

## Republic of the Philippines Department of Health FOOD AND DRUG ADMINISTRATION



2 6 NOV 2021

FDA ADVISORY No. 2021-2704-A

TO:

ALL HEALTHCARE PROFESSIONALS AND THE

GENERAL PUBLIC

SUBJECT:

Lifting of the FDA Advisory No. 2021-2704 entitled "Public

Health Warning Against the Purchase and Use of the Unregistered Medical Device Product MEDICA® FLEXIBLE DIGITAL THERMOMETER WITH

FLEXIBLE TIP"

The Food and Drug Administration (FDA) informs all healthcare professionals and the general public that the medical device product, MEDICA® FLEXIBLE DIGITAL THERMOMETER WITH FLEXIBLE TIP, has been issued an FDA Certificate of Product Registration Variation Approval Letter to the Market Authorization Holder (MAH), Medical Center Trading Corporation, in accordance to existing FDA rules and regulations.

Accordingly, the warning against the purchase and use of the product as mentioned in FDA Advisory No. 2021-2704 dated 2 November 2021 is hereby lifted.

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 existing laws, rules, and regulations.

For more information and inquiries, kindly contact the FDA Center for Device Regulation, Radiation Health, and Research through e-mail at <a href="mailto:cdrrhr@fda.gov.ph">cdrrhr@fda.gov.ph</a> or call (02) 8857-1900 loc. 8301.

To report any sale or distribution of unnotified medical device, contact the online reporting facility, **eReport**, through e-mail at **ereport@fda.gov.ph**.

Dissemination of this advisory to all concerned is hereby requested.

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