Certificate No: 803 / GP-QLD

MINISTRY OF HEALTH VIETNAM
DRUG ADMINISTRATION

Certificate of a Pharmaceutical Product

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

Exporting (certifying) country: Vietnam

Importing (requesting) country: Republic of Moldova

Exporting name: TETRACYCLIN 1% EYE OINTMENT

Proprietary Names (if applicable) and dosage form: TETRACYCLIN 1%, eye ointment

Active Ingredient (s) and amount (s) per unit dose: Each tube of 5g eye ointment contains: Tetracycline hydrochloride 50mg

1. Is this product licensed to be placed on the market for use in the exporting country? yes ☑ no ☐ If yes, complete Box A, if no, complete Box B.

A. Product licence holder:
Medipharco Pharmaceutical Joint Stock Company
Address: 08 Nguyen Truong To street, Hue City, Vietnam
Tel: 02343.823099
Status of licence holder: Manufacturer
Number of product licence and date of issue: VD-26395-17 Dated: 06/02/2017
Date of renewal: December 31st 2024 (according to the letter No. 62/QĐ-QLD dated 08/02/2023)
The name and address of manufacturer producing the dosage form:
Medipharco Pharmaceutical Joint Stock Company
Address: 08 Nguyen Truong To street, Hue City, Vietnam
Tel: 02343.823099
Is an approved technical summary appended? yes ☐ no ☑
Is the attached product information complete and consonant with the licence?
yes ☐ no ☐ not provided ☑
Applicant for certificate if different from the licence holder: No

B. Applicant for certificate:

Status of applicant: Manufacturer

Why is authorization lacking?
not required ☐ not requested ☐ under consideration ☐ refused ☐

2. Does the certifying authority arrange for the periodic inspection of the manufacturing plant in which the dosage form is produced? yes ☑
no ☐ If no, proceed to question 3

Periodicity of routine inspection (years): 3 years

Has the manufacturer of this type of dosage form been inspected: yes ☑ no ☐

Do the facilities and operations conforms GMP as recommend by the World Health Organization? yes ☑ no ☐

3. Does the information submitted by the applicant satisfy the certifying authority on all aspects of the manufacture of the product undertaken by another party?
yes ☐ no ☐ if no, explain:

Address of the certifying authority:
Drug Administration - Ministry of Health Vietnam
138A, Giang Vo - Ha Noi - Viet Nam

Name of authorized person:
Signature:

Stamp and date
31/7/2023

CỤC QUẢN LÝ DƯỢC

TL. CỤC TRƯỞNG
TRƯỞNG PHÒNG
Nguyễn Văn Lợi

Digitally signed by Sili Oleg
Date: 2025.11.21 10:20:06 EEST
Reason: Mold-Trust Signature
Location: Moldova
MOLDOVA-EUROPEANA

BỘ NGOẠI GIAO NƯỚC CHXHCN VIỆT NAM
MINISTRY OF FOREIGN AFFAIRS OF THE S.R. OF VIETNAM

CHỨNG NHẬN / HỢP PHÁP HÓA LÃNH SỰ
CONSULAR AUTHENTICATION

1. Quốc gia ..... Viet Nam .....
Country

Giấy tờ, tài liệu này
This public document

2. do Ông (Bà) ..... Nguyễn Văn Lợi ..... ký
has been signed by

3. với chức danh ..... Head of Division .....
acting in the capacity of

4. và con dấu của ..... Drug Administration of Vietnam .....
bears the seal/stamp of

.....

được chứng nhận / hợp pháp hóa lãnh sự
Certified

5. tại ..... Hanoi ..... 6. ngày 10 08/ 2023
at *the* *(dd/mm/yyyy)*

7. Cơ quan cấp .... Consular Department .....
by

8. Số 0222950 / CLS
N°

Ký tên và đóng dấu
Signature and seal/stamp
Head of Division

Trần Thanh Vân