Nazarbayev University Institutional Research Ethics Committee (NU IREC)

Appendix F: NU IREC Request for Continuing Approval Form

Part 1 - Adm	inistrative Inform	ation		
1. Protocol II	nformation			
Project Title:				
2. Contact in	formation			
2.1. Principa Name:	l Investigator (PI)			
Email addres	SS:			
School:				
Department/	Unit:			
Status: 🗆 Uı	ndergraduate Studer	nt 🗆 Graduate	Student Facult	ty 🗆 Staff 🗆 Oth
2.2. Please lis	st current member	s of the resear	rch team:	
Name	Email Address	NU/ Non-NU	School and Dept (If NU)	Name of organization (If Non-NU)
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Part 2 - Study Overview

	Please provide a lay summary of the study purpose and the general research questions/objectives.
s a H	Progress report since the last approval (please explain the progress of the study since the initial NU IREC approval or last approval, excluding amendment approvals) Please select all the types of research activities that were conducted on this protocol since last NU IREC approval (excluding amendment approvals):
\Box A	Active collection of data (not human biological materials or psychological data)
\Box A	Active collection and use of human biological materials or psychological data
agei	Use of psychological or biomedical devices, or drugs, biologics, or chemical nts
J	Jse of existing data (not human biological materials)
\Box A	Active collection of data (not human biological materials or psychological data)
IJ □	Jse of existing human biological materials
	ase summarize the research activities since last NU IREC approval (excluding endment approvals):
	be the last NU IREC approval (excluding amendment approvals), were there any icipant withdrawals from the study or complaints about the research activities? Wes \square No
	the last NU IREC approval (excluding amendment approvals), were there any expected problems or adverse events involving risks to participants? Wes \square No



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Since the last NU IREC approval (excluding amendment approvals), were there any				
changes to your study (including with recruitment, informed consent, study design				
and/or research procedures, research personnel, study location, etc.)?				
□ Yes □ No				
3. Research activities planned for the next year:				
3.1 Do you plan to recruit new participants?				
□ Yes □ No				
3.2 Do you plan to collect new or additional data from current research participants? \square Yes \square No				
Part 3 – Proposed Changes to Study Design 1. Plance calcut ALL the entegories of amandment(s) you are requesting:				
1. Flease scient ALL the categories of amendment(s) you are requesting.				
1. Please select ALL the categories of amendment(s) you are requesting: ☐ Change in Study Title				
☐ Change in Study Title ☐ Change in Principal Investigator				
☐ Change in Study Title☐ Change in Principal Investigator				
☐ Change in Study Title				
 □ Change in Study Title □ Change in Principal Investigator □ Addition of/change in research personnel 				
 □ Change in Study Title □ Change in Principal Investigator □ Addition of/change in research personnel □ Addition of/change in funding source □ Change to research/study design, methods, or procedures (e.g., observations, interventions, collection of biological samples or biometric information, 				



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\square Addition of/change to survey(s), questionnaire(s), or other research instruments - please attach the revised instrument/s
\Box Addition of/change to the identifiers collected in the study, or any others that would impact the privacy and confidentiality of the study participants
$\hfill\Box$ Addition of/change to informed consent/assent document(s) and/or procedures - please attach all related documents
☐ Other changes. Please specify:
2. Change in Principal InvestigatorName:
Email address:
School:
Department/Unit:
Status: \square Undergraduate Student \square Graduate Student \square Faculty \square Staff \square Other
3. Please state the reasons you are making amendments to the study.
 4. Are any of these changes the result of something that occurred during human participant interaction or an unexpected event? ☐ Yes ☐ No
If Yes, please explain:
5. How will the proposed changes have an impact on the risks or benefits to research participants?



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6. Do these changes involve information that a continue to take part in the research?☐ Yes ☐ No	might relate to a subject's willingness to
Signature	
This page is to be signed by the Principal Invean undergraduate or graduate student, the fallower box.	
Principal Investigator	
I certify that the information I provide in this also pledge that I will not change any of the plant this study without first seeking review and approximational Review Ethics Committee.	procedures, forms, or protocols used in
☐ Attestation of Principal Investigator	
Name / Signature of Drive in all Investigators	Data
Name / Signature of Principal Investigator	Date
Name / Signature of Faculty Supervisor	Date