



GOVERNMENT OF THE PEOPLE'S REPUBLIC OF BANGLADESH

MINISTRY OF HEALTH & FAMILY WELFARE
DIRECTORATE GENERAL OF DRUG ADMINISTRATION
OUSHAD BHABAN, MOHAKHALI
DHAKA-1212, BANGLADESH
www.dgda.gov.bd



CERTIFICATE OF GOOD MANUFACTURING PRACTICE (GMP)
FOR PHARMACEUTICAL (PRODUCT(S))

This certificate conforms to the format recommended by the World Health Organization (WHO)

Certificate Number: DGDA/6-28/2025/9759

Date: 13 JUL 2025

It is hereby certified that M/S. Zhaofeng Gelatin Limited a drug (Pharmaceutical Product) manufacturing and marketing organization, has been given license to manufacture and sell its product freely in the People's Republic of Bangladesh as lawfully required and granted in pursuance of the provisions Chapter-Vii, of the Drug & Cosmetics Act, 2023 (xxix of 2023) or any rule made there under.

On the basis of the inspection carried out on 23.08.2024 & 27.11.2024 we certify that the site indicated on this certificate complies with Good Manufacturing Practices for the dosage forms, categories and activities listed in Table 1.

1. Name & address of site : M/S Zhaofeng Gelatin Limited
: Sutipara, Kalampur-1351, Dhamrai, Dhaka, Bangladesh.

2. Manufacturer's License No : Non Biological License: 537; Date of Issue: 27.07.2022

3. Table: 1

Product(s)	Category(ies)	Activity (ies)
Empty Hard Gelatin Capsules Shell (Manufactured from Animal Bone Gelatin)	Capsule Shell	• Sourcing & Procurement of RM/PM • Formulation design and development • Dispensing of RM/PM • Production • Packaging • Quality Assurance: - Manufacturing Environment Control - In-process Quality Control - Finished Product Analysis - Batch & Lot Release

Certificate of Good Manufacturing Practices (GMP) for Pharmaceutical Product(s) Continued to page-2



The responsibility for ensuring the quality of the individual batches of the pharmaceutical products manufactured through this process lies with the manufacturer.

The manufacturing plant in which the pharmaceutical products are produced is subject to inspection at suitable intervals.

The manufacturer conforms to the requirements for good practices in the manufacture and quality control (GMP) of drugs, as required under law of this country, as well as recommended by the World Health Organization (WHO) in respect of pharmaceutical products to be manufactured, sold or distributed within the country of origin or to be exported.

This Certificate remains valid for a period of 2 (Two) years from the date of issue. It becomes invalid if the activities and/or categories certified, herewith are changed or if the site is no longer considered to be in compliance with GMP.

Name of Authorized Person

: Major General Md. Shameem Haidar

Address of the certifying Authority

: Directorate General of Drug Administration.
Aushad Bhaban, Mohakhali, Dhaka-1212

Telephone

: +880-02-2222-80803

Fax No

: +880-02-9880854

E-mail

: dgda.gov@gmail.com

Web-site

: www.dgda.gov.bd

Stamp and Date:



Major General Md. Shameem Haidar

Director General 09 JUL 2025

Directorate General of Drug Administration

&

Licensing Authority (Drugs)

Government of the Peoples Republic of Bangladesh