

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



FDA ADVISORY No. 2019-208

19 6 JUL 2019

TO:

THE GENERAL PUBLIC

SUBJECT:

Public Health Warning Against the Purchase and Consumption of the Following Unregistered Food Supplements:

- 1. RAPIDCELL THERAPY Vitamin E
- 2. RAPIDCELL THERAPY Calcium & Vitamin D Strawberry Flavor
- 3. RAPIDCELL THERAPY Multi Vitamins for Kids
- 4. THYRONOL Dietary Supplement
- 5. RAPIDCELL THERAPY Magnesium Orange Flavor

The Food and Drug Administration (FDA) warns the public from purchasing and consuming the following unregistered food supplements:







Figure 1. Unregistered RAPIDCELL THERAPY Vitamin E





Figure 2. Unregistered RAPIDCELL THERAPY Calcium & Vitamin D Strawberry Flavor



Figure 3. Unregistered RAPIDCELL THERAPY Multi Vitamins for Kids



Figure 4. Unregistered THYRONOL Dietary Supplement



Figure 5. Unregistered RAPIDCELL THERAPY Magnesium Orange Flavor

The FDA verified through post-marketing surveillance that the abovementioned food supplements are not authorized and the Certificates of Product Registration (CPR) have not been issued. Pursuant to the Republic Act No. 9711, otherwise known as the "Food and Drug Administration Act of 2009", the manufacture, importation, exportation, sale, offering for sale, distribution, transfer, non-consumer use, promotion, advertising or sponsorship of health products without the proper authorization is prohibited.

Since these unregistered food supplements have not gone through evaluation process of the FDA, the agency cannot assure their quality and safety. The consumption of such violative products may pose health risks to consumers.

In light of the foregoing, the public is advised not to purchase the aforementioned violative products. Moreover, the public is advised to always check if a food supplement is registered with the FDA. The FDA website (www.fda.gov.ph) has a Search feature which may be used by typing in the name of the product before purchasing.

All concerned establishments are warned not to distribute violative food supplements until they have been issued the appropriate authorization, a License to Operate (LTO) for the establishment, and a CPR for the food supplement.

All FDA Regional Field Offices and Regulatory Enforcement Units, in coordination with law enforcement agencies and Local Government Units, are requested to ensure that violative products are not sold or made available in the market or areas of jurisdiction.

Kindly contact the FDA Center for Food Regulation and Research through e-mail at cfrr@fda.gov.ph, or call (02) 857-1900 loc. 8112 or 8105.

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

Dissemination of this advisory to all concerned is hereby requested.

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