

**ETHICS REVIEW COMMITTEE FOR THE RESEARCH  
IN LIBRARY & INFORMATION SCIENCE (LIS-ERC)**

<b>For Office Use Only</b>
<b>Application Number: LIS-ERC/ NLSL/2023 / 001</b>
<b>Date Received: ____ / ____ /20 ____</b>

**APPLICATION FORM**

This application should be filled and signed by the principal investigator who requests ethical approval for a research project. All co- investigators should provide consent to submit the application to LIS-ERC by signing the application. Please read the guidelines for application available at NLSL website carefully before filling out the application.

**Please note that applications cannot be considered for ongoing projects. Along with this application, please submit a detailed research proposal (synopsis)/protocol, consent forms, checklist and all other relevant documents necessary.**

**This application form consists of three sections:**

SECTION A: GENERAL INFORMATION – RESEARCHER DETAILS

SECTION B: RESEARCH DETAILS – ETHICAL ISSUES

SECTION C: RESEARCH DETAILS- REVIEWER CHECK LIST

**Applicant must complete all three sections and attach research proposal (according to the guidelines provided by the NLSL).**

## SECTION A – GENERAL INFORMATION - RESEARCHER DETAILS

### A.1. TITLE OF THE RESEARCH:

Location of research (e.g. Institutions, communities)	Expected Date of Commencement of Research:  Expected Date of Completion of Research:

### A.2. Details of the Investigators (Should provide principal investigator's name first).

	Title: (Rev./Prof./Dr./Mr./Ms): Name, Designation and Affiliation	Role (State whether principal investigator, co-investigator, supervisor).	Signature
01.			
02.			
03.			
04.			

### A.3. Contact Details of the Principal Investigator

3.1	Postal Address	
3.2	Email Address	
3.3	Telephone	Official:  Mobile:

**SECTION B – DETAILS OF THE RESEARCH – ETHICAL ISSUES**

**Please provide answers to all of the following questions.**

**B.1. Provide a brief description of your research (Background of the study, aims and objectives, data collection, sampling, methodology, chronological/ geographical distribution etc.)**

**Use additional sheet if necessary.**

**B.2. Has the relevant board of study approved the research project (if applicable)?**

Yes:                   No:                    Pending Approval

**If Yes, Details:**

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**B.3. Has ethics approval for this study been requested earlier from LIS-ERC of NLSL?**

Yes :                   No:

**If Yes, ERC NO: .....**

**B.4. Funding (if any)**

Yes :  No:

Name and address of the funding source: (**Attach** relevant details)

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.....

**B.5. Do you believe the proposed project has conflicts of interest?**

Yes:  No:

**If Yes, Details:**

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.....  
.....

## SECTION C: RESEARCH DETAILS - REVIEWER CHECK LIST

(Applicant should indicate the number of the protocol section where each issue is addressed in their research proposal. If a particular issue is not relevant to your study indicate that as ‘NA’) (Eg: if your objectives appear in pg.no.3, of the proposal then mention 3 as the protocol pages).

		For Applicant	For Reviewer			
C.1	Social Value	Protocol Page/s	Reviewer Evaluation			
			Acceptable			Comments
			Yes	No	NA	
I	Benefits of the study to the community / society		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
II	Plan for dissemination of study findings		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
III	Scientific importance of the study		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	

C.2	Scientific validity	Protocol Page/s	Reviewer Evaluation			
			Acceptable			Comments
			Yes	No	NA	
I	Title		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
II	Research problem		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
III	Research questions/hypothesis		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
IV	Objectives		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	

V	Study setting		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
VI	Study design		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
VII	Study population (giving inclusion exclusion criteria)		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
VIII	Sample size		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
XI	Sampling method		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
X	Measurements / variables		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
XI	Study instruments (data collection method/s)		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
XII	Procedures to ensure quality of data		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
XIII	Plan for analysis		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
XIV	Ethical considerations		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
XV	Budget (if relevant)		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
XVI	Work plan and time frame		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	

XVII	Justification for a replication study, if your study is a replication.		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
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**\*NA- Not Applicable**

C.3	Fair participant selection and vulnerability*	Protocol Page/s	Reviewer Evaluation			
			Acceptable			Comments
			Yes	No	NA	
I	Justification for selection of study population		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
II	Justification for conducting the study in a vulnerable population		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	

**\*children, women, people with disabilities, people with psycho-social issues, sexually marginalized populations etc..**

C.4	Participants rights and consent	Protocol Page/s	Reviewer Evaluation			
			Acceptable			Comments
			Yes	No	NA	
I	Procedure for recruiting the participants		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
II	Information provided to the participants		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
5	Procedure for obtaining informed consent		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
IV	Procedure for obtaining proxy consent		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	

V	Procedure for withdrawing consent		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
VI	Incentives provided to participants		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
VII	Procedure for participants to ask questions / lodge complaints		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
VIII	Participants right to decline consent		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	

C.5	Confidentiality and Privacy	Protocol Page/s	Reviewer Evaluation			
			Acceptable			Comments
			Yes	No	NA	
I	Steps to ensure confidentiality of data		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
II	Justification for collecting personal identification data		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
III	Steps taken to ensure privacy during data collection		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
IV	How long data and samples will be kept		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
V	Who will have access to the data		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
VI	Procedure for storage of data and samples		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	



