OECD GLP Registration Procedure Chart

กรมวิทยาศาสตร์การแพทย์
Department of Medical Sciences

BLQS
BUREAU OF LABORATORY QUALITY STANDARDS
DEPARTMENT OF MEDICAL SCIENCES

1 Application submission documents

Requirements and Conditions for >>
GLP compliance test facility

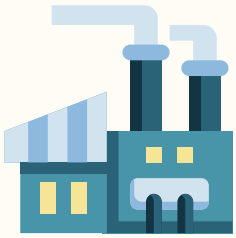
Application form No. 5 - 7 >>

2 Payment of fee

<< ระเบียบกรมวิทยาศาสตร์การแพทย์ ว่าด้วยอัตราค่าบำรุง การตรวจวิเคราะห์และให้บริการ พ.ศ. 2562
"การตรวจสอบ Test Facility ตามหลักการ OECD GLP " (หน้า 192)

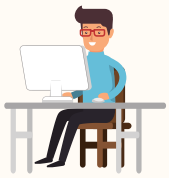
3 Pre-inspection (1 day)

Corrective action 12 month
(6-12 month)



4 Full inspection/ Study audit (2-5 days)

5 Corrective action by Test facility



Corrective action
- Major deviation (30 working days)
- Minor deviation (90 working days)

In case of follow up inspection corrective action will take within 30 working days

6 Approval inspection report by Lead inspector

7 Certificate by Director of BLQS

Category of test item contained in products

1) Pharmaceuticals
2) Pesticides
3) Cosmetic products
4) Veterinary drug products
5) Food additives
6) Feed additives
7) Industrial chemicals products
8) Others

Area of Expertise

1) Physical-chemical testing
2) Toxicity studies
3) Mutagenicity studies
4) Environmental toxicity studies on aquatic and terrestrial organisms
5) Studies on behavior in water, soil and air; bioaccumulation
6) Residue studies
7) Studies on effects on mesocosms and natural ecosystems
8) Analytical and clinical chemistry testing
9) Other studies

Bureau of Laboratory Quality Standards (BLQS)

Certificate of Compliance to OECD Principles of GLP

GLP number
Test Facility
Address
Category of test item contained in products
Area of Expertise
Date of first Inspection
Date of last Inspection
Registration Number

The above mention test facility has been inspected and found to be operating in compliance to OECD Principles of Good Laboratory Practice and BLQS Compliance Programme. The inspection of compliance is conducted at regular basis.

(.....)
Director
Bureau of Laboratory Quality Standards

Date of 1st Compliant
(first GLP compliant awarded)

Date of certificate issued:

After the last granting date, the test facility shall have at least two new completed studies in every year.

Bureau of Laboratory Quality Standards, Department of Medical Sciences
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