DCA
DRUGS CONTROL ADMINISTRATION
TELANGANA

DRUGS CONTROL ADMINISTRATION
GOVERNMENT OF TELANGANA

Certificate of Pharmaceutical Product[1]

L.Dis. No. **883778/E1/2021**
WHO GMP Valid up to 01.07.2022

Exporting (Certifying) Country: **INDIA**
Importing (Requesting) Country: **Burkina Faso**

1. Name and Dosage form of the Product: **Rabies Antiserum 1000 I.U. I.H.S. (5 mL Liquid Vial) - VINRAB**

1.1 Active Ingredient(s) [2] and amount(s) per unit dose[3]: Each mL contains Enzyme Refined; Equine Antirabies immunoglobulin fragments not less than 200 I.U.

For Complete qualitative composition including excipients see attached[4]: Attached

1.2 Is this product Licensed to be placed on the market for use in the exporting country? [5] if yes, complete box A, if no complete box B

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

2.A.1 Number of product license[7] and date of issue: **01/MN/AP/2003/Sera/G,Dt. 17.12.2003,** **valid up to 16.12.2023** 2. A.2 Product License Holder: **VINS Bioproducts Limited, Sy. No. 117, Thimmapur (V), Kothur(M), Ranga Reddy(Dist.), - 509325, Telangana, India.** 2.A.3 Status of License Holder : [8] |a √ | b | c | d | 2.A.3.1 For categories b and c the name and address of the Manufacturer producing the dosage form is [9] : 2. A.4 Is a summary basis for approval appended? **Yes √ /** No 2.A.5 Is the attached, officially approved product information complete And consonant with the license? **Yes √** | No | Not Provided 2. A.6 Applicant for certificate, if different from the license holder[12] NA.	2.B.1 Applicant for certificate 2.B.2 Status of applicant : | a | b| c | d | 2.B.S.1 For categories (b) and (c) the name and address of the Manufacturer producing the dosage form is : [9] 2. B.3 Why is marketing authorization lacking? Not Required |Not Requested |Under Consideration | Refused 2.B.4 Remarks [13]

3. Does the certifying authority arrange for periodic inspection of the manufacturing plant in which the dosage form is produced? [14]**Yes √**\| No if no proceed to question 4

3.1 Periodicity of routine inspections (years): Not less than 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 \|

4. Does the information submitted by the applicant satisfy the certifying authority on all aspects of the manufacture of the product? [16]**Yes √ \|** No \| Not applicable \| if no, explain \|

Address of Certifying Authority: Name of the authorized person: **RAMDHAN GUGULOTH**
Deputy Director (FAC) & Licensing Authority

Drugs Control Administration,
Govt., of Telangana, Vengalraonagar,
Hyderabad 500 038.
Telephone/Fax Number 0091-40-23814119/0091-40-23814360

Signature: 25/10/2021

Stamp: DEPUTY DIRECTOR-I (FAC), LICENSING & CONTROLLING AUTHORITY DRUGS CONTROL ADMINISTRATION HYDERABAD TELANGANA STATE

Deputy Director Drugs Control Administration Government of Telangana HYDERABAD

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

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Date: 2021.11.12 20:06:18 EET
Reason: MoldSign Signature
Location: Moldova

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Explanatory notes

1. This certificate, which is in the format recommended by WHO, establishes the status of the pharmaceutical product and of the applicant for the certificate in the exporting country. It is for a single product only since manufacturing arrangements and approved information for different dosage forms and different strengths can vary.
2. Use, whenever possible, International Nonproprietary Names (INNs) or national nonproprietary names.
3. The formula (complete composition) of the dosage form should be given on the certificate or be appended.
4. Details of quantitative composition are preferred but their provision is subject to the agreement of the product-licence holder.
5. When applicable, append details of any restriction applied to the sale, distribution or administration of the product that is specified in the product licence.
6. Sections 2A and 2B are mutually exclusive.
7. Indicate, when applicable, if the licence is provisional, or the product has not yet been approved.
8. Specify whether the person responsible for placing the product on the market:
    a) manufactures the dosage form;
    b) packages and/or labels a dosage form manufactured by an independent company; or
    c) is involved in none of the above.
9. This information can only be provided with the consent of the product-licence holder or, in the case of non-registered products, the applicant. Non-completion of this section indicates that the party concerned has not agreed to inclusion of this information. It should be noted that information concerning the site of production is part of the product licence. If the production site is changed, the licence has to be updated or it is no longer valid.
10. This refers to the document, prepared by some national regulatory authorities, that summarizes the technical basis on which the product has been licensed.
11. This refers to product information approved by the competent national regulatory authority, such as Summary Product Characteristics (SPC)
12. In this circumstance, permission for issuing the certificate is required from the product-licence holder. This permission has to be provided to the authority by the applicant.
13. Please indicate the reason that the applicant has provided for not requesting registration.
    a) the product has been developed exclusively for the treatment of conditions — particularly tropical diseases — not endemic in the country of export;
    b) the product has been reformulated with a view to improving its stability under tropical conditions;
    c) the product has been reformulated to exclude excipients not approved for use in pharmaceutical products in the country of import;
    d) the product has been reformulated to meet a different maximum dosage limit for an active ingredient;
    e) any other reason, please specify.
14. Not applicable means the manufacture is taking place in a country other than that issuing the product certificate and inspection is conducted under the aegis of the country of manufacture.
15. The requirements for good practices in the manufacture and quality control of drugs referred to in the certificate are those included in the thirty-second report of the Expert Committee on Specifications for Pharmaceutical Preparations, WHO Technical Report Series No. 823, 1992, Annex 1. Recommendations specifically applicable to biological products have been formulated by the WHO Expert Committee on Biological Standardization (WHO Technical Report Series, No. 822, 1992, Annex 1).
16. This section is to be completed when the product-licence holder or applicant conforms to status (b) or (c) as described in note 8 above. It is of particular importance when foreign contractors are involved in the manufacture of the product. In these circumstances the applicant should supply the certifying authority with information to identify the contracting parties responsible for each stage of manufacture of the finished dosage form, and the extent and nature of any controls exercised over each of these parties.

DEPUTY DIRECTOR-I (HAC),
LICENSING & CONTROLLING AUTHORITY
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Ref: Rabies Antiserum 1000 I.U. I.H.S (5 mL Liquid Vial) -- VINRAB

Sl. No.	Ingredients	Quantity	Application	Pharmacopoeial Standard
	ACTIVE INGREDIENTS			
1.	Enzyme Refined, Equine Antirabies Immunoglobulin fragments	Not less than 200 I.U./mL	Active ingredient	IHS
	EXCIPIENTS			
2.	Cresol	0.0075mL	Preservative	BP
3.	Glycine	112.5mg/mL	Stabilizer	BP
4.	Sodium Chloride	45mg/mL	Isotonic	BP

Name of the Authorized person: **RAMDHAN GUGULOTH**
Deputy Director (FAC) & Licensing Authority

Signature:

25/02/2021

Deputy Director Drugs Control Administration Government Of Telangana HYDERABAD

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