



FDA ADVISORY  
No. 2021-2033

170 AUG 2021

TO: ALL HEALTHCARE PROFESSIONALS AND THE GENERAL PUBLIC

SUBJECT: Public Health Warning Against the Purchase and Use of the following Unnotified Medical Device Products:

1. XIAO RU DISPOSABLE FACE MASK
2. HENG DE KN95 PM2.5 PARTICULATE RESPIRATOR
3. DISPOSABLE FACE MASK
4. BO HUI DISPOSABLE DAILY PROTECTIVE FACE MASK

The Food and Drug Administration (FDA) warns all healthcare professionals and the general public **NOT TO PURCHASE AND USE** the unnotified medical device products:



Figure 1. Unnotified Xiao Ru Disposable Face Mask





Figure 2. Unnotified Heng De KN95 PM2.5 Particulate Respirator

3-ply Surgical Facemask (FDA APPROVED)

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Figure 3. Unnotified Disposable Face Mask



Figure 4. Unnotified Bo Hui Disposable Protective Mask

The FDA verified through post-marketing surveillance that the above mentioned medical device products are not notified and no corresponding Product Notification Certificates have 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 unnotified medical device products have not gone through evaluation process of the FDA, the agency cannot assure their quality and safety.

All concerned establishments are warned not to distribute, advertise, or sell the said violative medical device products until the Product Notification Certificates are issued, otherwise, regulatory actions and sanctions shall be strictly pursued. Always check if a product has been notified with the FDA before purchasing it by making use of the embedded Search feature of the FDA website accessible at [www.fda.gov.ph](http://www.fda.gov.ph). You may also look for the FDA Notification number on the product label in the form of CMDN-xxx.

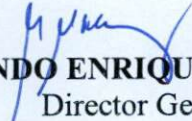
All Law Enforcement Agencies (LEAs) and Local Government Units (LGUs) are requested to ensure that these products are not sold or made available in the market or areas of jurisdiction.

The Bureau of Customs is urged to restrain the entry of these unnotified products.

For more information and inquiries about this advisory, kindly contact the FDA CDRRHR through e-mail at [cdrhr@fda.gov.ph](mailto:cdrhr@fda.gov.ph) indicating on the subject the concerned Advisory, or call (02) 8857-1900 loc. 8301.

To report any sale or distribution of unnotified medical device, the online reporting facility, eReport can be accessed at [www.fda.gov.ph/ereport](http://www.fda.gov.ph/ereport).

Dissemination of this advisory to all concerned is hereby requested.

  
**ROLANDO ENRIQUE D. DOMINGO, MD**  
Director General

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