

Republic of the Philippines Department of Health FOOD AND DRUG ADMINISTRATION



FDA ADVISORY No. 2023-0132

1:7 JAN 2023

TO

: ALL HEALTHCARE PROFESSIONALS AND THE GENERAL

PUBLIC

SUBJECT:

Public Health Warning Against the Purchase and Use of the

Following Unregistered Veterinary Drug Products:

1. PaWell Veterinarian's Choice OB Supplement with Probiotics 120mL [Vanilla Flavor]

2. PaWell Veterinarian's Choice Multivitamins + Minerals + Lysine with Electrolytes 120 mL [Vanilla Flavor]

3. PaWell Veterinarian's Choice Sodium Ascorbate + Calcium Ascorbate + Zinc (Calzinc) 120 mL [Vanilla Flavor]

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



PaWell Veterinarian's Choice OB Supplement with Probiotics 120mL [Vanilla Flavor]

Distributed by: Pets J Veterinary Products Inc. – Unit E, Ground Floor, Westgate Tower, Investment Drive Madrigal Park 1, Brgy. Ayala Alabang Muntinlupa City 1780

Figure 1. Unregistered veterinary drug product

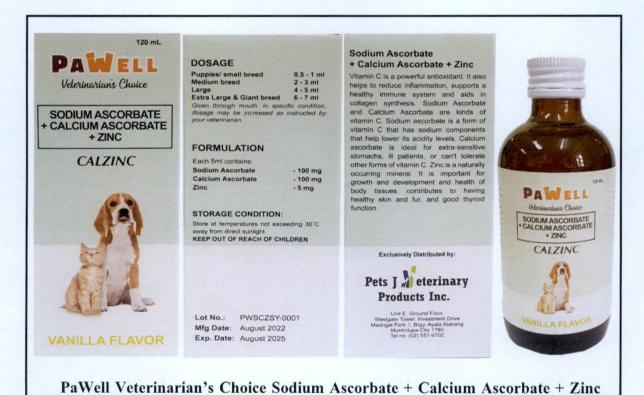




PaWell Veterinarian's Choice Multivitamins + Minerals + Lysine with Electrolytes 120 mL [Vanilla Flavor]

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Figure 2. Unregistered drug product



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Figure 3. Unregistered drug product

(Calzine) 120 mL [Vanilla Flavor]

FDA Post-Marketing Surveillance (PMS) activities have verified that the abovementioned drug products have not gone through the registration process of the Agency and have not been issued with proper authorization in the form of Certificate of Product Registration. Thus, the Agency cannot guarantee their quality, safety and efficacy. Therefore, consumption of such violative products may pose potential danger or injury to health.

Pursuant to 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, promotion, advertising or sponsorship of health products without proper authorization from FDA is prohibited.

All concerned establishments and/or entities are warned not to distribute the above-identified violative drug products until they have already been covered by the appropriate authorization, otherwise, regulatory actions and sanctions shall be strictly pursued. Always check if the products are registered with the FDA by using the FDA Verification Portal feature accessible at https://verification.fda.gov.ph.

All Local Government Units (LGUs) and Law Enforcement Agencies (LEAs) are requested to ensure that these products are 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 unregistered health products, kindly e-mail us via ereport@fda.gov.ph. You may also call the Center for Drug Regulation and Research at telephone number (02) 8809-5596. For any suspected adverse drug reaction (ADR), report immediately to FDA through this link: https://primaryreporting.who-umc.org/Reporting/Reporter?OrganizationID=PH and fill out all the required fields.

Dissemination of the information to all concerned is requested.

Director General

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