

Republic of the Philippines Department of Health FOOD AND DRUG ADMINISTRATION



09 June 2015

FDA ADVISORY
No. 2015 - 035

SUBJECT: <u>Public Health Warning Against the Use of the Following</u>
Unregistered Drug Products:

- 1. Erythromycin Eye Ointment, Marketing Authorization Holder: In Foreign Language
- 2. Liver Aid Sugar-Coated Tablet
- 3. Butiao Tablets
- 4. Anticancerlin 0.25 g Film-Coated Tablet
- 5. Monophosphoric Ester B1 Vitamin + Pyridoxine HCl + Cyanocobalamin + Indomethacin (Bevidan One + Six + Twelve) 300 mg/ 200 mg/ 250 mcg/ 25 mg Capsule
- 6. Pulmonary Tonic Tablets 500 mg
- 7. I-Ching-Sung Laxative Tablets 5 mg
- 8. Snake Bone Capsules
- 9. UA-Block Uric Acid Block Capsule
- 10. Kidney Wash Capsule

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



ERYTHROMYCIN EYE OINTMENT

Marketing Authorization Holder: In Foreign Language









LIVER AID SUGAR-COATED TABLET

Marketing Authorization Holder: China National Medicines & Health Products Import and Export Co.-Heilongjiang Branch





BUTIAO TABLETS

Prepared by The United Pharmaceutical Manufactory - Kwangchow, China



ANTICANCERLIN 0.25 g FILM-COATED TABLET

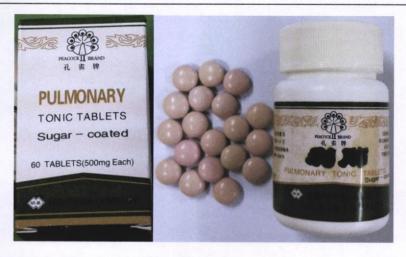
Manufactured by Shanghai Sine Jiahua Pharmaceutical Co., Ltd.

Sole Distributor: Jin Ling Enterprises



MONOPHOSPHORIC ESTER B1 VITAMIN + PYRIDOXINE HCl + CYANOCOBALAMIN + INDOMETHACIN (BEVIDAN ONE + SIX + TWELVE) 300 mg/ 200 mg/ 250 mcg/ 25 mg CAPSULE

Sole Agent for Asia: Wah-Tai Co. - Macau



PULMONARY TONIC TABLETS 500 mg

Marketing Authorization Holder: Ming Wah Company

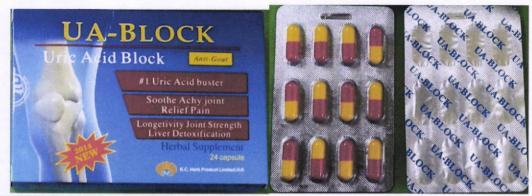


I-CHING-SUNG LAXATIVE TABLETS 5 mg Manufactured by Jintan Pharmaceutical Factory – Jiangsu China



SNAKE BONE CAPSULES

Marketing Authorization Holder: K.C. Herb Product Ltd. H.k.



UA-BLOCK URIC ACID BLOCK CAPSULE

Marketing Authorization Holder: K.C. Herb Product Limited. H.K.



KIDNEY WASH CAPSULE

Manufactured by: D.K. Traditional Medicines & Natural Health Products Limited - Hongkong

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 of this Office 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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DTN:20150521093549

Pursuant to DPO 2015-1845