Republic of the Philippines
Department of Health
FOOD AND DRUG ADMINISTRATION
Civic Drive, Filinvest Corporate City, Alabang, Muntinlupa City

LICENSE TO OPERATE

as

Drug Manufacturer

is hereby granted to

AM-EUROPHARMA CORPORATION

Barangay Saimsim, Calamba, Laguna

Owner:	**AM-EUROPHARMA CORPORATION**
License Number:	**LTO-3000003151687**
Application Type:	**Renewal**
Validity of License:	**09 January 2025**

This LTO shall be renewed within **90 days** before its expiration, upon compliance with relevant laws, rules and regulations and the payment of fees. This LTO may be suspended, cancelled or revoked by this Office for cause if found violating RA 9711 and related issuances.

Furthermore, with this LTO, the FDA allows the establishment to apply for a market authorization [i.e. registration (CPR) or notification] for health products prior to manufacture, importation, sale or offer for sale, distribution, transfer, advertisement and/or promotion as the case may be.

BY AUTHORITY OF THE DIRECTOR GENERAL

JESUSA JOYCE N. CIRUNAY, RPh
Director IV
Center for Drug Regulation and Research

DISPLAY IN PUBLIC VIEW

Additional information required under FDA Circular 2016-006 are reflected at the second page of this LTO

This electronic-LTO (eLTO) is computer generated and does not require signature.
Verify at http://www.fda.gov.ph/industry-corner/e-lto/560767-LTO-3000003151687

FDA-20212B3IWZIJQI29A7IX4C57

Republic of the Philippines
Department of Health
FOOD AND DRUG ADMINISTRATION

Food and Drug Administration PHILIPPINES

CERTIFICATE OF CURRENT GOOD MANUFACTURING PRACTICE

This is to certify that

NAME	:	**AM-EUROPHARMA CORPORATION**
PLANT ADDRESS	:	**Barangay Saimsim, Calamba, Laguna**
OWNER	:	**AM-EUROPHARMA CORPORATION**
AUTHORIZEDPERSON FOR PRODUCT BATCH RELEASE	:	**Edna O. Saur**
QUALITY ASSURANCE	:	**Ma Zaida De Jesus Benavidez**
QUALITY CONTROL	:	**Kevin Bossie Sanchez Davis**
PRODUCTION	:	**Celerina V. Samonte**
PHARMACIST	:	**Olivia Y. Licuanan**
NAME OF PRODUCTS	:	**Non-Penicillin (Capsules, Liquid, Drops, Powder for Suspension, Syrup & Tablet); Veterinary products - Non-Penicillin (Tablet & Capsule)**

with a valid License to Operate bearing number **LTO-3000003151687** as drug manufacturer has complied to the requirements of **Administrative Order No. 2020-0017 or the Revised Guidelines on the Unified Licensing Requirements and Procedures** and to **Administrative Order No. 2012-0008 s. 2012 or the Adoption and Implementation of the Pharmaceutical Inspection Cooperation Scheme (PIC/S) Guides for the Good Manufacturing Practice (GMP) for Medicinal Products** for the manufacture of the above product.

This certification is **valid until 09 January 2025**. Notwithstanding this certification, the above establishment shall be subject to inspection at anytime to validate its continuous compliance with relevant laws, rules and regulations. Any violation thereof, this Office reserves the right to suspend, cancel or revoke this certification.

Issued this 9th day of November 2021 at Alabang, Muntinlupa City, Philippines.

BY AUTHORITY OF THE DIRECTOR GENERAL
Per FDA Order No. 2016-005

JESUSA JOYCE N. CIRUNAY, RPh
Director IV
Center for Drug Regulation and Research

Certificate No.: DM-2021-124
O.R # Ref# 1932-11022021-981051
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