Republic of the Philippines
Department of Health
FOOD AND DRUG ADMINISTRATION

FDA ADVISORY

No. **2021-0909**

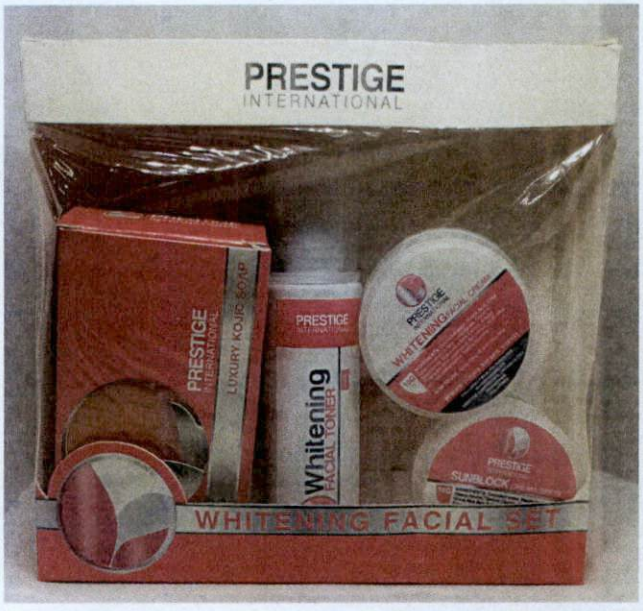
29 APR 2021

TO: **GENERAL CONSUMING PUBLIC**

SUBJECT: **Public Health Warning Against the Purchase and Use of the Non-Compliant Cosmetic Product " PRESTIGE INTERNATIONAL WHITENING FACIAL SET "**

1. **PRESTIGE INTERNATIONAL WHITENING FACIAL CREAM**
2. **PRESTIGE INTERNATIONAL LUXURY KOJIC SOAP**
3. **PRESTIGE INTERNATIONAL WHITENING FACIAL TONER**
4. **PRESTIGE INTERNATIONAL SUNBLOCK CREAM SPF 15**

The Food and Drug Administration (FDA) warns the general public from purchasing and using the non-compliant cosmetic product **PRESTIGE INTERNATIONAL WHITENING FACIAL SET** with details specified below:

PRODUCT DETAILS	PRODUCT IMAGE/S
Local Company Responsible for Placing the Product in the Market: **MANNIX CARANCHO PRESTIGE CORPORATION** Address: **Alexander St. Corner Rosario St. Poblacion, Urdaneta, Pangasinan, Ilocos (Region I), 2428, Alexander St. Corner Rosario St. Poblacion, Urdaneta, Urdaneta, Pangasinan, Ilocos (Region I), 2428, Urdaneta, Pangasinan** Country of Manufacture: **Philippines** Lot/Batch Number: **Not Indicated** Manufacturing Date: **01282019** Expiration Date: **01282021**	

NON-COMPLIANCE/S
1.The ingredients listed on the label are inconsistent with the information declared in the acknowledged product notification. 2.Three components of the kit are not notified as a combination product 3. Failure to declare on the label the manufacturer's batch number and the manufacturing or the expiry date

The Food and Drug Administration has verified that the abovementioned product is **NON-COMPLIANT** through its postmarketing surveillance (PMS) pursuant to Book I, Article II, Section 2 of the Rules and Regulations Implementing Republic Act No. 9711, otherwise known as the "Food

and Drug Administration Act of 2009" which provides for the relevant functions, powers and duties of the agency, including the conduct of PMS activities in the monitoring of health products.

Based on the Certificate of Product Notification issued to the company, any subsequent changes to the information previously submitted to the FDA will render the notification invalid.

In light of the foregoing, the public is advised not to purchase the aforementioned violative cosmetic product. Always check if a product is notified with the FDA by using the FDA Verification Portal feature accessible at https://verification.fda.gov.ph. which may be used by typing in the name of the product before the purchase and/or using the cosmetic products.

All concerned establishments are warned not to distribute violative cosmetic product until they have fully complied with the rules and regulation of the FDA.

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 their jurisdiction.

To report any sale, distribution, complaint and/or adverse event on the use of the violative cosmetic product, the online reporting facility, **eReport** can be accessed at **ereport@fda.gov.ph,** or call us at the Center for Cosmetics and Household/Urban Hazardous Substances Regulation and Research (CCHUHSRR) hotline **(02) 8857-1900 loc. 8113 or 8107.**

Dissemination of this advisory to all concerned is hereby requested.

ROLANDO ENRIQUE D. DOMINGO, MD
Director General

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