

(Translation)
(Official Seal)

Letter No. MOI 0714/31429

Thai Industrial Standards Institute
Rama 6 Road, Rajthevi
Bangkok 10400

29 August 2018

Re: Appointment of the National Compliance Monitoring Authority to the Department of Medical Sciences

To: Director General, Department of Medical Sciences,

Refer to: Letter of the Ministry of Public Health, No. MOPH 0621/1026 dated 22 March 2018

This letter refers to the letter of the Ministry of Public Health requesting the National Standardization Council to appoint the Bureau of Laboratory Quality Standards, Department of Medical Sciences, Ministry of Public Health to act as the National Compliance Monitoring Authority with its duty for inspecting and monitoring the compliance of the test facilities to the principles of good laboratory practice (GLP) according to the Organisation for Economic Cooperation Development (OECD).

The Thai Industrial Standards Institute would like to inform that the National Standardization Council in its meeting No. 8-1/2018 on 27 August 2018 had considered and approved the followings:

1. Appoint the Department of Medical Sciences as the responsible organization according to the Article 30 of the National Standardization Act, B.E. 2551 (A.D. 2008) to act as the National OECD GLP Compliance Monitoring Authority having the duty in inspecting and monitoring the compliance of the test facilities with OECD GLP in the scope of 1) pharmaceuticals, 2) pesticides, 3) cosmetic products, 4) veterinary drugs, 5) food additives, 6) feed additives, 7) industrial chemicals and others (to be added or amended).

2. The Department of Medical Sciences must operate according to the National Standardization Council Announcement on the Rules, Procedures and Conditions for Reporting by the Responsible Organization No. 2 that it must submit the annual operation report of the registered test facilities to the Thai Industrial Standards Institute which is in charge of the Council's Secretary within December of each year for reporting to the council.

3. The Department of Medical Sciences must report on its progress in the development of the plan to promote the competitiveness of the commercial products for the non-human safety testing according to the OECD GLP to the council.

Please be informed accordingly.

Respectfully yours,

Suthon Nikomketh

Signature

(Mr.Suthon Nikomketh)

Deputy Secretary-General of TISI

for Secretary-General of TISI