Republic of the Philippines
Department of Health
FOOD AND DRUG ADMINISTRATION

FDA ADVISORY
No. 2017-235

04 AUG 2017

TO: **GENERAL CONSUMING PUBLIC AND OTHER CONCERNED STAKEHOLDERS AND PARTIES**

SUBJECT: **Cancellation of the Certificate of Product Registration for Baoma® Black Mosquito Coil with FDA Registration No. HSR-7656**

The Food and Drug Administration (FDA) advises the public that the Certificate of Product Registration for the product **Baoma® Black Mosquito Coil** with **FDA Registration No. HSR-7656** issued to Marketing Authorization Holder (MAH) **Targetline Marketing Corp.** located at No. 349 Mac Arthur Highway, Malinta, Valenzuela City, is hereby **CANCELLED**.

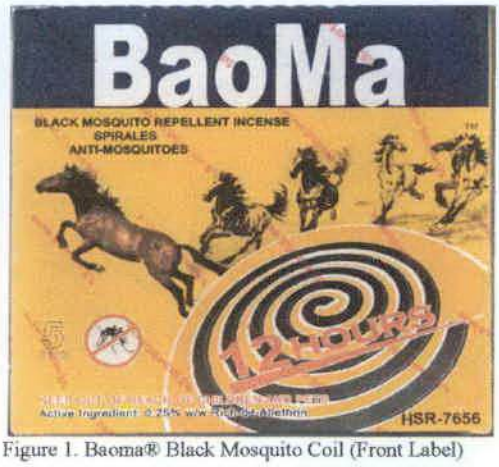

Figure 1. Baoma® Black Mosquito Coil (Front Label)

Figure 2. Baoma® Black Mosquito Coil (Back Label)

After a judicious review of the records of the administrative case and the evidence presented, this Office rules against MAH and respondent **Targetline Marketing Corp.** for violation of Sec. V (A) (a.3), AO 2014-0038, in relation to Sec. 11 (a), (b) and (j) of RA 3720, as amended by RA 9711, for importing, distributing, selling and offering for sale of adulterated, misbranded and unregistered Household/Urban Hazardous Substances.

In light of the above, the public is advised not to purchase and use the aforementioned violative product. The safety, efficacy and quality of any health product that is adulterated, misbranded, and/or unregistered cannot be guaranteed. The use and exposure to such products may pose imminent danger to human and animal health.

All concerned establishments are warned not to distribute the above-identified violative product. Non-compliance thereto shall warrant regulatory actions and sanctions to be strictly pursued.

All Local Government Units (LGUs) and Law Enforcement Agencies (LEAs) are requested to ensure that this product is 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 the above violative product, utilize our online reporting facility, **eReport**, at **www.fda.gov.ph/ereport**, or e-mail us via **report@fda.gov.ph**, or call us at the Center for Cosmetics Regulation and Research (CCRR) hotline **(02) 857-1979 or (02) 857-1984**.

Dissemination of the information to all concerned is requested.

NELA CHARADE G. PUNO, RPh.
Director General

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