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The Notification of the Food and Drug Administration

Practical Guidelines to submit Health Product for Safety Assessment and Toxicity Testing Procedures in the Preclinical Phase according to the OECD Principles of GLP

Some of health products including food additives, dietary supplement, new formula for used in cosmetic products, pharmaceutical products, veterinary drugs and active ingredients for hazardous substances used in household or public health, that may be required to go through toxicity test for safety assessment in the Preclinical Phase according to OECD GLP. Therefore, in order to allow the toxicity test and safety assessment comply with standard, it is required that the test facilities shall be carried out in compliance with the OECD GLP. In this regard, the Food and Drug Administration would like to request for applicants' cooperation to submit health products for toxicity test at test facilities inspected by the Bureau of Laboratory Quality Standards, Department of Medical Sciences, Ministry of Public Health, as OECD GLP Compliance Monitoring Authority.

This shall be in effort from now onwards.

Announced on 17 December B.E. 2558 (2015)

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