

SRI LANKA ACCREDITATION BOARD for CONFORMITY ASSESSMENT

APPLICATION FORM for RECOGNITION of GOOD LABORATORY PRACTICE

Instructions to the Applicant:

Please submit this application along with the questionnaire, duly filled, Laboratory's GLP Manual and associated documents referred in the application and questionnaire.

Director /CEO, Sri Lanka Accreditation Board for Conformity Assessment, No. 104/A, Kitulwatte Road, Borella



	APPLICATION FOR RECO	GNITION OF GOOD	LABORATORY PRAC	TICE
We a _l	apply for recognition of Good Laboratory Pra	octice/s of our laboratory a	as per the details given below	r:
	First Recognition Sc	ope Extension	Renewal of Recognition	
1.	Laboratory Details			
1.1	Name of the Laboratory			
	Address			
	Telephone			
	Fax No	e-mail		
1.2	Name of Parent Organization (if laboratory is a part of an organization)			
	Telephone No.	Fax No	e-mail	
1.3	Legal status and date of establishment (please give Registration No. and name of authority)			
2.	Recognition Details			
2.1	Field of studies for which recognition (please tick the appropriate box, a separate applic		ine)	
	Physical-chemical studies	Studies	related to ecosystems	
	Toxicity studies	Studies	related to medicines	
	Residue studies	• Ergono	mics studies	
	Safety studies	Other (Please specify)	

2.2 If the Laboratory is already recognized for GLP, indicate the Scope & Good Laboratory Practice for which recognition has been granted.

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2.3		Laboratory is already accredited for ISO/IEC 17025/ISO 15189, indicate the fields of testing and for which accreditation has been granted.				
2.4	Scope	e of Recognition s	ought for GLP			
	no n	Group of products, materials or items studied	Specific or types of tests/studies performed	Method of Study Ref. No, Code No.	Range of measurement/ Limit of detection	
	Note	e 1. Laboratories perfor		clearly identify the specific te on	product(s)/ material performed at	
3.	Orga	nization				
3.1	Labor	ratory Managemen	t (Name, Designation, telephone,	Fax, e-mail)		
	3.1.1	Chief Executive of	of the laboratory			
	3.1.2	Person responsib	ole for the test system			
	3.1.3	Study Director				
	3.1.4	Authorized Repre	esentative for SLAB ——			

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3.1.5	prepared towards SLAB a. Development of Labor b. Development of Techr c. Specific Training: d. Conducting Internal Au	accreditation; atory Management System nical Operations: ————	tion that has provided cons	, .		
Organ	ization Chart					
	Indicate in an organization is being sought (please app		ments of the testing facility for	which recognition		
	Indicate how the testing fa (where applicable)	acility is related to external	organizations or to its own pa	arent organization		
Emplo	pyees					
3.3.1	Total number in the testing	ng facility for the specific fie	eld applied			
3.3.2	Total number in the testin		tion is being sought			
3.3.3	Details of staff (please clear	rly indicate the staff responsible fo	or site testing)			
SI no	Name	Designation (Study Director, Principle Investigator/ Investigator/other)	Academic and Professional Qualifications*	Experience related to present work (in years)		
	* Please clearly indicate the field o					

3.2

3.3.

3.3.4 If Trainees or Contracted persons are employed, Please indicate the details of them.

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^{*} Please clearly indicate the field of specialization

4. Equipment and Reference Materials

4.1 Equipment List

please list down all significant items of equipment, providing details of make, model, serial number, range, if applicable and calibration status (date of last calibration, name of calibrating authority), if available.

The preferred order is: a) Reference equipment - Weights, balances, thermometers etc;

- b) Testing equipment spectrophotometers, testing machines etc;
- c) Ancillary equipment sieves, autoclaves, etc;

SI no	Name of equipment	Model/ type/ year of make	Receipt date & date placed in service	Range and accuracy	Date of last calibration	Calibration due on *	Traceability**

4.2 List of reference materials

please list down all reference materials used for verification or validation of test method or technique applied for Accreditation

SI.	Name of reference material/	Source	Date of expiry/ validity	Source of
no.	strain/ culture			Traceability

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5. Quality Assurance Programme

Please list down the details of Quality Assurance programmes currently employed by the Laboratory

SI. no.	Study method or group of studies applied for recognition	QA programme	Service provider	Frequency

6. Willingness to undergo Assessment

We declare that

- 6.1 We are familiar with and will abide by the terms and conditions of maintaining SLAB recognition for GLP included in the agreement to be signed by both parties, which is enclosed.
- 6.2 We agree to comply fully with SLAB GLP Criteria for recognition of test facilities.
- 6.3 We agree to comply with GLP procedures, pay all costs for assessment, verification visit (if any), surveillance and reassessment irrespective of the result.
- 6.4 We agree to co-operate with the assessment team appointed by SLAB for examination of all relevant documents by them and their visits to those parts of the laboratory that are part of the scope of recognition.

Signature of Chief Executive or his authorized representative	
Name & Designation	
Date & Place	

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