Data Clearing Office at the Medical University of Innsbruck

Application form concerning the transfer of personal data to external recipients

Organizational Unit		
Applicant		
Name	First Name	Academic Title
Phone	Email	
Respondent¹ (if different	from Applicant)	
Name	First Name	Academic Title
Phone	Email	
2. Recipient		
Name		Address
Respondent ¹		
Name	First Name	Academic Title
Phone	Email	
		elated to each other in this context? Please spec also employed by the data Recipient?
	projects or the applicant t	mso employed by the data needplent.

page 1 of 5

¹ A person who is able to provide information on procedural and/or technical matters.

3. Project			
Title			
Description The project description must be attached	to the application (see also item 6.	Attachments).	
Participating organizational units			
Project duration Start	Termination		
Is there an Ethics Committee Vote (IRB)		YES 🗆	NO □
If "Yes", please provide the reference nu	Imper of the Ethics Committee decis	ion	
4. Data Sharing			
Are there already any contracts, draft or regarding data sharing between the co	•	ents	
If yes, please specify below:			
Has the data already been transferred			
part of this research activity or project?	?	YES 🗌	NO 🗌
If "Yes":			
Date of the Data Sharing			

page 2 of 5

Does the aforementioned personal data have been	Is ther	e an agreement in place concerning this very project?	YES 🗆	NO □
Please specify the ratio and the legal basis of Data Sharing (Cooperation, Spin-Off, multicentric Study Does these or any related personal data already have been shared with other recipients in the past? Does the aforementioned personal data have been published in at least pseudonymized form? YES NO NO NO NO NO Storage and Storage location Data types (images, electronic documents, videos, etc) Data formats (DICOM, JPG, PDF, Excel, proprietary, non-proprietary) Data volume (GB, TB, number of examinations, number of patients included,) Has the Data been anonymised/pseudonymised beforehand? Are there requirements regarding the pseudonymisation of the data arising from contracts or from the studies themselves (keyword: re-identification of patients)? Is the data entered into a system outside the jurisdiction/sphere of influence of the Medical University of Innsbruck regarding the and correlating to the study's data documentation? Please specify and illustrate accordingly the data processing at the Medical University of Innsbruck as well as at the Recipients end in a schematic way How (technically) will the data be shared with the recipient? What TOMS (technical and organisational measures) are present on the recipient's side	If "Yes", p	please specify below:		
Please specify the ratio and the legal basis of Data Sharing (Cooperation, Spin-Off, multicentric Study Does these or any related personal data already have been shared with other recipients in the past? Does the aforementioned personal data have been published in at least pseudonymized form? YES NO NO NO NO NO Storage and Storage location Data types (images, electronic documents, videos, etc) Data formats (DICOM, JPG, PDF, Excel, proprietary, non-proprietary) Data volume (GB, TB, number of examinations, number of patients included,) Has the Data been anonymised/pseudonymised beforehand? Are there requirements regarding the pseudonymisation of the data arising from contracts or from the studies themselves (keyword: re-identification of patients)? Is the data entered into a system outside the jurisdiction/sphere of influence of the Medical University of Innsbruck regarding the pseudonymisation of the study's data documentation? Please specify and illustrate accordingly the data processing at the Medical University of Innsbruck as well as at the Recipients end in a schematic way How (technically) will the data be shared with the recipient? What TOMS (technical and organisational measures) are present on the recipient's side				
Does these or any related personal data already have been shared with other recipients in the past? Does the aforementioned personal data have been published in at least pseudonymized form? YES NO	Please sp	ecify the Data that has been shared.		
been shared with other recipients in the past? Does the aforementioned personal data have been published in at least pseudonymized form? YES NO	Please sp	ecify the ratio and the legal basis of Data Sharing (Cooperation, Sp	in-Off, multicentr	ic Study,)
been shared with other recipients in the past? Does the aforementioned personal data have been published in at least pseudonymized form? YES NO				
Does the aforementioned personal data have been published in at least pseudonymized form? YES NO	Does t	hese or any related personal data already have		
