Research proposal for ethical evaluation by the FGGA Ethics Committee

In accordance with the <u>VSNU Code of Conduct for Scientific Practice</u> (2018), Leiden University ensures that researchers can address their questions regarding research integrity to an ethics committee and have their research proposals ethically reviewed. The FGGA Ethics Committee evaluates <u>all new</u> research projects that directly involve human subjects and are designed by FGGA researchers or where the main applicant is a researcher of the FGGA, using the most recent, nationally recognized version of the <u>DSW Code of Ethics Social and Behavioral Science</u> or other relevant guidelines for ethical research conduct.

Before using this form and for further guidelines on the procedure for submitting an application for ethical review, please consult the *FGGA Ethics Committee Guidelines*

Name and contact details
Please include email address and telephone number (office)
Name of project and group
Name your supervisors/co-researchers
Name principal investigator/co-researchers/supervisor (if relevant)
Title of research project
Title of Teesal on project
Project duration
Projected starting date and duration
r rejected claraning date and alleran
Funding body/bodies
If applicable: 1st money stream (Leiden University), 2nd money stream (NWO/KNAW/EU), 3rd
money stream (all other)
Grant number
If applicable. A grant number provides unique identification for the grant.

	tner organizations oplicable. These may be research partners that use your data, or that you use data from.	
	If ye	a icate whether your research directly involves human subjects es, please answer all questions in the template format below o, please answer questions 1 through 3 and 8 in the template below
]	If ap	dical-Ethical Evaluation Committee oplicable. Is or will this research proposal be evaluated by a Medical-Ethical Evaluation mmittee?
1.		search topic and research question Please describe in max. 100 words what your research topic and preliminary research question(s)/objective(s) are.
2.		mmary of Proposed methods and brief work plan Please provide a brief statement of your methodology and clarify your theoretical and methodological premises

b.	Are your prepared to make your theoretical premises and main questions open to the Ethics Committee (available for internal review)?	Yes No
C.	Are your prepared to make your theoretical premises and main questions open to the public after publication of your results?	Yes No
d.	What methods do you intend to use to access data to answer you (e.g. survey, qualitative interviews, observation, focus group interother)? i. State the estimated number of participants of your research response rate, if applicable) ii. State the estimated percentage of non-response of your research response of your research response rate.	views, life histories,
	iii. What is your recruitment strategy for the target population iv. How do you plan to get access to your research target population	
e.	Do you have assistance (e.g. research partner organizations) in access to your research target population? i. If yes, from what kinds of organizations are you having assistance from to gain access to your research target population?	Yes No

3. lmլ	partiality and independence ¹				
a.	What procedures do you have to ensure your impartiality and independence in the way				
	you set up and conduct your research?				
b. Please designate a colleague from the FGGA (at associate or full professor leve					
	not involved in the research) to serve as your advisor in case of o	uestions of impartiality			
	and scientific independence in relation to your funder, in the cour	•			
	specifically in relation to your conclusions	•			
	·				
		Yes			
C.	Are you aware of any potential issues that your funding body				
	could use to influence your research?	No			
	i. If yes, please specify what actions would you				
	undertake to ensure your independence?				

4. Location, setting and target population

Describe in max. 200 words:

- a. In which country/countries your research will take place and why there;
- b. What the precise setting is in which you intend to conduct the research (i.e. workplace, shelters, a clinic, people's homes, streets); and
- c. If known, please also specify further which categories of people are directly involved in your research

process as well as the dissemination of research findings.

¹ Impartiality refers to ensuring that the research or experiment be conducted in a fair, honest, impartial, and transparent manner after full disclosure is made by those associated with the research or experiment of each aspect of their interest in the research, and any conflict of interest that may exist.

Independence refers to freedom of research and professional integrity that is maintained throughout the research

Application form for ethical review of research proposals, version December 2018		
5. Confidentiality, informed consent and data protection		
Discuss in max. 200 words: a. Do you have a written consent form or do you intend oral consent participants?	of your research	
 Please give an example of your personalized Informed Congiven in the document <u>FGGA Ethics Committee Guidelines</u> IC form, so include it in an appendix to the application for example send the IC form as a separate file.). Please append the	
b. Do you intend to obtain consent in case of semi-structured	Yes	
interviews?	No	
c. Is it possible to ensure the anonymity ² of your respondents if	Yes	
you publish your analysis?	No	
d. Do you intend to keep the information your research	Yes	
participants will provide for your project confidential in your resulting publications/reports?	No	

 $^{^2}$ Anonymity implies that no one can trace back the original respondent; confidentiality means that it is still possible to trace back the original respondent.

6. Vulnerability and protection of research participants/precautionary measures to ensure your own and your participants' safety Describe in max. 200 words: a. What are your precautionary measures to ensure your own safety? Yes b. Are the people you intend to collect data from or on part of a group that can be considered vulnerable, in their context? No c. Will your participants potentially be put at risk as a result of Yes participating in your research? No d. Are there any concerns for the safety of persons whom you Yes may contact, due to a.o. the political or cultural context of your Nο research? If yes to questions c or d, how do you intend to mitigate those risks; how will you ensure that your research will not cause harm to the people you involve?

7. Official permission in host country, and embeddedness (Only in case you plan to carry out research abroad)

Describe in max. 150 words:

a. Have you familiarized yourself with the official requirements around research in the country/on the site where you want to conduct the research? (E.g., do you have to apply for specific visa, and/or do you have to be embedded in/affiliated with an organization?)

[b.	-	have an organization carrying out research on your , are you familiar with the requirements for their safety? If yes, please briefly list the requirements as well as the risks involved in carrying out the research on your behalf.	Yes No
L				
	Reminder: Please check whether or not you are allowed to use personal data. In accordance with European General Data Protection Regulation, researchers should indicate which of th acknowledged foundations justifies their use of personal data. For help or information, may contact the data protection officer via privacy@bb.leidenuniv.nl . See https://www.stoff.upiversiteitleiden.pl/ict/privacy.and.deta.protection/general.deta.			dicate which of the p or information, you
	See			

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Click the button to save and submit your form:

You can send all your questions regarding research integrity in general to ethicscommittee@fgga.leidenuniv.nl.

Reminder:

Have you written a data management plan? Please contact the Centre for Digital Scholarship at the University Library if you need help: datamanagement@library.leidenuniv.nl