



Institutional Research Ethics Checklist

This checklist must be completed and submitted through email (oric@cmhlahore.edu.pk) along with the research synopsis to the ORIC-CMH LMC & IOD.				
Undergraduate Student □	For office use only			
Postgraduate Resident:				
Faculty:	ORIC No:			
Other:				
Other.	ERC No:			
Department/School/Other Unit:				
Program of study (if applicable):				
Name of Principal investigator:				
Name of 1 fincipal investigator.				
Name of Supervisor:				
Corresponding E-mail:				
Name/s of all co-investigators:				
Project Title:				
Start date:				
End date (cannot be retrospective):				
Number of participants (if applicable):				
Funding Source (if any):				
1. Participant Information and Consent				
• Are you gathering data from people? If yes:		YES \square NO \square		
		YES □ NO □		
(a) Have you attached a <u>participant</u> information to their involvement in your research and mainta data?				
(b) Have you attached a consent form?		YES \square NO \square		





2. Data Source and Protection		
•	Are you gathering data from secondary sources such as websites, archive material, and research datasets?	YES □ NO □
•	Have you read the data protection guidelines? (https://www.isaca.org/resources/news-and-trends/industry-news/2024/achieving-ethical-protection-of-data-privacy#f6)	YES □ NO □
. ,	Have you considered and addressed data protection issues relating to storing disposing of data?	YES □ NO □
	Is this an auditable form (can you trace the use of the data from collection to posal)?	YES □ NO □
•	Have you planned to share data with third parties, have you obtained appropriate consent and safeguards?	YES □ NO □
3. Vulnerable Participants		
•	Are you gathering data from people who are considered vulnerable participants (e.g., children, pregnant women, prisoners)?	YES □ NO □
If yes:		
•	Have you read the guidelines for proceeding with research and obtaining consent from vulnerable participants (e.g., children, people with learning disabilities, your students)? (http://nbcpakistan.org.pk/assets/national-guidelines-for-collection%2c-usage%2c-storage-and-export-of-human-biological-materials-20202.pdf)	YES □ NO □
4. Sensitive Data and Study Protocol		
•	Have you read the Helsinki – Ethical Principles for Medical Research Involving Human Participants (https://www.wma.net/policies-post/wma-declaration-of-helsinki/)	YES □ NO □
•	Will the study involve using participants' images or sensitive data (e.g., personal details stored electronically, image capture techniques)?	YES □ NO □
•	Did the participants specifically consent to the related images or sensitive data?	YES □ NO □
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•	Could the study induce psychological stress or anxiety in participants or those associated with the research, however unlikely you think that risk is?	YES □	NO 🗆
•	Have you ensured that your research is culturally sensitive and respectful of diverse perspectives?	YES □	NO □
5. Biological Samples and Invasive Procedures			
•	Will blood or tissue samples be obtained from participants?	YES □	NO □
•	Is your research governed by the Ionizing Radiation (Medical Exposure) Regulations (IRMER) 2000?	YES □	NO □
•	Are drugs, placebos, or other substances (e.g., food substances, vitamins) to be administered to the study participants, or will the study involve invasive, intrusive, or potentially harmful procedures of any kind?	YES □	NO 🗆
•	Is pain or more than mild discomfort likely to result from the study? Please attach the pain assessment tool you will be using.	YES □	NO □
•	Will the study involve prolonged or repetitive testing or does it include a physical intervention?	YES □	NO □
•	Are you gathering data from laboratory animals?	YES □	NO □
•	Have you read ethical guidelines for using laboratory animals for medical research purposes? (https://grants.nih.gov/grants/olaw/guide-for-the-care-and-use-of-laboratory-animals.pdf)	YES □	NO □
6.	Incentives and Risk Assessments		
•	Will financial inducements (other than reasonable expenses and compensation for time) be offered to participants?	YES □	NO □
•	Have you undertaken Risk Assessments for each of the procedures that you are undertaking?	YES □	NO 🗆
7. Researcher-Participant Relationship and Conflicts of Interest			
•	Is there an existing relationship between the researcher(s) and the participant(s) that needs to be considered? For instance, a lecturer	YES □	NO 🗆





	researching his/her students, or a manager interviewing her/his staff, if yes,		
	please declare.		
		YES \square	NO □
•	Will there be transparency regarding funding sources and potential conflicts		
	of interest?		
8.]	Ethical Guidelines and Dissemination		
•	Is the research compliant with national ethical guidelines and regulations? (http://nbcpakistan.org.pk/assets/hbm-nbc-guidelines-final-18june-	YES □	NO □
	<u>2016.pdf</u>)		
•	Is there a plan to disseminate findings to participants and the wider community?	YES □	NO 🗆
9.	Contact for Ethical Reporting		
•	Have contact details been included for reporting and addressing ethical issues that may arise during the research?	YES □	NO □
10.	Collaboration and Partnership:		
	•		
•	Are you collaborating with other researchers or institutions?	YES \square	NO □
•	If yes, then mention the details of collaboration in the research synopsis.	YES \square	NO □
•	Do you have clear ethical guidelines and expectations established for your collaborators?	YES □	NO □
•	Have you considered involving the community in the research process to ensure that it is relevant and beneficial to their needs?	YES □	NO □
•	Is any of the research activity taking place outside of Pakistan?	YES □	NO □
11. Security-Sensitive Research			
•	Does your research fit into any of the following security-sensitive categories?		NO □
٥)	Commissioned by the military	$\mathbf{YES} \ \Box$	NO □
a)	· ·	YES □	NO □
b)	Involves the acquisition of security clearances	-	· • —
c)	Concerns terrorist or extreme groups	YES \square	NO □