

Research Proposal Approval Format

Research Title:
Institutional Review Committee (IRC)
Annapurna Neurological Institute & Allied Sciences
Maitighar, Kathmandu
Phone: 01-4256656/4256568, Hotline: 9801203340
Email: ircanias2018@gmail.com
For Official Use Only
(Please see the check list before Registration of the application form)
Registration No:
Date of registration:
Date of approval:
Name of Principal Investigator::
Total budget of the Project:
Site for the research work:
Tentative date for the start of the Project:
Duration of the Research Project:
Name of Internal Reviewer:
Name of External Reviewer:
Signature & Seal of IRC:

Part – I

Passport size photograph

Administrative Information

1. Research Title:
2. Name and title of the Principal Investigator responsible for the proposed research:
Last (Surname) Middle (if any) First name
Nationality:
Citizenship Number with the name of the district (only for Nepali)
Passport Number (only for non Nepali citizen):
Tussport Tussiaer (omy for non Tuspan etazen).
Signature: Date:
Postal Address:
Telephone No.:
Mobile No.:
Fax No.:
E-mail:
Alternate E-mail:

3. Full name of the Institution associated with the Principal Investigator (if	
applicable):	
Postal Address (if different from the address given above):	
Talanhana Na	
Telephone No.: Fax No.:	
e-mail:	
Website:	
4. Declaration of the head of the Institution (if applicable)	
If the proposed research is approved, we will allow him/her to conduct the research in this institution.	
Signature: Date:	
Last (Surname) Middle (if any) First name	
Designation:	
Signature: Date:	
Postal Address (if different from the address given above):	
Telephone No.:	
Fax No.:	
E -mail	

5. Name and Title of Co-investigators responsible for the proposed research (Use the similar format if more than one):

CO-INVESTIGATOR 1

Title:				
]
Last (Sur	name) Middle	e (if any)	First name	
Nationalit	y:			
Citizenship Nun	nber with name of the d	istrict from	where it was obtained (only	y for
Nepali)				
Passport Number	er (only for non Nepali o	citizen):		
Affiliated Institu	ution (if applicable):			
Designation:				
Signature:			Date:	
Postal Address	(if different from the ad	dress given	above):	
Telephone No.:				
Fax No.:				
E -mail:	·			

CO-INVESTIGATOR 2

Title:			
Last (S	Surname) Middle (if any) First na	ime	
Nation	ality:		
Citizenship N	Number with name of the district from where it	was obtained (only fo	r
Nepali)			
Passport Nur	mber (only for non Nepali citizen):		
Affiliated Ins	stitution (if applicable):		
Designation:			
Signature:		Date:	
Postal Addre	ess (if different from the address given above):		
Telephone N	o.:		
Fax No.:			
E -mail:			

(Use additional sheet if necessary)

6. List the name(s) and institutional affiliation to the researcher(s) (other than co-

Name	Institution and Address
(a)	
(b)	
	(Use additional sheet if necessary)
List the name(s) of Ne	epali researcher(s) (other than co-investigator) or Nepalese
Institution/hospital/No	GO(s) etc. from whom you may seek co-operation (if any)
(a)	
(b)	
(b)	
(b)	(Use additional sheet if necessary)
List major equipment(s	s) in relation to your research project you plan to bring/import
	s) in relation to your research project you plan to bring/import
. List major equipment(s	s) in relation to your research project you plan to bring/import
. List major equipment(s	s) in relation to your research project you plan to bring/import

	8.1 List details of all specimen(s) (if any) that you may transport from Nepal
	in relation to your research.
	(a)
	(b)
	(c)
	(d)
	8.2 Country of Destination:
	Name of Institution:
	8.3 Mode of Transportation of Specimen
	8.4 How will you ensure duplicate specimens remain in the country?
	(If necessary use additional sheet)
9.	Is this research part of your Thesis?
	Yes No
	If yes,
	For what degree and in which subject?
	From which university?
	From which country?

Part – II

Financial Information

10. Research T	l'itle:				
10. Name of th	ne funding o	organization:			
Contact infor	rmation of fu	nding organizati	on or agency	y:	
Postal Ad	ldress: Teleph	none No.:			
Fax No.:					
E-mail:					
Contact p	erson at the f	unding organiza	tion or agen	ey:	
Last (Sur	name)	Middle (if ar	ny)	First name	
Designation	on:				
Total amount	t of funds (in	NRs / US \$) allo	ocated for th	ne proposed research pr	oject:
Itemized bud research wor	-	•	resources re	quired for the proposed	1

Part – III

Research Proposal Description

12.	. Research Title:	
13	3. Proposal Summary (maximum 500 words):	

