



FDA ADVISORY
No. 2022-1748

04 OCT 2022

TO : ALL HEALTHCARE PROFESSIONALS AND THE GENERAL PUBLIC

SUBJECT : Public Health Warning Against the Purchase and Use of the Unregistered Drug Product "MD® Skin Fighter 8 g"

The Food and Drug Administration (FDA) advises the public against the purchase and use of the unregistered drug product:

MD® Skin fighter
Net content: 8g

SKIN FIGHTER

- Using range: hand / feet ringworm, Body moss, Psoriasis, Moss, Mosses, eczema, Scabies, Blisters, Diaper rash, Skin itching, Ichthyosis, Scleroderma, Mosquito bites, Balanitis, Genital itching, Hemorrhoids, The skin problems caused by Staphylococcus aureus, Escherichia coli and Candida albicans
- Ingredient: Sophora flavescens 0.05%, Stemonia tubrosa Lour. 0.05%, Cnidium monnieri(L.) Cuss. 0.05%, Latin Tille. 0.05%, Mentha haplo-calyx Brip. 0.05%, Chlorhexidin* Acetas 0.138%(w/w).
- Usage: topical, take this product covered with hand, foot, body skin, three times a day.
- Note: Topical, not for the eyes and oral.
- Hong Kong Huayuan Pharmaceutical Development Co., Ltd. Producer
Newport Centre, 30 Canton Road, Tsim Sha Tsui, Yau Tsim Mong District, Hong Kong Special Administrative Region.

MFG. 20220103
EXP. 20270102

MD® Skin Fighter 8 g
Manufactured by: Hong Kong Pharmaceutical Development Co., Ltd. - Newport Centre, 30 Canton Road, Tsim Sha Tsui, Yau Tsim Mong District, Hong Kong Special Administrative Region

Figure 1: Unregistered drug product



FDA Post-Marketing Surveillance (PMS) activities have verified that the abovementioned drug product has not gone through the registration process of the Agency and has not been issued with proper authorization in the form of Certificate of Product Registration. Thus, the Agency cannot guarantee its quality and safety. Therefore, consumption of such violative product may pose potential danger or injury to health.

Pursuant to 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, promotion, advertising or sponsorship of health products without proper authorization from FDA is prohibited.

All concerned establishments and/or entities are warned not to distribute the above-identified violative drug product until it has been covered by the appropriate authorization, otherwise, regulatory actions and sanctions shall be strictly pursued. Always check if a product is registered with the FDA by using the **FDA Verification Portal** feature accessible at <https://verification.fda.gov.ph>.

All Local Government Units (LGUs) and Law Enforcement Agencies (LEAs) are requested to ensure that this product is not sold or made available in their localities or areas of jurisdiction.

For more information and inquiries, please e-mail us at info@fda.gov.ph. To report continuous sale or distribution of unregistered health products, kindly e-mail us via ereport@fda.gov.ph. You may also call the Center for Drug Regulation and Research at telephone number (02) 8809-5596. For any suspected adverse drug reaction (ADR), report immediately to FDA through this link: <https://primaryreporting.who-umc.org/Reporting/Reporter?OrganizationID=PH> and fill out all the required fields.

Dissemination of the information to all concerned is requested.


DR. SAMUEL A. ZACATE
Director General

DTN:



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