Index.*
6. Does unilateral external funding occur in the cooperation? - *Check AFI*

No limitations – cooperation can take place

7. There are no obstacles for the collaboration from a knowledge security perspective

Legal Requirements

Valorisation & Impact

Contact IXA

- Helps with intellectual property by setting up [patents and copyrights](#)
- Assists in setting up [contracts](#) with third parties (e.g. consortium agreements)

Medical Research

If conducting medical research, determine:

- How your research is defined by [Dutch law](#)
- Whether any additional [medical research laws](#) apply

3

Register & Prepare



Pre-register

What?

Practice of documenting your **research plans** at the beginning of the study

- **Time-stamped, read-only, publicly available**

Why?

Improves **findability, transparency, reproducibility, and collaboration**

When?

After **you have a plan** for your study, but before **you get started**

How?

- First create an [ORCID](#) (a unique, persistant identifier for you as a researcher)
- [Create a pre-registration](#) on OSF

Data Classification

Factors that will impact the risk classification for your data:



3

Personal Data Detour



Read up on policies

Consult the [Privacy Five Step Plan](#)

Read up on the GDPR, beginning with this [GDPR primer](#)



Assess your data's privacy risk level

Use the [Data Classification Tool](#) and follow the necessary measures for risk level



Learn about de-identification

Do your best to de-identify your participants' data.

(See this helpful [guide](#) from University of Groningen)



Write your ICF appropriately

Use the [ICF generator tool](#) to create a faculty approved form

Or check your [faculty ethics page](#) for ICF templates

Forms you may need

Assessments



[Data Protection Impact Assessment \(DPIA\)](#)

Pre-DPIA

BIV

Legal Requirements



[Privacy Statement](#)

[GDPR registration](#)

Informed Consent Form

External Agreements



Joint Controllers Agreement
Data Processing Agreement (DPA)

Data Transfer Agreement (DTA)

