



Republic of the Philippines
Department of Health
FOOD AND DRUG ADMINISTRATION



FDA ADVISORY
No. **2022-1899**

07 DEC 2022

TO : ALL HEALTHCARE PROFESSIONALS AND THE GENERAL PUBLIC

SUBJECT : Public Health Warning Against the Purchase and Use of the Following Unregistered Drug Products:

1. **Natural Green Leaf® Max Strength Gel Patch 1 Patch 8 x 12 cm**
2. **Natural Green Leaf® Quick Action Vapor Inhaler [Primary Packaging] Natural Green Leaf® Vapor Inhaler [Secondary Packaging]**
3. **Natural Green Leaf® 2-in-1 Nasal Inhaler [Orange Breeze] [Primary Packaging] Natural Green Leaf® 2 Ways Inhaler (Inhale and Apply) [Secondary Packaging]**

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

Natural Green Leaf® Max Strength Gel Patch 1 Patch 8 x 12 cm
 Manufacturer: ShanDong Pharmaceuticals Co., Ltd – Lashan Industrial Park, Shizong District, Jinan 250000
 Imported by: Cha-Ken Consumer Goods Trading – 602A, 611 Tytana Bldg. Tytana St, Plaza Lorenzo Ruiz, Binondo, Manila

Figure 1. Unregistered drug product





Figure 2. Unregistered drug product



Figure 3. Unregistered drug product

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

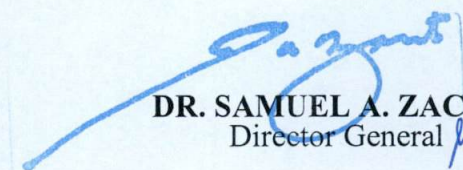
Pursuant to Republic Act No. 9711, otherwise known as the "Food and Drug Administration Act of 2009", the manufacture, importation, exportation, sale, offering for sale, distribution, transfer, promotion, advertising or sponsorship of health products without proper authorization from FDA is prohibited.

All concerned establishments and/or entities are warned not to distribute the above-identified violative drug products until they have already been covered by the appropriate authorization, otherwise, regulatory actions and sanctions shall be strictly pursued. Always check if the products are 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 these products are 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

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