Certificate No: 4571/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: Viet Nam

Importing (requesting) country: Azerbaijan

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

Exporting name: 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: 02343.832814
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 2022 (according to the letter No. 4781/QLD-ĐK dated 02/06/2022)
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: 02343.832814
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 06/7/2022

CỤC QUẢN LÝ DƯỢC

KT. CỤC TRƯỞNG
TRƯỞNG PHÒNG
Nguyễn Văn Lợi

Digitally signed by Sîli Oleg
Date: 2023.01.04 09:28:55 EET
Reason: Moldsign Signature
Location: Moldova