Confidentiality agreement



Determine
storage &
technical
solutions

Storage Solutions



Yoda

Good for:
large volumes of data
that don't need
frequent
processing/analysing

Recommended for:
archiving



ResearchDrive

Good for:
large volumes of data
that need regular
access

Easy collaboration

Recommended for:
active stage



SciStor

Good for:
very large volumes of
data that need regular
access

Recommended for:
high-performance
computing



Teams/Sharepoint

Good for:
collaboration

Recommended if:
no other option is
possible and data is
not high sensitivity

Use [Storage Finder](#) to choose an appropriate solution

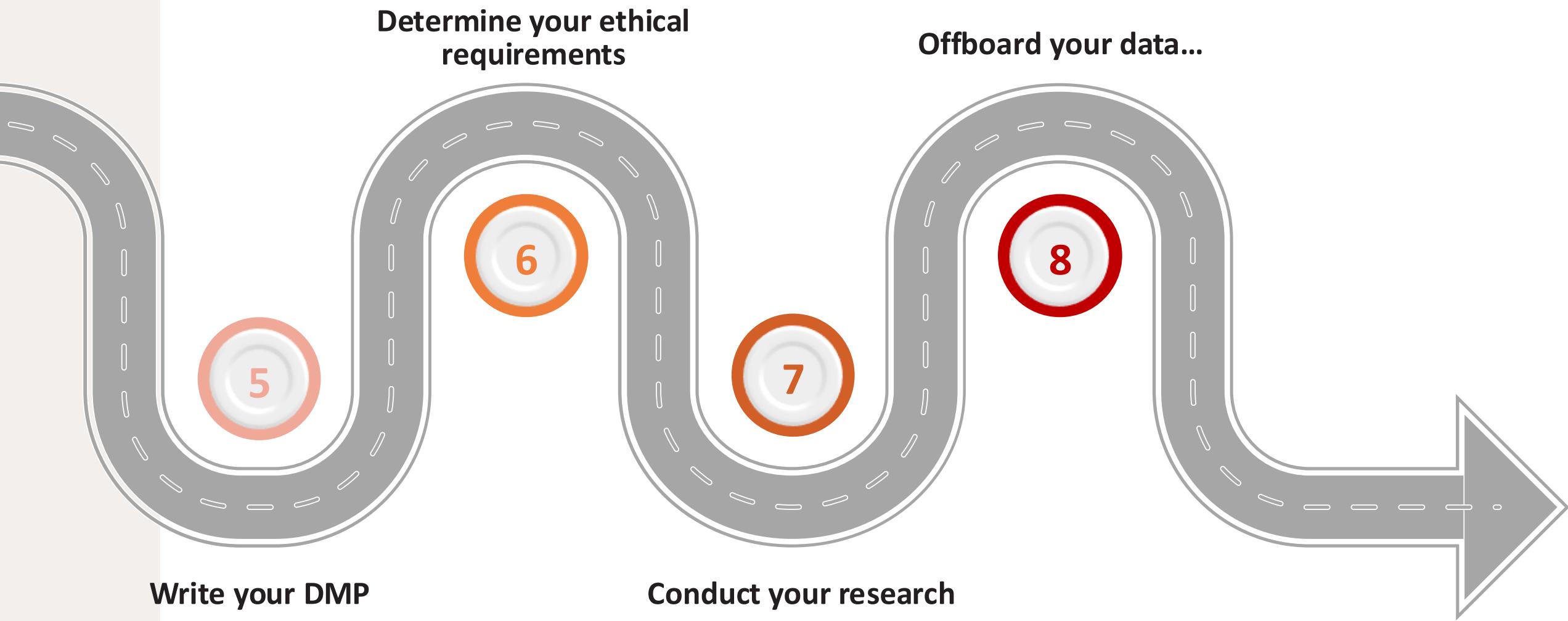
ITvO: IT for Research

Services:

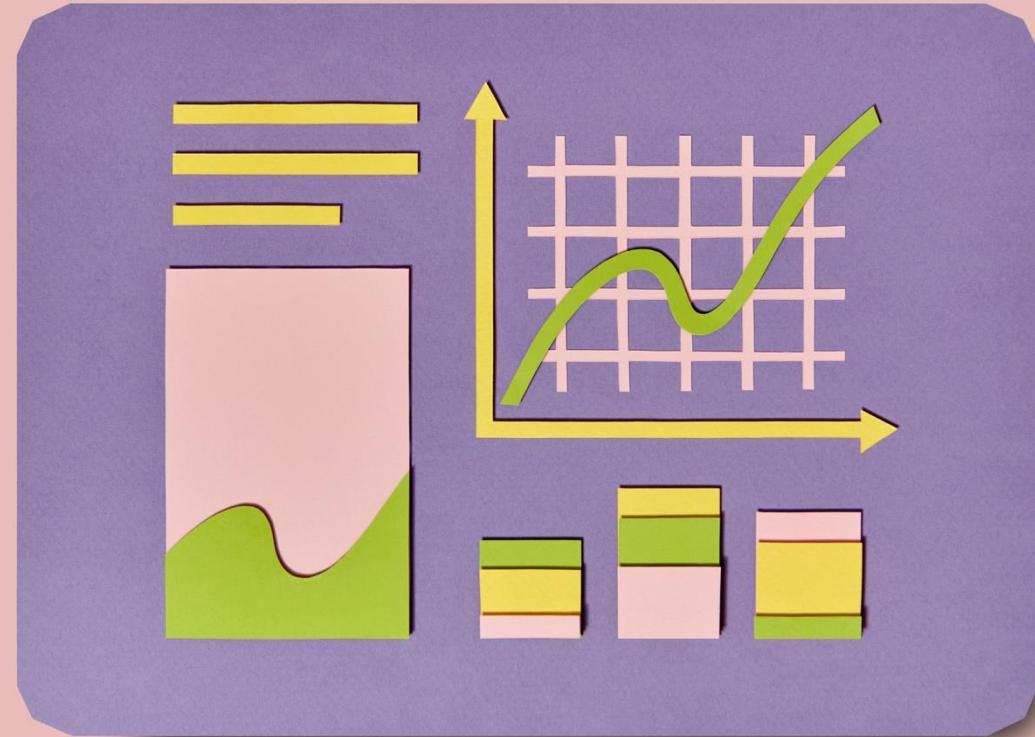
- SciStor: scientific storage service of VU
- SciCloud: flexible platform for virtual machine management
- Ada HPC: high performance computing cluster of VU
- Consultancy & technical support
- Advice on data management, security and analysis workflows

When to reach out (examples):

- Your research requires specific IT infrastructure
- You need to set up a public web application
- Your laptop is too slow to run your analysis
- Your lab instrument requires specialised software
- You're not sure if you'll need technical support



Write a DMP



Write a DMP

What?

