Republic of the Philippines
Department of Health
FOOD AND DRUG ADMINISTRATION

09 AUG 2021

FDA ADVISORY
No. 2019-525-A

TO: **ALL HEALTHCARE PROFESSIONALS AND THE GENERAL PUBLIC**

SUBJECT: **Lifting the Advisory of the Registered Medical Device under FDA Advisory No. 2019-525 entitled Public Health Warning Against the Purchase and Use of the Unregistered Medical Device "SURGITECH CLINICAL FIEBER THERMOMETER MT-101"**

The Food and Drug Administration (FDA) informs all healthcare professionals and the public that the medical device product "Surgitech Clinical Fieber Thermometer MT-101" has been registered by the Market Authorization Holder (MAH), Medasia Medical Product Corporation, with FDA registration number DVR-8795 in accordance to FDA rules and regulations.

Accordingly, the warning against the purchase and use of the product as mentioned in FDA Advisory No. 2019-525 is hereby lifted.

For more information and inquiries, kindly contact the FDA Center for Device Regulation, Radiation Health, and Research through e-mail at **cdrrhr@fda.gov.ph**, or call **(02) 8857-1900 loc. 8301.**

To report any sale or distribution of unregistered medical device products, the online reporting facility, **eReport** can be accessed at **www.fda.gov.ph/ereport.**

The issuance of this advisory shall not in any manner preclude this Office from issuing subsequent orders it may deem necessary and appropriate, should there be findings of any violation of the company to the existing laws, rules and regulation.

Dissemination of this advisory to all concerned is hereby requested.

ROLANDO ENRIQUE D. DOMINGO, MD
Director General

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