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/6-36/2026/ 6770 Date: 27 /04/2026

It is hereby certified that **N H Laboratories (Unani),** a unani drug and marketing organization, has been given license to manufacture and sell its products freely in the People's Republic of Bangladesh, as lawfully required and granted in pursuance of the provisions of chapter VII of The Drug & Cosmetics Act, 2023 (Act No. XXIX of 2023) and the rules made there under:

On the basis of the inspection carried out on 10.10.2024 & 18.02.2026 is hereby confirmed that the manufacturing facility specified below has been found to be in compliance with the principles of Good Manufacturing Practices (GMP), as recommended by the World Health Organization (WHO), for the dosage forms, product categories, and manufacturing activities listed in Table 1 of this certificate.

The Lincese information are as below:

1. Name and Address of site : **N H Laboratories (Unani)**
   39/C, BSCIC I/A, Gotatikor, Kadamtoli, Sylhet, Bangladesh.

2. Drug Manufacturing License No. : U-216 Date of Issue: 05.01.1988

3. Table-1

Dosage Form (s)	***Category***	***Activity (ies)***
Capsule	Antifungal, Antioxidant, Antidepressant, Analgesics, Hepatoprotective, General tonic, Memory enhancer, Immunity enhancer, Probiotics, Dietary supplements, Stimulant, Anti hypertensive, Antiulcer, Anti diabetic, Anti piles	• Procurement of raw materials from approved sources/vendors • Quarantine of raw materials in warehouse • Sampling and testing of raw materials/packaging materials • Dispensing of raw material/packaging materials • Control of manufacturing environment • In-process control of intermediate, bulk and finished products • Control of packaging and labeling operations • QC & QA of incoming materials and finished products • Storage of finished products • Documentation and archiving • Compliance of safety, health and environments • Change control procedure • Deviations and OOS handling • Complain handling • Training • Self-inspection/Internal auditing • Process validation • Annual Product Quality Review (APQR)
Ointment/Syrup/Semi-Solid	Topical antifungal, Topical antiprotozoal, Analgesics, Cough expectorant, Supplements, Blood purifier, Warm carminative, Digestive, Antidiarrheal, Liver tonic, Stimulant, Blood alkaly, Antiulcer	

The responsibility for the quality of each batch of the Unani (Medicine) manufactured by this process lies with the manufacturer. The manufacturing site at which the pharmaceutical product(s) are produced is subject to inspection by the competent regulatory authority at suitable intervals.

The manufacturer complies with the requirements of Good Manufacturing Practices (GMP) in respect of the manufacture and quality control of pharmaceutical products, as laid down in the applicable national legislation of the country of origin, and as recommended by the World Health Organization (WHO).

This certificate is valid for a period of 2 (two) years from the date of issue, unless suspended or withdrawn earlier. It shall become invalid in the event of any change to the certified activities and/or product categories, or if the manufacturing site is no longer considered to be in compliance with GMP requirements.

Name of the authorized person	:	**Major General Md. Shameem Haidar**
Address of Certifying Authority	:	Directorate General of Drug Administration Aushadh Bhaban, Mohakhali, Dhaka-1212, Bangladesh
Telephone	:	022222-80803
Fax	:	+880-(0)2-9880924
E-mail	:	dgda.gov@gmail.com
Web	:	www.dgda.gov.bd

Stamp and Date:

MINISTRY OF HEALTH & FAMILY WELFARE ★ DIRECTORATE GENERAL OF DRUG ADMINISTRATION ★ GOVT. OF THE PEOPLES REPUBLIC OF BANGLADESH ★ DHAKA

Major General Md. Shameem Haidar
Director General
Directorate General of Drug Administration
&
Licensing Authority (Drugs)
Government of the People's Republic of Bangladesh

27 APR 2026