A document outlining how **research data will be handled** throughout the **research life cycle**

Why?

Identify challenges early, receive support while it's still possible, implement **FAIR** practices from the start, **reduce administrative overload** at the end of the project

When?

Begin after **you have a plan** for your study, but before **you get started**

- This is a **living document**: update with any changes as you go

How?

Use [**DMPonline**](#) to write your DMP

- tip: use the VU DMP Template as it's approved by most grant providers and automatically registers you with the GDPR if necessary



Determine your
ethical
requirements

Ethical Requirements

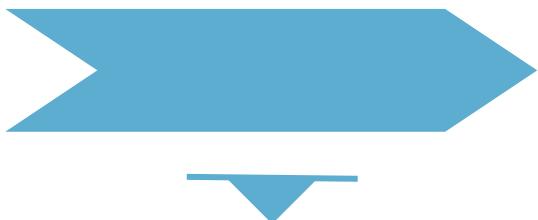
1



Investigate

Faculty requirements
NETHICS
WMO/METc

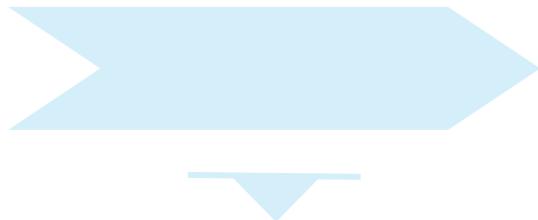
2



Prepare

The necessary documents as required by your faculty
e.g. DMP, GDPR registration, Privacy Statement

3



Submit

The required documents and ethics form as per your faculty procedure
e.g. Informed Consent Form, Participant Information Sheet

WMO/METc

Research falls under the WMO if the following two conditions are met:

- it concerns medical-scientific research; and
- people are subjected to procedures or are required to follow rules of behaviour

Some research may require a non-WMO declaration:

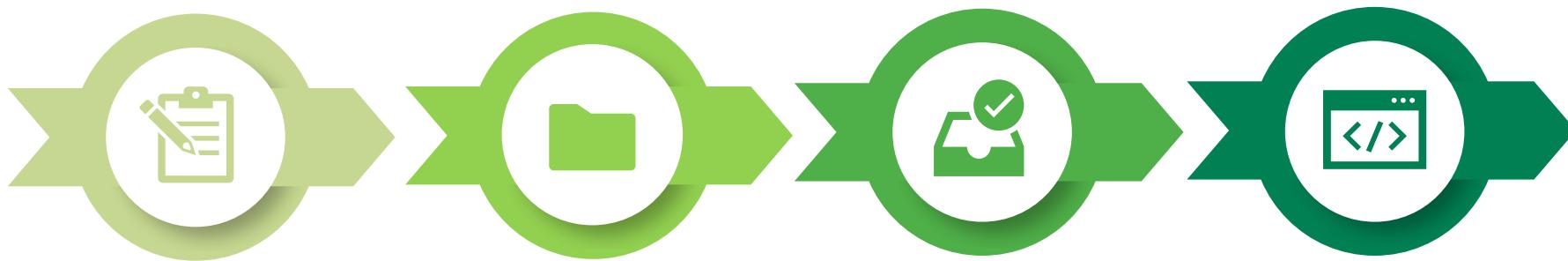
- You need a non-WMO statement/approval from a recognised METC, for example for publication.
- Non-WMO statement is not ethical approval, check with your faculty ethics committee if you need to undergo a full ethical review

Conduct your research



Data Collection

Good practices to maintain throughout data collection:



Documentation

Keep up to date [documentation](#) on your research
e.g. *codebook, e-lab journal, README file*

Organisation

Maintain a clear file structure
See SURF's guideline for an example of good folder structuring

Data Quality

Follow field specific [protocol](#) and [clean](#) your data

Metadata

Provide [metadata](#) alongside your research. Be sure to follow domain standards if applicable

Collaboration



Roles & Responsibilities

Define the roles & responsibilities **early and clearly**.