 published in at least pseudonymized form? S. Data The description of the data must be attached to the application and must include at least the following criteria and information: Source of the Data (Organizational Unit, Clinic TK, device) Storage and Storage location Data types (images, electronic documents, videos, etc) Data formats (DICOM, JPG, PDF, Excel, proprietary, non-proprietary) Data volume (GB, TB, number of examinations, number of patients included,) Has the Data been anonymised/pseudonymised beforehand? Are there requirements regarding the pseudonymisation of the data arising from contracts or from the studies themselves (keyword: re-identification of patients)? Is the data entered into a system outside the jurisdiction/sphere of influence of the Medical University of Innsbruck regarding the and correlating to the study's data documentation? Please specify and illustrate accordingly the data processing at the Medical University of Innsbruck as well as at the Recipients end in a schematic way How (technically) will the data be shared with the recipient? What TOMs (technical and organisational measures) are present on the recipient's side 	been s	hared with other recipients in the past?	YES 🗀	NO 🗀
 5. Data The description of the data must be attached to the application and must include at least the following criteria and information: Source of the Data (Organizational Unit, Clinic TK, device) Storage and Storage location Data types (images, electronic documents, videos, etc) Data formats (DICOM, JPG, PDF, Excel, proprietary, non-proprietary) Data volume (GB, TB, number of examinations, number of patients included,) Has the Data been anonymised/pseudonymised beforehand? Are there requirements regarding the pseudonymisation of the data arising from contracts or from the studies themselves (keyword: re-identification of patients)? Is the data entered into a system outside the jurisdiction/sphere of influence of the Medical University of Innsbruck regarding the and correlating to the study's data documentation? Please specify and illustrate accordingly the data processing at the Medical University of Innsbruck as well as at the Recipients end in a schematic way How (technically) will the data be shared with the recipient? What TOMs (technical and organisational measures) are present on the recipient's side 				
 The description of the data must be attached to the application and must include at least the following criteria and information: Source of the Data (Organizational Unit, Clinic TK, device) Storage and Storage location Data types (images, electronic documents, videos, etc) Data formats (DICOM, JPG, PDF, Excel, proprietary, non-proprietary) Data volume (GB, TB, number of examinations, number of patients included,) Has the Data been anonymised/pseudonymised beforehand? Are there requirements regarding the pseudonymisation of the data arising from contracts or from the studies themselves (keyword: re-identification of patients)? Is the data entered into a system outside the jurisdiction/sphere of influence of the Medical University of Innsbruck regarding the and correlating to the study's data documentation? Please specify and illustrate accordingly the data processing at the Medical University of Innsbruck as well as at the Recipients end in a schematic way How (technically) will the data be shared with the recipient? What TOMs (technical and organisational measures) are present on the recipient's side 	publish	ned in at least pseudonymized form?	YES 🗀	NO 🗀
	The defollowing	scription of the data must be attached to the application and must ng criteria and information: Source of the Data (Organizational Unit, Clinic TK, device) Storage and Storage location Data types (images, electronic documents, videos, etc) Data formats (DICOM, JPG, PDF, Excel, proprietary, non-proprietar Data volume (GB, TB, number of examinations, number of patients Has the Data been anonymised/pseudonymised beforehand? Are there requirements regarding the pseudonymisation of the contracts or from the studies themselves (keyword: re-identificated is the data entered into a system outside the jurisdiction/sphere Medical University of Innsbruck regarding the and correlating to documentation? Please specify and illustrate accordingly the data processing at the Innsbruck as well as at the Recipients end in a schematic way How (technically) will the data be shared with the recipient?	y) included,) lata arising from tion of patients)? of influence of ti the study's data ne Medical Unive	he rsity of
		to ensure the security of processing.	_	
Is a later re-identification required? YES L NO	Is a lat	er re-identification required?	YES 🗀	NO L
If "Yes": for what purpose?	If "Yes": f	or what purpose?		

page 3 of 5 version 1.4_en

YES □	NO 🗆
_	YES □

6. Attachments

Compulsory:

- Project description
- Description of data
- Contracts (finalised/drafted) or other agreements; if the latter please provide the protocols or correspondences that reproduce the circumstances
- (Test)-datasets of at least 10 cases (examinations, patients, etc.)
- Consent Form if present
- If applicable and present:
- Study Plan, Study Protocol
- Ethics Committee application and/or vote (reference number)

Date	J	Signature Applicant
		Printed name
Date		Signature Head of Organizational Unit
		Printed name