



Republic of the Philippines
Department of Health
FOOD AND DRUG ADMINISTRATION



FDA ADVISORY
No. **2015-054**

30 JUL 2015

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

1. **Kidney Guard Capsule**
2. **Cordyceps Lung Roborant Capsule**
3. **Prostate Gland Pills (Specific Kai Kit Pill)**
4. **Pollen Qian Lie Tong Tablet**
5. **Tong Lin Huashidan (Urinary Organs Salve) Pills**
6. **Butiao Tablets**
7. **Cow's Head Brand Tung Shueh Pills**
8. **Gejie Da Bu Wan Pills**
9. **Skinhelp Cream**
10. **Bi Yan Pian Tablets**
11. **Snake's Bone Anti Rheumatism Capsule**
12. **She Wang Bui Capsule**
13. **Clindamycin Hydrochloride (Eurocin-A Gel 1%)**

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

 <p>KIDNEY GUARD CAPSULE Marketing Authorization Holder: D.K. Traditional Medicines & Natural Health Products Limited, Hong Kong</p>	 <p>CORDYCEPS LUNG ROBORANT CAPSULE Marketing Authorization Holder: D.K. Traditional Medicines & Natural Health Products Limited, Hong Kong</p>
 <p>PROSTATE GLAND PILLS (SPECIFIC KAI KIT PILL) Marketing Authorization Holder: Kwangchow Pharmaceutical Industry Co., Kwangchow, China</p>	 <p>POLLEN QIAN LIE TONG TABLET Marketing Authorization Holder: D.K. Traditional Medicines & Natural Health Products Limited, Hong Kong</p>





TONG LIN HUASHIDAN (URINARY ORGANS SALVE) PILLS

Marketing Authorization Holder: D.K. Traditional Medicines & Natural Health Products Limited, Hong Kong



BUTIAO TABLETS

Marketing Authorization Holder: D.K. Traditional Medicines & Natural Health Products Limited, Hong Kong



COW'S HEAD BRAND TUNG SHUEH PILLS

Marketing Authorization Holder: Ta Ang Pharmaceutical Company, Hong Kong



GEJIE DA BU WAN PILLS

Export by: China Huikang Medicines & Health Product Imp. & Exp. Co., Ltd.



SKINHELP CREAM

Marketing Authorization Holder: Europharmax (Japan) Int'l Ltd.



BI YAN PIAN TABLETS

Marketing Authorization Holder: WuHan ZhongLian Pharmaceutical Group Co. Ltd. Works, Hankow China

 <p>SNAKE'S BONE ANTI RHEUMATISM CAPSULE Marketing Authorization Holder: Zhong Hua Medicines Biology Institute, China</p>	 <p>SHE WANG BUI CAPSULE Marketing Authorization Holder: Zhong Hua Medicines Biology Institute, China</p>
	<p>CLINDAMYCIN HYDROCHLORIDE (EUROCIN-A GEL 1%) Manufactured by Europharm Laboratories Co., Ltd. – 1 2-14 Dai Wang Street, Tai Po Industrial Estate, Hong Kong</p>

Table 1. Unregistered Drug Products

The abovementioned drug products pose 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.

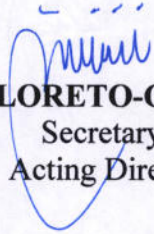
In the interest of protecting public health and safety, and pursuant to Section 12 of R.A. No. 9711, the Field Regulatory Operations Office (FROO) Officers are hereby ordered to seize the aforementioned unregistered drug products found in the market.

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

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

All consumers are advised to purchase their medications only from FDA-licensed establishments. Please note that, 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. Moreover, please 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 email us at info@fda.gov.ph. To report continuous sale or distribution of unregistered health products, kindly email us via report@fda.gov.ph or you may call the telephone number (02)807-8275. For any suspected adverse drug reaction (ADR), report immediately to FDA through this link: www.fda.gov.ph/adr-report-new and fill out all the required fields.



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¹ Pursuant to DPO 2015-1845