

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



FDA ADVISORY No. 2016-131 F-1 NOV 2046

TO:

ALL HEALTHCARE PROFESSIONALS AND THE

GENERAL PUBLIC

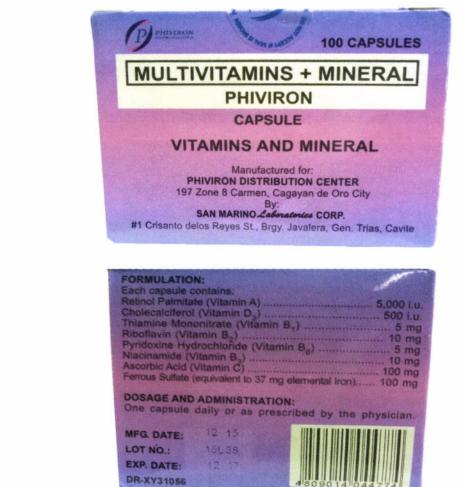
SUBJECT:

Public Health Warning Against the Use of the Unregistered

Drug Product "Multivitamins + Mineral (Phiviron)

Capsule"

The Food and Drug Administration (FDA) advises the public against the use of the unregistered drug product Multivitamins + Mineral (Phiviron) Capsule:



## Multivitamins + Minerals (Phiviron) Capsule

Manufactured for: PHIVIRON DISTRIBUTION CENTER - 197 Zone 8 Carmen, Cagayan

de Oro City

By: SAN MARINO Laboratories CORP. - #1 Crisanto delos Reyes St., Brgy. Javalera, Gen. Trias, Cavite

Figure 1. Box of the unregistered Multivitamins + Minerals (Phiviron) Capsule





Figure 2. Blister packs and capsule of unregistered Multivitamins + Minerals (Phiviron)
Capsule

Please be informed that the issued Certificate of Product Registration for the abovementioned product with Registration Number DR-XY31056 has already expired last 2010.

All healthcare professionals and the general public are hereby warned to be vigilant of the abovementioned unregistered drug product. This product poses potential danger or injury to the consuming public and the importation, selling or offering for sale of such is in direct violation of Republic Act No. 9711 or the Food and Drug Administration Act of 2009.

In the interest of protecting public health and safety, the Officers of the Field Regulatory Operations Office (FROO) are hereby mandated to exercise their regulatory function pursuant to Section 14 of Republic Act. No. 9711.

All establishments and outlets are hereby warned against selling and/or dispensing the unregistered product identified above. Anyone found selling the said product will be penalized.

Likewise, all local government units and law enforcement agencies are requested to ensure that this unregistered product is not sold or offered for sale in their localities or area of jurisdiction.

Consumers are advised to purchase their medications only from FDA-licensed establishments. In addition to inspection of establishments, product evaluation, registration, and testing are measures that the government undertakes to ensure the quality, safety, and efficacy of health products. Please look for the FDA Registration number on the product label. Be reminded that drug products registered with FDA Philippines bear in their labels information in English and/or in Filipino for all consumers to understand.

For more information and inquiries, please e-mail us at <a href="mailto:info@fda.gov.ph">info@fda.gov.ph</a>. To report continuous sale or distribution of unregistered health products, kindly e-mail us via <a href="mailto:report@fda.gov.ph">report@fda.gov.ph</a> or you may call telephone number (02)807-8275. For any suspected adverse drug reaction (ADR), report immediately to FDA through this link: <a href="https://www.fda.gov.ph/adr-report-new">www.fda.gov.ph/adr-report-new</a> and fill out all the required fields.

Dissemination of the information to all concerned is requested.

NELA CHARAPE G. PUNO, RPh Director General

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