

MedUni Vienna Data Clearing House

Request for disclosure of personal data or previously anonymised data to external recipients

1. Applicant at the Medical University of Vienna

Organisational unit

Applicant			
Family name	First name	Title	
Telephone/Pager	Email		
Contact person ¹ (if not the app	licant)		
Family name	First name	Title	
Telephone/Pager	Email		
2. Data recipient			
Name of the company / partn	er organisation		
Institute, department, clinic, v	vorking group		
Is the recipient of the data loc	ated within the EU?	yes	no
Contact person ¹			
Family name	First name	Title	

Telephone/Pager

Email

¹ An individual who can provide responses to organisational or technical questions.



What is the relationship between the applicant and the data recipient? Are the applicant or employees of the applicant also employed by the data recipient?

3. Project

Title

Description: A description of the project is to be attached to the request (refer to Point 6, Annexes).

Organisational units at the MedUni Vienna which are involved

4. Data disclosure

Do (actual / draft) written or oral agreements relating to the data disclosure already exist between the cooperation partners?

If yes: which agreements? (Please provide relevant details if the agreement is already stored in the contracts database of MedUni Vienna)

yes no

yes

no

Is there an informed consent for this project? (If yes, please attach it to the annex)



The data disclosure will be

once only	Is the intention to disclose only once as a single data packet or
	is it expected that data will be disclosed multiple times at different
m ultiple disclosure	points in time (e.g. When new source data are available, follow-ups,
	etc.)?

yes no

Have data already been disclosed in the course of this research activity (or this research project)

If yes:

Date of data disclosure

Is there an agreement?

Was the data disclosure authorised by the data clearing house?

If yes, DC-number / Date

Which data were disclosed?

Reason for disclosure (cooperation, spin-off, multi-centre trial, etc.)

Have these or associated data of the same patient already been yes no published or disclosed to other recipients in the past?

5. Data

Description: A description of the data is to be attached to the request (refer to Point 6, Annexes).

yes no

Is it necessary to be able to re-identify the patients later

What is the purpose of this?



6. Annexes

Attached

Project description (see Point 3)

Data description (see Point 5), including the following information

- Source (OU, clinic, device)
- Storage location
- Data types (images, electronic documents, videos, etc.)
- Data formats (DICOM, JPG, PDF, Excel, proprietary, etc.)
- Data volume (GB, TB, number of investigations, number of patients, etc.)
- Will the data be transferred to one or more recipients? Which data is transmitted to which recipients?
- Do the data contain personal details (name, occupation, date of birth, medical history, social insurance number, etc.)
- Have the data been anonymised / pseudonymised?
- Are there requirements in terms of the pseudonymisation of the data?
- During the data documentation of a study / trial, are the data entered into a system other than a MedUni Vienna system?
- A description of the data storage and processing at the MedUni Vienna and on the part of the data recipient, including a diagram of this.
- Which data are extracted, linked, etc.?
- How is it intended to disclose the data to the recipient?
- Do technical requirements exist on the part of the recipient?

Contracts, draft contracts and, in the case of other agreements, records or correspondence which state the facts should be attached

Data records covering at least 10 cases (investigations, patients, etc.)

If applicable: Study protokoll, clinical trial plan, trial-related record

If applicable: Applicant to the ethics commissions and the voting (current version)

If applicable: Text of the patient consent form (current version)





Signature of applicant



OU Head: Name in block capitals

Signature of OU Head