



## CERTIFICATE OF GMP COMPLIANCE

This is to certify that the foreign manufacturer:

Name : **RUSAN PHARMA LIMITED**

Plant Address : **KHASRA NO. 122 MI, CENTRAL HOPE TOWN,  
SELAQUI, DISTRICT DEHRADUN, 248197,  
UTTARAKHAND, INDIA**

engaged in the manufacture of the following pharmaceutical dosage form/s:

- I. Human
  1. Non-sterile\*
    - 1.1. General
      - 1.1.1. Liquids for internal use
      - 1.1.2. Tablets

*\*Primary and Secondary packaging, and Quality control testing: Chemical/Physical and Microbiological*

made available in the Philippines through **AGLOBAL CARE, INC.**, with valid License to Operate No. CDRR-NCR-DI/W-101728, has complied with the requirements of **current Good Manufacturing Practice (cGMP)** after **desktop evaluation of the submitted GMP documentary evidence**, consistent with Administrative Order No. 2012-008, "Adoption and Implementation of the Pharmaceutical Inspection Cooperation Scheme (PIC/S) Guides for the Good Manufacturing Practice (GMP) for Medicinal Products" and Administrative Order No. 2013-0022, "Guidelines for Current Good Manufacturing Practice (cGMP) Clearance and inspection of Foreign Drug Manufacturers."

This Certificate is valid until **18 April 2026**. Notwithstanding this certification, the foreign manufacturer shall be subject to inspection at any time to validate its continuous compliance with relevant FDA laws, rules, and regulations. Any violation thereof, this Office reserves the right to suspend, cancel or revoke this certificate.

Issued this 14 November 2023 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 & Research

CERTIFICATE NO. : CDRR-CGMP-5165  
REG. STATUS : INITIAL  
TRACK NUMBER : 20230606092316  
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