G	. C.1. D	11	ID C	1 / T .**	···			
Statemen	t of the P	roblem an	nd Rations	ale / Justif	ication (m	aximum 500) words)	
Statemen	t of the P	roblem an	nd Ration	ale / Justii	ication (m	aximum 500) words)	
Statemen	t of the P	roblem an	nd Ration	ale / Justii	ication (m	aximum 500) words)	
Statemen	t of the Pa	roblem an	nd Rations	ale / Justif	ication (m	aximum 500) words)	
Statemen	t of the P	roblem an	nd Ration	ale / Justii	cation (m	aximum 500) words)	
Statemen	t of the P	roblem an	nd Ration	ale / Justin	ication (m	aximum 500) words)	
Statemen	t of the P	roblem an	nd Ration	ale / Justin	ication (m	aximum 500) words)	
Statemen	t of the P	roblem an	nd Ration	ale / Justit	ication (m	aximum 500) words)	
Statemen	t of the Pa	roblem an	nd Ration	ale / Justif	ication (m	aximum 500) words)	
Statemen	t of the Pa	roblem an	nd Ration	ale / Justif	ication (m	aximum 500) words)	
Statemen	t of the P	roblem an	nd Ration	ale / Justif	ication (m	aximum 500) words)	

Research Objectives / purpose / aim of the study:
General
Specific

15. Research Design and Methodology Research Method Qualitative Quantitative Combined Study Variables: Type of Study (Specify): Study Site and Its Justification: Study Population (Specify): Study Unit: Sampling Methods / Techniques (Specify): Sample size (with justification):

teria for Sa	ample Selection:	
a Collecti	on Technique / Methods (Specify):	
a Collecti	on Tools: (please attached in annex)	
Pre-tes	ting the Data Collection Tools (if applicable):	
X7 1' 1'4		
Validity	and Reliability of the Study Tools:	
Potenti	al Biases (if applicable):	
Limitor	ion of the Study:	
	ion of the Study:	

16. Plan for Supervision and Monitoring:	
17 Plan for Data Managament and Analysis:	
17. Plan for Data Management and Analysis:	
18. Expected Outcome of the Research:	
_	
10. Dien fen Dissemination of Research Besults:	
19. Plan for Dissemination of Research Results:	
20. Plan for Utilization of the Research Findings (optional):	
20. I tan 101 e amzan of the research I memgs (optional).	
How is the research project going to strengthen the research capability of the h	ost
institution: Nepali Researcher (if submitted from aboard):	
21. Work Plan (should include duration of study, tentative date of starting the	
project and work schedule / Gantt chart):	
project and work schedule / Ganit chart).	

Part – IV

Ethical Consideration

Are human participants required in this research? If yes, provide justification. Yes (provide justification) No How many participants are required for the research? Explain. What is the frequency of the participant's involvement in the research? Explain. Clearly indicate the participant's responsibilities in the research. What is expected of the research participants during the research? Are vulnerable members of the population required for this research? If yes, provide justification. there any risks involved for the participants? If yes, identify clearly what are the expected risks for the human participants in the research and provide a	22. Regarding the human participants:					
How many participants are required for the research? Explain. What is the frequency of the participant's involvement in the research? Explain. Clearly indicate the participant's responsibilities in the research. What is expected of the research participants during the research? Are vulnerable members of the population required for this research? If yes, provide justification. there any risks involved for the participants? If yes, identify clearly what are the expected risks for the human participants in the research and provide a	Are human participants required in this research? If yes, provide justification.					
What is the frequency of the participant's involvement in the research? Explain. Clearly indicate the participant's responsibilities in the research. What is expected of the research participants during the research? Are vulnerable members of the population required for this research? If yes, provide justification. there any risks involved for the participants? If yes, identify clearly what are the expected risks for the human participants in the research and provide a	Yes (provide justification) No					
What is the frequency of the participant's involvement in the research? Explain. Clearly indicate the participant's responsibilities in the research. What is expected of the research participants during the research? Are vulnerable members of the population required for this research? If yes, provide justification. there any risks involved for the participants? If yes, identify clearly what are the expected risks for the human participants in the research and provide a						
Clearly indicate the participant's responsibilities in the research. What is expected of the research participants during the research? Are vulnerable members of the population required for this research? If yes, provide justification. there any risks involved for the participants? If yes, identify clearly what are the expected risks for the human participants in the research and provide a	How many participants are required for the research? Explain.					
Clearly indicate the participant's responsibilities in the research. What is expected of the research participants during the research? Are vulnerable members of the population required for this research? If yes, provide justification. there any risks involved for the participants? If yes, identify clearly what are the expected risks for the human participants in the research and provide a						
Clearly indicate the participant's responsibilities in the research. What is expected of the research participants during the research? Are vulnerable members of the population required for this research? If yes, provide justification. there any risks involved for the participants? If yes, identify clearly what are the expected risks for the human participants in the research and provide a						
Are vulnerable members of the population required for this research? If yes, provide justification. there any risks involved for the participants? If yes, identify clearly what are the expected risks for the human participants in the research and provide a	What is the frequency of the participant's involvement in the research? Explain.					
Are vulnerable members of the population required for this research? If yes, provide justification. there any risks involved for the participants? If yes, identify clearly what are the expected risks for the human participants in the research and provide a						
Are vulnerable members of the population required for this research? If yes, provide justification. there any risks involved for the participants? If yes, identify clearly what are the expected risks for the human participants in the research and provide a						
provide justification. there any risks involved for the participants? If yes, identify clearly what are the expected risks for the human participants in the research and provide a						
provide justification. there any risks involved for the participants? If yes, identify clearly what are the expected risks for the human participants in the research and provide a						
provide justification. there any risks involved for the participants? If yes, identify clearly what are the expected risks for the human participants in the research and provide a						
expected risks for the human participants in the research and provide a						
expected risks for the human participants in the research and provide a						
expected risks for the human participants in the research and provide a						
expected risks for the human participants in the research and provide a						
justification for these risks.						

