

Research Management and Processing System

is a program designed to manage the submission of research proposal and its' processing. It also provides a platform for the users to manage their daily tasks and generate reports as needs.

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System Services

- Submission of a new research work
- Dashboard
- Requesting a specific service (study extension , closure , amendment and reactivate)
- Print out Approval Memos
- Provide Active Timeline
- Ability to Report personal research activities
- User support (on progress)

First time use

- MNGH member :
 - 1. Access the system using your login credential
 - 2. Upload your CV
 - 3. Request P.I role (in case you will submit a proposal)
 - 4. Upon approval of P.I role then you will be able to submit a research proposal
- Non-MNGH member:
 - Should sign up first

Login Page

All MNGH users have a default accounts that are accessible using organization credential to Login. Users are grouped in to four main categories, the fourth category for non MNGHA users who has not credential to Login and need to sign up

- 1. Hospital users (who has hospital credential)
- 2. University users (who has University credential)
- 3. KAIMRC users (who has KAIMRC credential)
- 4. External users (non MNGHA staff)



eSubmission

Select User Type.	
Please Select User Type	~
Enter email and password.	
🔤 Email	
Password	
F	orgot Password?
	orgot i assirora.
RXKN	
ERXKN	
ERXKN	

Don't have an account yet? Sign Up!

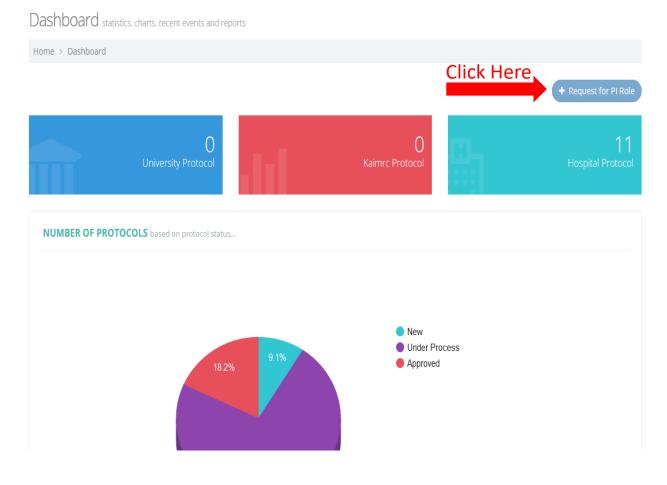
Uploading/Updating CV

- First step after accessing the system is to upload CV, researchers cant proceed with proposal unless they have approved P.I role which required CV review by the team.
- Research team still can update their CV any time by accessing 'CV document'at tab and uploading the new CV.

ubmission	
Dashboard	CV DOCUMENT
rotocol <	UPLOAD CV*
CV Document	Browse No file selected.
Clinical Trial Document General Document	+ Upload Attachment
Jseful Links and Policy	File Name 🔺 View 🔶 Delete 👙
Logout	72442.pdf View Delete
	<u>(</u> 1)
	 Acceptable formats: pdf and docx File Size can't exceed 30 MB

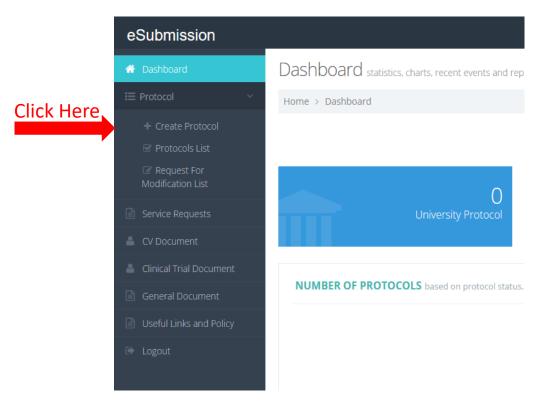
Requesting P.I role

P.I role request is considered the second step after uploading CV, upon this function approval the P.I can create and submit a research work.



Submitting a Research Proposal/Protocol

Step 1: Once the P.I role is accepted, then the P.I can access the function create protocol from the sidebar by clicking on protocol option



Submitting a Research Proposal/Protocol

Step 2: the P.I will fill up the basic information, then click on save protocol information

PROTOCOL INFORMATION					
Title *					
Organization *		Department *			
Please Select	~	Please Select		~	
Site *		Type of Project *			Area of Study *
Please Select	v	Please Select		v	Please Select
Study Design *		Grant *			Amount
Please Select	~	Please Select		~	
Aim of the Study *			Scope *		
			Please Select	-	
Specific Objectives *			Secondary Objectiv	ves	
Specific Objectives *			Secondary Objectiv	ves	

Submitting a Research Proposal/Protocol

Step 3: a new field including co-authors information will be available. the P.I is required to fill up all mandatory information and upload all necessary documents then click on the bottom submit protocol

SUBMIT PROTOCOL

Adding co-author

• The P.I can search for the co-author the this section

RESEARCH TEA	M: SUB-INVESTIGATOR *			
Search By	Proposed Name	Email	• Badge	T Search
Scarch by				× Clear

• Then choose the applicable co-author and click on save sub-investigator, one by one.

