HEALTH & FAMILY WELFARE DEPARTMENT BADDI, HIMACHAL PRADESH -173205
CERTIFICATE OF A PHARMACEUTICAL PRODUCT[1]

This certificate conforms to the format recommended by the World Health Organization
(General instructions and explanatory notes attached)

No. of Certificate : HFW-H [Drugs] 185/05/20-216 Valid Upto 05/03/2023
Exporting (certifying) Country : INDIA
Importing (requesting) Country : Syria
1. Name and Dosage form of Product : RABEVAN
1.1 Active ingredient (s)[2] and : (Combipack of Rabeprazole for Injection (Lyophilized)
And Sterilised Water for Injection BP 10 mL)
Amount (s) per unit dose[3] Each Combipack Contains:-
1) One Vial of Rabeprazole For Injection 20 mg
Each Vial Contains:-
Rabeprazole Sodium BP 20 mg
Excipients q.s.
2) With One 10mL Ampoule of Sterilized Water For Injection BP

Digitally signed by Chitic Ecaterina
Date: 2023.01.04 10:11:12 EET
Reason: MoldSign Signature
Location: Moldova

For Complete Qualitative Composition Including Excipients: NA

1.2 Is this Product licensed to be placed on the market for use in the exporting country?[5]

Yes ☒ No

1.3 Is this product actually on the market in exporting country?

YES ☒ No Unknown

If the Answer to 1.2 is YES, continue with section 2A and omit section 2B.
If the Answer to 1.2 is NO, omit section 2A and continue with section 2B.[6]

2A.

A.1 No. of Product Licence[7] : MB/05/158 in form No. 28
And date of Issue : 13.10.2020

A2. Product Licence holder : M/s Health Biotech Ltd.
Vill. Sandoli, Nalagarh Road,
Baddi, Distt. Solan [H.P.] India

A.3. Status of the Product-licence Holder[8] :

a. ☒ b. c.

A.3.1 For Categories b and c, The name and address of the Manufacturer producing the dosage form are[9]
Not Applicable

A.4. Is summary Basis of approval appended?[10] :

YES NO ☒

A.5. Is the attached, officially approved product information complete and consonant with the licence?[11] :

YES NO Not Approved ☒

A.6. Applicant for certificate if different from Licence holder[12]:
Not Applicable

2. B.

B. 1. Applicant for Certificate
(name and address)

B.2. Status of the Applicant:

a. b. c.

B.2.1. For categories b and c the name and address of the manufacture producing the dosage form are

B.3. Why is marketing authorization Lacking?

Not Required | Not Requested | Under Consideration | Refused

B.4. Remark[13]:

3. Does the certifying authority arrange for periodic inspection of the Manufacturing plant in which the dosage form is produced : YES ☒ NO Not Applicable[14]
If No or Not Applicable, proceed to Question 4

3.1 Periodicity of routine inspection (Years) : Once in a Year

3.2 Has the manufacturer of this type of dosage form been inspected? : YES ☒ NO

3.3 Do the facilities and operations conform to GMP as recommended By the World Health Organization?[15] : YES ☒ NO Not Applicable

4. Does the information submitted by the applicant satisfy the certifying Authority on all aspects of the manufacture of the product?[16] If No, explain: : YES ☒ NO

Licensing Authority cum Drug Controller * Himachal Pradesh * सत्यमेव जयते

Address of Certifying Authority:
State Drugs Controller
Controlling Cum Licensing Authority
Baddi Distt. Solan (H.P.) 173205 India
01795 244288, sdc4hp@gmail.com

Name of the Authorized Person : Mr. Navneet Marwaha
Designation : State Drugs Controller
Signature
Stamp and Date :

23-12-2020
(NAVNEET MARWAHA)
State Drugs Controller
Controlling cum Licensing Authority
Baddi Distt.Solan (H.P.)-173205
01795-244288,sdc4hp@gmail.com