mci ethics assessment.

TO THE MCI/ETHICS COMMITTEE

Please complete the following pages (digital only, no handwriting allowed) and submit to your department. mci ethics assessment part 1.

Title of the research project:

First name of the student:

Second name of the student:

Student ID:

Department/Study program:

Email: @mci4me.at

Name of the supervisor:

What method do you use in your study?	Empirical analysis	
	Pure literature work	
	If your study is based	
	solely on literature work,	
	questions 3, 4 and 5 in the	
	second part of the assess-	
	ment are not relevant to	
	you.	
	Desktop research /	
	Secondary data analysis	
	If your study is based ex-	
	clusively on a secondary	
	data analysis, questions 4	
	and 5 in the second part of	
	the assessment are not rel-	
	evant to you.	



Please indicate which of the following apply to	o the research project/study in question	Check all boxes that apply
Was the study previously denied ethical cleara	nce by another review board?	
Has the study already been granted ethical clear If yes, a copy of this clearance must be attache		
Could there be any potential conflicts of intere the research, the public, your own interests)?	st (between clients, those affected by	
Will secondary data analysis be conducted as p data will be processed?	part of the study, during which sensitive	
Will the study intentionally mislead participants in any way? If yes, provide details of this in question 5 and explain how this misleading is handled.		
Could the study pose a risk of exposing particip stress situations or discomfort (either as a rest publication of the results)? <i>If yes, please provid</i> <i>what instructions/advice you will give participa</i> <i>who to ask for help in such cases</i>).	ult of participating in the study or by le details on the next page and tell us	
Will the study require access to personal and/o genetic, biological, socio-administrative inform ual orientation?		
Does the study indirectly or directly affect any of the following groups?	• Children and teenagers (under 18 years)	
	Persons with cognitive or com- munication impairments	
	 vulnerable groups of people (e.g. children and adults with physical or cognitive impairments, ethnic minorities, persons in asym- metrical or dependent relation- ships, pregnant women, mar- ginalized persons e.g. home- less persons, LBTIQ* people, refugees, etc.). 	
	Patients	
	 Persons who are particularly exposed and/or identifiable due to their position. 	



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part 2.

1) Aim of the study including academic reasoning - motivation, purpose, objectives, etc.:

2) Preliminary Research Question(s):



3) Brief description of the methods planned for data collection, storage, analysis and -interpretation:

(e.g. Which research methods will be used? How and where will the data be collected? Which tools will be used? Where will the data be stored? When will the data be deleted? What precautions will be taken to ensure results remain confidential and anonymous?

In the case of secondary data analysis: How is the data provided and transmitted? How is the selection of the data basis carried out? How is the processed data secured and archived? How is the analysis documented?)

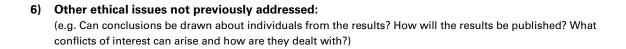
4) Selection of participants/sampling – Age, gender, etc., exclusion/inclusion criteria, numbers, reasoning for respective numbers of participants:
(a.g. Who should participate and why? What is the target sample size?)

(e.g. Who should participate and why? What is the target sample size?)



5) **Dealing with participants**:

(e.g. How is contact established? How are participants informed about the research project? How will informed consent be obtained? Will participants have the option NOT to participate or to withdraw their participation during the research process? How will potential risks to participants or the researcher related to the research be handled? How will feedback of results be provided to participants? What precautions are taken to deal with risks to participants? To what extent are people with communication impairments considered in the design of research? What precautions are taken to avoid drawing conclusions about individuals?





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acknowledgement.

I hereby confirm that I will (where applicable):

- Provide participants with an information sheet (or in the case of an online study, a web page) to explain the basic procedures (see "Example information sheet"),
- Inform participants that participation is not mandatory,
- In case of an observational study, obtain the participants' permission to observe them,
- Obtain permission from the participants to observe them,
- Inform the participants that they can interrupt the study at any time and for any reason without suffering any disadvantage,
- If a questionnaire is used, offer participants the option of omitting any questions they do not wish to answer,
- Assure the participants that their data will be treated confidentially and will not be personally identifiable in the event of publication,
- Inform participants that any recordings, such as audio or video recordings and photographs, cannot be traced back to individuals unless written consent is given in advance, and
- If requested, provide a debriefing (i.e., an explanation of the outcomes of the study) after the study is completed.

Signature:

Date:

Student

.....

Signature:

Supervisor

Date:

