USER MANUAL FOR

Researcher

Prepared BY



Nepal Health Research Council Kathmandu, Nepal

1.1.1 Dashboard

Note*:User need to complete each section before preceeding to next section .Document required should be uploaded.

	Government of Nepal Nepal Health Resea	arch Council				Q Notifica	tions 🚺	🖹 My Proposal 15	Account 👻
Das	shboard / My Dashboar	ď							
We	Icome, Bikash								
				Subm	it Another Proposal				
S.N	Created Date	Title	Applied Date	Status	Payment Status	Assigned Reviewers	Action		
1.	25th September 2016	Health Related	Not Applied Yet	Draft	Unpaid	N/A	View A	pply Payment Revisi	on Request
2.	12th September 2016	New Proposal14	Not Applied Yet	Draft	Unpaid	N/A	View A	pply Payment Revisi	on Request
3.	12th September 2016	Health	Not Applied Yet	Draft	Unpaid	N/A	View A	pply Payment Revisi	on Request
4.	11th September 2016	New Proposal	Not Applied Yet	Draft	Unpaid	N/A	View A	pply Payment Revisi	on Request
5.	11th September 2016	New Proposal	Not Applied Yet	Draft	Unpaid	N/A	View A	pply Payment Revisi	on Request

This is the main dashboard page where user can submit new proposal by clicking "submit new proposal" button. Proposal will be automatically saved after clicking "submit new proposal" button. Also, after the proposal has been created user can update their proposal and can apply whenever user wants. There are several steps for the proposal submission which will be described below:

- User can see their proposal status.
- They can view their proposal in brief by clicking view button. In notification user will get notified about proposal status, details, comments etc.
- In account module, user can view their profile; edit profile, change account password and logout.

1.1.2 Screening

Govern Nepa	^{ment of Nepal} al Health Research Council			Q Notifications 4	🕒 My Prop	osal 15 A	ccount +	
Dashboard	d / Proposals / Screening Infor	mation						
Step 1	Step 2	Step 3	Step 4	Step 5	Step 6	Step 7	Step 8	
Screening	Administrative Information	Financial Details	Technical Details	Ethical Consideration	Documents	Time line	Apply	
Screer	Screening Information							
	Does Your Research Relate	d To Health ? 🛞 Y	∕es © No					
	is This '	/our Thesis ? 🛞 Y	∕es © No					
	Educat	ional Level *	Select Educational Level		٣			
University Name* Please Enter Your University Name								
	Con	untry Name*	Select Your Country		Ŧ			
		S	ave and Continue Cancel					

Note: Field marked with * are Mandatory

Description

After clicking submit new proposal in dashboard, user will be redirect to this screening page. Here, users have to fill according to required inputs shown in user screen

Action : **save and continue** – will save information into database and redirect to administrative information page.

Cancel- will redirect to dashboard.

1.1.3 Administrative information page

Dashboar	rd / New Proposal / II	rwestigator					
Step 1	Step 2	Step 3	Step 4	Step 5	Step 6	Step 7	Step 8
Screening	Administrative Inforr	nation Financial Details	Technical Details	Ethical Consideration	Documents	Time line +Add I	Apply
	Investigator Type *	 Principal Investigator Co-Investigator 		Country *	Select Your Count	try	,
	First Name *	Enter your first name		Mobile No.*	Enter Mobile Num	ber	
				Phone No.*	Enter Phone Num	ber	
	Middle Name	Enter your middle name		Fax No.	Enter Fax Number		
	Last Name *	Enter your Last name		Email address.*	Enter email addre	22	
	Identification *	Select Identification Type	•	Alternative Email	Enter alternative o	mail address	
	Identification No. *	Enter your Identification No:		Investigator Image *	Choose File No f	le chosen	
	Institute Name	Enter your Institute name	in	vestigator Signature *	Choose File No f	le chosen	
De	signation of Principal Investigator	Enter your designation			No f		
Ins	titute Postal Address	Enter Institute Postal Address					
	Institute Fax No.	Enter Institute Fax Number					
	Institute Phone No.	Enter Institute Phone Number					
	Institute Email	Enter Institute Email Address					
	Institute Website	Enter Institute website					
	Save Note: I	Back Field marked with * are Mandatory					

Here, user can add investigator (either principal investigator or co- investigator). User can add investigator by fulfilling the required form.