VII	Procedure for disposal of data		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
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C.6	Risk Benefit Assessment	Protocol Page/s	Reviewer Evaluation			
			Acceptable			Comments
			Yes	No	NA	
I	Potential risks to the participants		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
II	Potential benefits to the participants		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
III	Justification for risks against benefits		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
IV	Steps taken to minimize risks		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	

C.7	Responsibilities of the researcher	Protocol Page/s	Reviewer Evaluation			
			Acceptable			Comments
			Yes	No	NA	
I	Ethical, legal, financial issues related to the study		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
II	Any conflicts of interest and how the researcher plans to manage them		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
III	Permissions from relevant institutions / authorities		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
IV	Collaborations with the relevant stakeholder		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
V	Qualifications of the research team to handle the research study		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	

C.8	Foreign funded studies	Protocol Page/s	Reviewer Evaluation			
			Acceptable			Comments
			Yes	No	NA	
I	Justification for conducting the study in SL		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
II	Relevance of the study to SL		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
III	Post research benefits to SL		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
IV	The sharing of intellectual property rights		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
V	How the results will be conveyed to authorities in SL		<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	

\* SL – Sri Lanka

**SECTION D- DECISION NOTIFICATION BY THE REVIEWERS**

**For Office Use Only - To be filled by the reviewers:**

Decision of the Reviewer:

Approved	<input type="checkbox"/>
Conditional Approval (please mention the conditions below)	<input type="checkbox"/>
Resubmit (please mention the reasons below)	<input type="checkbox"/>
Reject (Reasons can be given under the comments that can be communicated to the principal investigator)	<input type="checkbox"/>

Give reasons for your decisions:

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(Use additional sheets if necessary).

Name of the Reviewer/s	Signature of the Reviewer/s	Date

Confirmed by LIS-ERC secretary: Signature \_\_\_\_\_ Date \_\_\_\_\_

**LIS-ERC DECISION NOTIFICATION**

**Name of Applicant: Prof./ Dr./Mr./Ms.** .....

**Application Number: LIS-ERC/ NLSL/20** \_\_\_\_ / \_\_\_\_

**Date received:**

...../...../.....

**LIS-ERC Meeting Date:**

...../...../.....

Thank you for submitting the ERC application for the research title

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..... It was considered by the ethics review

committee at its meeting held on .....The LIS-ERC decision is as follows;

Approved	<input type="checkbox"/>
Conditional Approval	<input type="checkbox"/>
Resubmit	<input type="checkbox"/>
Reject	<input type="checkbox"/>

If LIS-ERC recommends conditional approval/rejection the comments of the reviewers are attached for your information.

.....  
Secretary  
LIS-ERC  
National Library of Sri Lanka

Date: .....

## Application Form for Ethics Review – Submission Check List

**For Office Use Only:**

Application Number: **LIS-ERC/ NLSL/20** \_\_\_\_\_ / \_\_\_\_\_

Date Received: \_\_\_/\_\_\_/20\_\_

**To be filled by the Applicant:**

**Title of the Research:**

**Name of the Principal Investigator:**

Document	Version	Date
1. Application form (3 copy)		
2. Detailed Research Proposal (3 copies)		
3. Summary of the Research Proposal - One page (3 copies)		
4. All study instruments (questionnaires/interview guides/checklist/data extraction forms) in English (3 copies)		
5. All study instruments – Sinhala Translation if Applicable (3 copies)		
6. All study instruments – Tamil Translation if Applicable (3 copies)		
7. Information Sheet in English (3 copies)		
8. Information Sheet -Sinhala Translation if Applicable (3 copies)		
9. Information Sheet - Tamil Translation if Applicable (3 copies)		
10. Consent Forms in English (3 copies)		
11. Consent Forms - Sinhala Translation if Applicable (3 copies)		
12. Consent Forms - Tamil Translation if Applicable (3 copies)		
13. Any other relevant documents in (3 copies)		
14. Any other relevant documents - Sinhala Translation if Applicable (3 copies)		
15. Any other relevant documents - Tamil Translation if Applicable (3 copies)		
16. Short curriculum vitae of all investigators (Maximum 2-3 pages) (3 copies)		

**I understand that the application for ethics clearance will not be accepted unless all documents are submitted.**

\_\_\_\_\_

**Signature of the Principal Investigator**

\_\_\_\_\_

**Date**

## Declaration

01. As the principal investigator of this project, my signature confirms that I will ensure that all procedures performed under the project will be conducted in accordance with all relevant national and international policies and regulations that govern research involving people.
02. I understand that if there is any significant deviation from the project as originally approved, I must submit an amendment to the LIS-ERC of the NLS for approval prior to its implementation.
03. I have submitted all significant previous decisions by this or any other ERC and /or regulatory authorities relevant to the proposed study.
04. I declare that I am not seeking approval for a study that has already been commenced or has already been completed.
05. I understand that at least two months are required for ethics review and granting ethics clearance.
06. I promise to report on all developments and drawbacks during the implementation of the research in accordance with the ERC guidelines.

\_\_\_\_\_  
Signature of principal investigator

\_\_\_\_\_  
Date

Full name of principal investigator:

\_\_\_\_\_  
\_\_\_\_\_

NIC/ Passport Number: \_\_\_\_\_ (Attach a copy of the NIC or pasport).

The application processing fee of Rs.2000/- should be paid to ERC fund (A/C No. 167-1-001-6-3170315 of people's bank. Town hall Branch.

(Attach the bank payment slip)

**Emil all relevant documents to [ethicsreview@lis.natlib.lk](mailto:ethicsreview@lis.natlib.lk)**