	IRC/RES/PROP/Approval		
	Are there any benefits involved for the participants? If yes, identify clearly what are the expected benefits for the participants.		
3.	Informed Consent Form / Ethical Issues:		
Statements required in the Informed Consent Form include:			
	A statement that the human participants can withdraw from the study at any time without giving reason and without fear. State clearly how the participants can op out the study.		
	A statement guaranteeing the confidentiality of the research participants.		
	If required, a statement on any compensation that might be given to the research participant and or their community.		
	A statement indicating that the participants has understood all the information in the consent form and is willing to volunteer / participate in the research.		
	Signature space for the research participants, a witness, and the date.		
	(Informed Consent form should be submitted in English and in the language appropriate to the research participants)		
	Obtaining the Consent How informed consent is obtained from the research participants? Verbal Written		
	Please indicate who is responsible for obtaining informed consent from the participants in this research study?		

informed consent is being sought?

If yes, explain
Is the research sensitive to the Nepali culture and the social values?
Yes No Explain.
Is health insurance (<i>if applicable</i>) being made available to the research participants? If yes, please provide the necessary insurance data.
(Include in consent form)

Part - V

ACCEPTANCE OF GENERAL CONDITIONS AND DECLARATION BY THE PRINCIPAL INVESTIGATOR

I hereby certify that the above mentioned statements are true, I have read and understood the regulation of the Institutional Review Committee (IRC), ANIAS on the approval of research proposal and will act in conformity with the said regulation in all respects.

If the research is terminated, for any reason, I will notify IRC of this decision and provide the reasons for such actions. I will provide IRC with a written notice upon the completion of the research as well as a final summary/full report of the research study. If I publish the results in a journal, I shall acknowledge the IRC and shall provide the Committee with three copies of any such articles.

Signature of Applicant	Date:

INFORMED CONSENT:

Describe the manner in which informed consent will be obtained.

Indicate what kind of consent (e.g. parental, child, adult, etc) will be used.

If the subjects are children/adolescents ages 7-18 years, an Assent Form must be included with the IRC application. The signed Assent Form along with the Parental/Guardian Consent Form must be retained on file for at least three years after completion of the research project.

If prisoners / pregnant women, or fetuses are to be included in the research sample, it is likely that a full IRC review will be required and additional human subjects' protections will be expected.

If the subjects do not read or comprehend English, you must provide a consent form in their language as well as in English for IRC review and approval.

If you are requesting a waiver of written consent (i.e. a signature on an informed consent form) from the subjects, you MUST justify this request by providing an explanation of why obtaining written consent would add additional risk to the subjects and your alternative provisions for informing them about the study.

If consent documents from another site will be used, you will have to indicate this and provide a copy of the authorized consent document and IRC approval with your application.

You will have to provide any other relevant information if necessary. Please be aware that the PI is legally required to retain all signed Informed Consent forms for at least three years after the project terminates

The Informed Consent form must be written at a level that the subjects will understand. Please use simple language, and avoid clinical jargon.

Attach a copy of the written informed consent form (assent or parental consent where applicable). Consent documents MUST be in format requested. See examples on line on NHRC (Nepal Health Research Council) web site.

If the study uses database or archival data the use of informed consent is not applicable.

CONFIDENTIALITY OF DATA: Confidentiality of data MUST be address for all studies.

Indicate the extent to which confidentiality of records identifying subjects will be maintained.

Describe the storage and disposal of information where applicable.

Check List

For all applicants

- 1. Covering letter addressed to the Member secretary indicating the submission of the approval of proposal.
- 2. Proposal will only be accepted if submitted in IRC format.
- 3. Both printed and electronic version of the proposal should be submitted.
- 4. Curriculum Vitae of the Principal Investigator & Co-Principal Investigator of the study team should be submitted.
- 5. If the Principal Investigator is a non Nepali citizen, at least one Co-investigator should be a Nepali citizen.
- 6. Source of funding for the proposed project.
- 7. If the research study is to be conducted in any hospitals/organization or institution/community, a letter of approval from the related hospital/organization or institution/district authority should be provided.
- 8. Consent form should be in Nepali & local language (if necessary).
- 9. Data collection tools should be in Nepali & local language (if necessary) including interview guideline, observation checklist, questionnaires etc.
- 10. Style of referencing should be in Harvard style.
- 11. List of abbreviations / acronyms should be provided.

For students' applicants

1. Recommendation letter from Academic Supervisor along with above all lists.