User must upload photo, signature and CV of investigator in pdf format before preceding to next section.

Action: **save** – will save information into database.

Continue - will redirect to financial details page.

Back - will redirect Screening page.

1.1.4 Financial details

Step 1	Step 2	Step 3		Step 4	Step 5	Step 6	Step 7	Step 8
Screening	Administrative Information	Financial De	stalls	Technical Details	Ethical Consideration	Documents	Timeline	Apply
Financi	al Details							
	Cho	ose currency*	© Doll	ar ® Rupees				
	Human Ri	esource Cost*	Rs	100				
		Field Cost*	Rs	0				
	Lab	oratory Cost*	Rs	0				
	Data mana	gement Cost*	Rs	0				
	Report writing and Dissen	ination Cost*	Rs	0				
		Logistic Cost*	Rs	0				
	Monitoring and Eva	luation Cost*	Rs	0				
	Miscell	aneous Cost*	Rs	0				
	Ethical a	pproval Cost*	Rs	10000				
			if budg	get is below rs 10,00,000, only rs 1	0000 will be charged			
	Total budget of t	he research *	Rs	10100				
	Is This a Funde	d Research ?*	0 Yes	® No				
				Save and Continue	Back			
				Note: Field marked with * a	ire Mandatory			

Description

Here, users have to choose currency first. User can choose either dollar or nepali currency. After choosing currency, users have to insert cost according to the form inputs.

Total budget and ethical approval cost will be automatically generated. Ethical approval cost is \$100 if total budget is below \$10,000 and 3% of total budget if total budget exceed \$10,000.

If the research is funded, you need to upload agreement letter with donor clearly specifying the approved budget for the research.

Action: **save and continue** – will save information into database and redirects to Technical details page.

Back - will redirect to administrative information page.

1.1.5 Technical details

Screening	Administrative Financia Information	d Details	Technical Details	Ethical Consi	deration Documents	Timeline	Apply
Technical I	Details (Introduction)						
	Title of the research *	New Pr	roposal				
	Research area +	Non co	ommunicable disease				
Summary of t	he proposal (Structured)						
	Background *	andfan	df				
			m words: 30) r 30 words left.				
	Rationale/ justification *	asdfad	if		7		
			m words: 30) r 30 words left.				
	Objective *	asdfas	df				
					/_		
	Methodology *	asdfsa	I		1		
			m words: 30) r 30 words left.				
D	ata management and analysis *	asdfa					
Introduction							
	Background *	andf					
	paceground -	and a					
			m words: 30) r 30 words left.				
	Rationale/justification *	asdfa					
	General Objective *	asdfa					
	Specific Objective *	asdf					
					/_		
	Research Hypothesis*	asdf			_		
	Research design*	Ouanti	itatioa				
	Research design subtype*	Clincal					
J	ustification of research design+	asdfsa	1				
					/_		
			Save and Continu	e Back			

Here, users have to fill the technical details information and summary of the proposal as shown in the form.

Note:

In research design label, there are options to choose quantitative, qualitative and mixed research area. The form will be generated accordingly. If user will choose quantitative research area, then form will be generated for the quantitative research area. Similarly, respective form will be generated for qualitative and mixed research area. If user select quantitative research area and research design subtype as clinical trial, then he/she have to add monitoring board after ethical consideration page.

Action: **save and continue** – will save information into database and redirects to Ethical consideration page.

Back – will redirect to the financial details page.

