## Research Ethics Board University of Peshawar



## **APPLICATION FORM**

#### Important:

- Please fill out every part of the form. Use simple words that anyone can understand.
- Submit a complete application with necessary documents (in original) to the office of the Secretary Research Ethics Board, University of Peshawar.
- Send a soft copy of the application to reb@uop.edu.pk

#### Required documents:

- i) Research Proposal
- iii) Consent form (English/Urdu)
- v) Proof of Funding Source
- ii) One Page CV of Applicant(s)/Supervisor/co-investigator(s)
- iv) Questionnaire (English and Urdu)

Section A	Application Details		
1. Project Title:			
2. Date of Submission: /	-	d Start Date: / d End Date: /	<i>I I</i>
3. Applicant (s) Details		Γ	
Full Name:			
Position held:			
(Researcher's designation/Graduate student's	level MPhil/PhD)		
Email:			
Contact No.		Signature	
Full Name:			
Position held:			
(Researcher's designation/Graduate student's	level MPhil/PhD)		
Email:			
Contact No.		Signature	
Name of Investigator/Supervi	isor (One-page CV must	be attached)	
Full Name:			
Position held:			
Affiliation:			
Address:			
Email:			
Contact No.		Signature	

Full Name: Position held: Affiliation: Address: Email: Contact No. Signature Section B **Details of the Project Brief Summary** of the Project in simple understandable language that a non-expert could understand (max 500 words)

Name of Co-investigator/ Co-supervisor (One-page CV must be attached)

Aim (s) and Objectives (max 200 words)		
Where will the study take place (Please pro	ovide area name for field work/ Department /Institu	ution)
, , , , , , , , , , , , , , , , , , ,		,
Details of subject/ participants (Tick √ w	here applicable)	
	here applicable)	
Details of subject/ participants (Tick √ w  1. Human  2. Animal	here applicable)	
<ol> <li>Human</li> <li>Animal</li> <li>Tissues</li> </ol>	here applicable)	
1. Human 2. Animal 3. Tissues 4. Genes	here applicable)	
<ol> <li>Human</li> <li>Animal</li> <li>Tissues</li> <li>Genes</li> <li>Genetically modified organisms</li> </ol>	here applicable)	
Human     Animal     Tissues     Genes     Genetically modified organisms     Cell Culture	here applicable)	
<ol> <li>Human</li> <li>Animal</li> <li>Tissues</li> <li>Genes</li> <li>Genetically modified organisms</li> </ol>	here applicable)	
Human     Animal     Tissues     Genes     Genetically modified organisms     Cell Culture     General		
Human     Animal     Tissues     Genes     Genetically modified organisms     Cell Culture		
<ol> <li>Human</li> <li>Animal</li> <li>Tissues</li> <li>Genes</li> <li>Genetically modified organisms</li> <li>Cell Culture</li> <li>General</li> </ol> Type of Animals' Research (Tick √ where		
<ol> <li>Human</li> <li>Animal</li> <li>Tissues</li> <li>Genes</li> <li>Genetically modified organisms</li> <li>Cell Culture</li> <li>General</li> </ol> Type of Animals' Research (Tick √ where Type of animals (Genus and Species)		
<ol> <li>Human</li> <li>Animal</li> <li>Tissues</li> <li>Genes</li> <li>Genetically modified organisms</li> <li>Cell Culture</li> <li>General</li> </ol> Type of Animals' Research (Tick √ where Type of animals (Genus and Species) Weight range		
<ol> <li>Human</li> <li>Animal</li> <li>Tissues</li> <li>Genes</li> <li>Genetically modified organisms</li> <li>Cell Culture</li> <li>General</li> </ol> Type of Animals' Research (Tick √ where Type of animals (Genus and Species)		

Detail of procedure using live animals (where ap	olicable)	
Complete procedure with reference if any		
Number of procedures		
Consent of the Human Subjects		
CONSENT		
In what form consent will be obtained (Mark √ the relevant)	Verbal	Written
Status of the subjects (Mark √ the relevant)	Healthy	Patient
State the reason if applying for waiving off the		
consent		
Major ethical issues (Attach extra sheets if required)		

## Participants Information (where applicable)

Participant Information and Consent	Yes	No
Will you inform participants that their participation is voluntary?		
Will you inform participants that they may withdraw from the research at		
any time and for any reason?		
Will you inform participants that their data will be treated with full		
confidentiality and that, if published, it will not be identifiable as theirs?		
Will you provide an information sheet that will include the contact details		
of the researcher/team?		
Will you obtain written consent for participation?		
Will you provide participants with written debriefing (i.e., a sheet that they		
can keep, showing your contact details and explanations of the study)?		
If the research is observational, will you ask participants for their consent		
to be observed?		

## Details and procedure of taking informed consent (where applicable)

Please describe the arrangements you are making to inform participants, before providing consent, of what is involved in participating in your study and the use of any identifiable data, and whether you have any reasons for withholding particular information. (No more than 200 words)		

# Potential risk and risk management procedures (where applicable) Identify, as far as possible, all potential risks (small and large) to participants (e.g. physical, psychological, etc.)

Identify, as far as possible, all potential risks (small and large) to participants (e.g. physical, psychological, etc.) that may be associated with the proposed research. Please explain any risk management procedures that will be put in place and attach any risk assessments or other supporting documents. Please answer within 200 words.

s of this project and will make sure to stick to the s. the approved research protocols without approval rovided in the project is correct and we agree to h subjects' rights and safety. nal reports after the completion of the study.
Signature of Investigator/Supervisor
Signature with official seal Head of the Department/Centre/College/Institution
r