The Organization for Economic Cooperation and Development (OECD) Principles of Good Laboratory Practice (GLP) is a quality system concerned with the organizational process and the conditions under which non-clinical health and environmental safety studies are planned, performed, monitored, recorded, archived and reported.

Nonclinical health and environmental safety studies covered by the principles of GLP include work conducted in the laboratory, in greenhouses, and in the field.

GLP principles are applied to the non-clinical safety testing of test items contained in pharmaceutical products, pesticide products, cosmetic products, veterinary drugs as well as food additives, feed additives and industrial chemicals.

The main duty of GLP monitoring agency is to monitor compliance with GLP Principles by conducting laboratory inspections and study audits. However, in many countries GLP-compliance is voluntary in nature.

ACCAB GLP Platinum Plus™ Accreditation is an independently delivered service requires that the Laboratory must conform to the most recent version of following standards/International Documents:

1. OECD Principles on Good Laboratory Practice.
2. Any additional requirements established by ACCAB from time to time.

Key Features & Advantages:

- May be useful to the Industries or laboratories who are dealing with the aforementioned chemicals;
- GLP embodies a set of principles that provides a framework within which laboratory studies are planned performed, monitored, reported and archived;
- GLP makes sure that the data submitted are a true reflection of the results that are obtained during the study;
- GLP makes sure that data is traceable;
- GLP promotes the quality and validity of test data used for determining the safety of chemicals and chemicals products;
- GLP Promotes international acceptance of tests.