1.1.6 Ethical Consideration



User manual

Collection Clas: X X Introduction to X X Upcoming even X X F Facebook - Log 3	X 📓 No Mountain w X 🔓 user manual sa X 🔭 Ethic	al Conside X G Google X Paras - 0 X
\leftrightarrow ∂ C 0 192.168.10.11:8001/researcher/ethical-consideration-technical-details/12	2	Q☆ (2 :
Who is responsible for obtaining informed consent? *	adf	·
Is there anything being withheld from the research participants at the time the informed consent is being sought? *	® Yes © No	
if yes, explain *	asdf	
is the research sensitive to the Nepali culture and the social values? *	® Yes © No	
is health insurance (if applicable) being made available to the research participants? *	asdfa /	
Regarding Clinical Trial		
The trial treatment *	asdfa /	
A detailed explanation of the trial procedures including all invasive procedures. *	asdfa 🦯	
The potential or direct benefits (if any) for the research participants *	asdfaa	
Alternative procedure(s) or treatment(s) that may be available *	asidfa /	
The risks, discontions, and inconveniences associated with the study \ast	asdfsa /	
Provisions for management of any adverse reactions *	adsfa	
The provisions of insurance coverage for any permanent disability or death *	asdf	
Address of contact person if adverse effects		
Contact Person Name *	adsfa	
🚱 🖉 🚺 Mi docs 🧿 Ethical 🔰 🍓 🕓 Skype 🔯 In	box 🔯 Inbox 📲 User 🔮 user 🞻	Ethical Kil do Ethical 🔺 🛱tl 🕪 4:44 PM

← → C ① 192.168.10.11:8001/researcher/ethical-consideration-technical-details/12	1	Q & Q :
Provisions for management of any adverse reactions *	adsfa	
The provisions of insurance coverage for any permanent disability or death *	asdf	
Address of contact person if adverse effects		
Contact Person Name *	adsfa	
Contact Person Mobile *	asdfa	
Contact Person Email *	b@gmail.com	
Name of affliated Organization *	ajskdfh	
Designation of the organization *	asdfh	
Phone number of the organization *	jasdfh	
Email address of the organization *	jkasdfh@gmail.com	
Is there going to be a transfer of any biological materials from the Country ? *	® Yes © No	
Type of sample to be transferred *	blood	
Country where sample will be taken *	NE – Niger	
Laboratory involved in testing samples *	adsfas //	
Test to be performed in the lab *	asdfsa //	
Main person involved in lab test *	asdfa //	
Qualification of the person involved in lab testing *	asidfas	
🛿 🚱 🔏 👖 Mildocs 👩 Ethical 🔉 🍪 🏾 S Skype 🔯 In	box Grund Information	I 💥 Mildo Ethical 🔺 🛱 📶 🌒 2015 PD45

Qualification of the person involved in lab testing st	asdfas
Upload CV of person involved in lab testing st	Choose File No file chosen
Does the study involve transfer of DNA sample? *	Yes No
Type of sample *	 Extracted Extracted and amplified Whole
Name of Laboratory in Nepal involved in extraction and amplification *	asdfa
Contact person of the laboratory st	asdfas
Mobile number of the contact person *	asdfa
Landline *	asda
Email address *	b@gmail.com
Does it have Data safety Monitoring Board ? *	Ø Yes O No
Number of members in data safety monitoring board st	1
	C Save and Continue Back

Here, user can full fill the form according to the required inputs as shown in the form.

Action: **save and continue** – will save information into database and If user had select quantitative research area and research design subtype as clinical trial, then he/she have to add monitoring board after this page else it will be redirects to documents page.

Back – will redirect to the technical details page.



1.1.7 Documents

Here, user have to upload documents according to form inputs. Users must upload data collection tool, informed consent form and work plan to go to the next section. If the study involves participants below 18 years of age, please upload assent form as well.

Action: save and continue – will save information into database and redirects to timeline page.Back – will redirect to the Ethical Consideration page.

1.1.8 Timeline



Here, users have to add start date and end date.

Action: save and continue – will save information into database and redirects to apply page.

Back – will redirect to the document page.

1.1.9 Apply



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Here, in this page there are two buttons (preview before apply and apply button). By clicking preview before apply button, user can see their proposal summary in brief and by clicking apply button proposal will be applied successfully. User can save the pdf summary of the proposal.