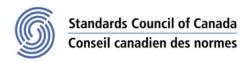


Application for Recognition of GLP Compliance

		Identify (√)
	Test Facility: Test Site ¹ :	Pesticides: Industrial Chemicals: Other (specify):
		Applicants Legal Name
		Street Address
	Postal Addr	ress (if distinct from street address)
	Cit	ty / Province / Postal Code
	Telephon	ne No. / Fax No. / E-mail address
(1998) fo above) u		nce to the <i>OECD Principles of Good Laboratory Practice</i> iple, sites or if the street address is distinct from that provided llow (¹ Field Sites are Test Sites):
1	Facility Legal Name	Street Address
	City / Province / Postal Code	Telephone No. / Fax No. / E-mail Address
2	Facility Legal Name	Street Address
	City / Province / Postal Code	Telephone No. / Fax No. / E-mail Address

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3.		
	Facility Legal Name	Street Address
	City / Province / Postal Code	Telephone No. / Fax No. / E-mail Address
4	Facility Local Name	Street Address
	Facility Legal Name	Street Address
	City / Province / Postal Code	Telephone No. / Fax No. / E-mail Address

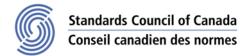
Area(s) of Expertise	Identify (//)			
	1	2	3	4
Physical-chemical testing				
Toxicity studies				
Genotoxicity/Mutagenicity studies				
Environmental toxicity studies on aquatic and terrestrial organisms				
Studies on behavior in water, soil and air; bioaccumulation				
Residue studies (field phase)				
Studies on effects on mesocosms and natural ecosystems				
Analytical and clinical chemistry testing (product preliminary analysis, residue and/or environmental)				
Toxicokinetics				
Other studies (specify)				

Recognition is being sought in accordance with the identified option²:

Recognition Options ²		Identify (//)			
	1	2	3	4	
A full inspection: a facility inspection and study audit(s) for facilities which have previously conducted GLP studies.					
A facility-only inspection without a corresponding study audit(s) for facilities which have not as yet conducted GLP studies.					

²Refer to CAN-P-1583, section 7.3.1 for description of recognition options.

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An application package is enclosed and includes:

- i) a completed Application for Recognition of GLP Compliance form;
- ii) a facility organizational chart and facility floor plan;
- iii) a current Master Schedule;
- iv) a list of all current SOPs under headings which are consistent with clause 7.4 of the *OECD Principles of Good Laboratory Practice* (1998);
- v) for test sites, a response to the questions found on Annex A; and
- vi) an appropriate application fee according to the published SCC GLP recognition fee structure.

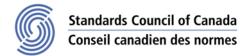
On granting of GLP recognition, the applicant agrees:

- i) to comply with the requirements and conditions contained in the latest edition of the Standards Council of Canada document, *SCC Monitoring Authority Requirements for the Recognition of GLP Compliant Test Facilities* (CAN-P-1583);
- ii) to pay the required annual and visit fees;
- that recognition does not imply that GLP studies are approved or endorsed by the SCC, or by any government agency or any certifying body; and
- iv) that GLP recognition may be withdrawn, on failure of the facility to continue to comply with the foregoing, subject to Standards Council of Canada appeal procedures as referenced in CAN-P-1583 and thoroughly described in the SCC appeals document, CAN-P-15.

SIGNATURE	
NAME	
TITLE	
DATE	

Application forms in electronic format are available upon submitting an email request to the SCC at: info.palcan@scc.ca. Cette forme est aussi disponible en français.

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ANNEX AGLP Test Site Questionnaire

Please respond to the following questions. The answers are particularly relevant to facilities operating as test sites in multi-site studies in order to establish how the corresponding requirements of the *OECD Principles of GLP* are met.

	nagement (test facility/test site), Study Director(s), Principal Investigator(s), Quality Assuran gram(s) and study personnel?
)	Do any personnel at your test site ever function in the role of a Study Director?
)	How are QAP functions conducted (e.g. sponsor, in-house; contract QA)?
	How are your records and materials archived (e.g. on-site, contract, transferred to Study Director)?

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