Which data does the recipient have access to? What can they do with the data?



Agreements

Based on the role of the recipient, draw up the necessary agreements:

See the '**Forms you may need**' slide to find the kind of external agreements you may need.



Security measures

Determine the necessary security measures given the **privacy risk** of the data.

Does the recipient need MFA? Full-disk encryption?



Offboard
your data

Archiving & Publishing

Data is properly archived when



**Persistent identifier – DOI
(Findable, linkable)**



**Reuse License – Terms and
Conditions**



**Locked and unable to be modified –
verification**

Data can be published as

- **Closed**
- **Restricted**
- **Open**



Provide with your research data



Standard metadata

See [YODA metadata](#)



Contact information for data access requests

(and long term, who should be contacted for data deletion after 10 years)



Is the project continuing at the VU, but you are leaving?

Then provide:



Project location data

i.e., where to find your data



Access to your data repositories

Deleting your data

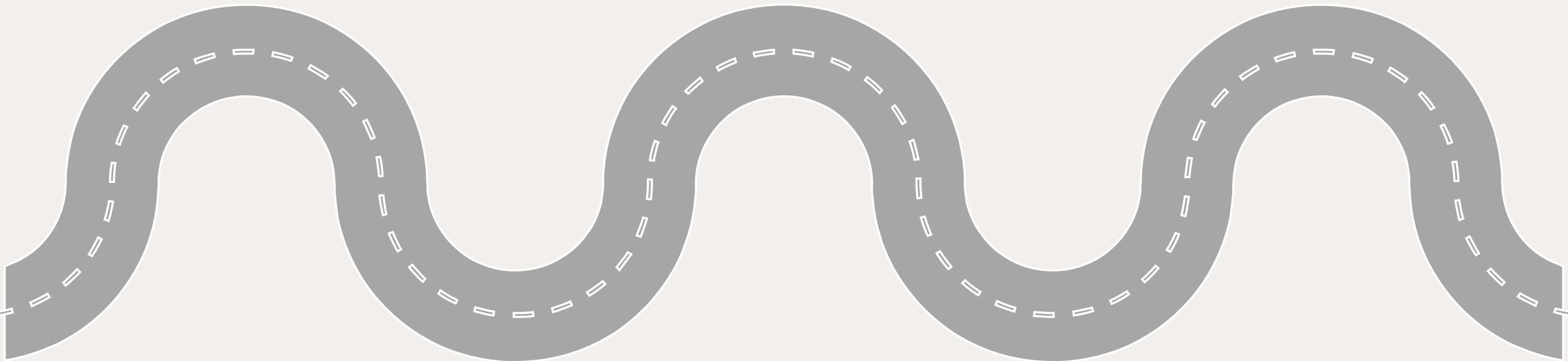
Delete your data from all non-archival repositories and software, including:

- Castor
- ResearchDrive
- Qualtrics
- Your local computer
- Any other software/tool/repository which the data can be stored and is not where you will be archiving

For most of these tools, contact rdm@vu.nl to be put in contact with the functional manager for deletion

Prior to deletion ensure all data you plan to delete has been exported and archived to the proper repository

Roadmap Checklist



Have you...

- Explored your funding & data?
- Written your DMP?
- Investigated your legal responsibilities?
- Determined your ethical requirements?
- Registered and prepared for your research?
- Conducted your research?
- Determined your technical & storage solutions?
- Offboarded your data?

General RDM support:
rdm@vu.nl

Faculty specific data stewards:
[Contact your data steward](#)

Faculty specific privacy champions:
[Privacy champions](#)