Select	Name	Department	Email	Badge	Job Position	cv
•	Potential co-author name	RESEARCH OFFICE				Download
					🖺 SAVE SUB INVE	STIGATOR

Print out Approval Memo

This function is available upon final decision by the final approval authority.

OTOCOI Information\ PPDU Approved					Click Here
		🔒 Print	O View Timeline	X Close Proposal	Show Mem
		🛓 Download II	RB Approval Memo-2	2 🕹 Download IRB	Approval Memo-
ROTOCOL INFORMATION					
PROTOCOL INFORMATION					
Protocol No. *	Submission Date		Study Status		
ICTR24/007/2	02/25/2024		Under Proces	S	
Title *					
clinical trail memo					
Organization *	Department *				
Hospital	∽ NGHA2	~			
PPDU Status	IRB Status		Fund Approval S	Status	
PPDU Approved	IRB Approved		N/A		
Site *	Type of Project *		Area of Study *		
Riyadh	← CT : Clinical Trial	~	Blood and car	ncer Research Program	n v

Timeline

Time is accessed through accessing the submitted protocol , then click on '**View Timeline**' option the upper right side of information screen. This option help the P.I to identify the progress on proposal processing classified per task and the assigned department.

OTOCOI Information\ N/A			Here	
		🔒 Print	O View Timeline	X Close Proposal
PROTOCOL INFORMATION				
PROTOCOL INFORMATION				
Study Status				
New				
Title *				
testing				
Organization *	Department *			
Hospital	NGHA2 ~			
PPDU Status	IRB Status	Fund Approval	Status	
N/A	N/A	N/A		
Site *	Type of Project *	Area of Study	*	
Riyadh	RM : Regular Research from western region – N \sim	Population H	lealth Research Progr	am 🗸

Timeline

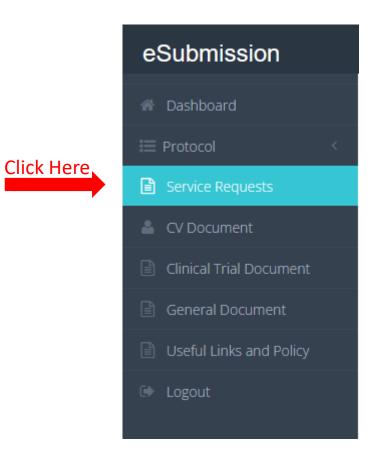
Proposal Time Line

© TIMELINE	
START AT :	
15/02/2024 01:47 PM	Proposal Created by Name of P.I Proposal # 1082
15/02/2024 01:47 PM	Proposal Submitted Submission date
18/02/2024 04:53 PM	PPDU Approved proposal Approval by the first authority
END	

Request a service

This option available once the proposal already processed or approved, the P.I can request a service of : study closure, By just accessing the target protocol then at the bottom click on request a service and select the needed service.

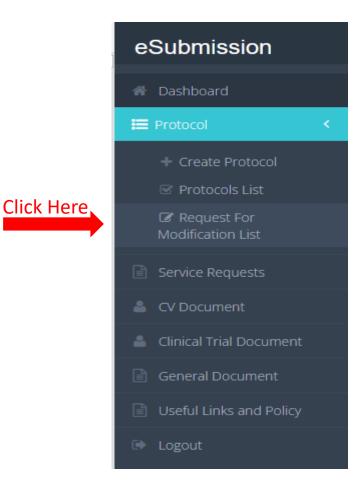
You can see your requested services by accesing 'Service requests'



Modification Request

If modification is needed, the P.I will find the request to modify the proposal on the sidebar by clicking over request for modification, then a response to the modification should be made

*Please note that if no response within 10 days to the modification request, your protocol will be deactivated automatically.



Necessary Document and educational links

To avoid any rejection or delaying in the approval process, the respected P.I should upload all necessary documents, please refer Clinical Trail Document, General document, and useful links and policy to download needed template. Clinical Trial Document
 General Document
 Useful Links and Policy