Good Laboratory Practice Compliance Monitoring Programme

What is the programme about?

The Singapore Good Laboratory Practice (GLP) Compliance Monitoring Programme recognises laboratory facilities that provide non-clinical safety testing in accordance with the Organisation for Economic Co-operation and Development (OECD) Principles of Good Laboratory Practice.

The programme ensures that a facility's processes and the conditions in which laboratory studies are planned, performed, monitored, recorded, and archived meet international requirements. It enables Singapore enterprises' research laboratories to gain acceptance for their environmental health and safety data in OECD countries.

Singapore has been a member of the OECD Mutual Acceptance of Data (MAD)[1] framework since 2010. Studies conducted in Singapore under GLP compliant conditions are accepted in more than 30 OECD and non-OECD member countries.

[1] *The principal tools for harmonization are a set of OECD Council Decisions which make up the OECD Mutual Acceptance of Data (MAD) system, including the OECD Guidelines for the Testing of Chemicals and OECD Principles of Good Laboratory Practice.*

Here are some of the GLP studies that a facility may seek registration on:

- Physical – Chemical Testing
- Toxicity Studies
- Mutagenicity Studies
- Environmental Toxicity Studies
- Studies on Behaviour in Water, Soil, Air, and Bioaccumulation
- Residual Studies
- Studies on the Effects of Mesocosms and Natural Ecosystems
- Analytical and Clinical Chemistry Testing

Benefits

- Enjoy greater market access and shorter time-to-market for your new products
- Increase the level of confidence in the integrity of your non-clinical study data and reports

Visit
www.sac-accreditation.gov.sg
to find out more.

Singapore Accreditation Council

Singapore Accreditation Council is managed by

Enterprise Singapore