

No. of Certificate: 26/1/Drug/20/2019/8456 | 11674

Exporting (Certifying) Country : INDIA

Date: .....

Importing (Requesting) Country: Afghanistan, Africa, Algeria, Azerbaijan, Bahrain, Belarus, Bhutan, Bolivia, Brazil, Burkina Faso, Cambodia, Cameroon, Chad, Costa Rica, D.R. Congo Egypt, Equatorial Guinea, Guyana, Gabon, Ghana, Gambia, Indonesia, Iran, Iraq, Israel, Ivory Coast, Jamaica, Jordan, Kazakhstan, Kenya, Korea, Kuwait, Kyrgyzstan, Madagascar, Malawi, Malaysia, Mongolia, Mozambique, Myanmar, Nepal, Nigeria, Oman, Panama, Peru, Philippines, Qatar, Republic of Congo, Republic of Guinea, Romania, Russia, Rwanda, Saudi Arabia, Senegal, Serbia, Singapore, South Africa, Sri Lanka, Sudan, Syria, Tajikistan, Tanzania, Thailand, Togo, Turkmenistan, UAE, Uganda, Ukraine, Uzbekistan, Venezuela, Vietnam, Yemen.

### CERTIFICATE OF A PHARMACEUTICAL PRODUCT<sup>1</sup>

1. Name & dosage form of product: **DIZIDOXINE TABLETS (Mecizine Hydrochloride & Pyridoxine Hydrochloride Tablets)**

1.1 Active ingredient (s)<sup>2</sup> and amount (s) per unit dose<sup>3</sup> (complete qualitative composition including excipients, see attached)

Composition:

Each uncoated tablet contains:

Mecizine Hydrochloride USP.....25 mg

Pyridoxine Hydrochloride (Vitamin B<sub>6</sub>).....50 mg

Excipients.....q.s.

1.2 Is this product licensed to be placed on the market for use in the exporting country? **YES**

If yes, complete box A, if no, complete box B:

1.3 Is this product actually on the market in the exporting country?<sup>5</sup> **YES**

A		B	
2A.1	Number of product licence <sup>7</sup> & date of issued: <b>65/UA/2016 &amp; 65/UA/SC/P-2016 dt. 24/08/2016</b>	2B.1	Applicant for certificate (name & address)
2A.2	Product licence holder: <b>M/s SIGNATURE PHYTOCHEMICAL INDUSTRIES. 122, MI, SELAQUI INDUSTRIAL AREA, DEHRADUN</b>	2B.2	Status of applicant: (a) <input type="checkbox"/> (b) <input type="checkbox"/> (c) <input type="checkbox"/> (d) <input type="checkbox"/>
2A.3	Status of product licence holder <sup>8</sup> : (a) <input checked="" type="checkbox"/> (b) <input type="checkbox"/> (c) <input type="checkbox"/>	2B.2.1	For categories b and c the names and address of the manufacturer producing the dosage form are <sup>9</sup> :
2A.3.1	For categories b and c the name and address of the manufacturer producing the dosage form are <sup>9</sup> : Not Applicable	2B.3	Why is marketing authorization lacking? Not <input type="checkbox"/> Not <input type="checkbox"/> Under <input type="checkbox"/> Refused <input type="checkbox"/> Required Requested Consideration
2A.4	Is summary basis of Approval appended <sup>10</sup> ? Yes <input type="checkbox"/> No <input checked="" type="checkbox"/>	2B.4	Remark: <sup>13</sup>
2A.5	Is the attached, officially approved product information complete and consonant with the licence <sup>11</sup> ? Yes <input type="checkbox"/> No <input type="checkbox"/> Not provided <input checked="" type="checkbox"/>		
2A.6	Applicant for certificate if different from licenced holder: Not Applicable <sup>12</sup>		
3.	Does the certifying authority arrange for periodic inspection of the manufacturing plant in which the dosage form is produced? Periodically of routine inspection (years): <b>Once in a year</b>	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> NA <input type="checkbox"/>	
3.2	Has the manufacture of this type of dosage form been inspected	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/>	
3.3	Do the facilities and operations conform to GMP as recommended by the World Health Organization? <sup>15</sup>	Yes <input checked="" type="checkbox"/> No <input type="checkbox"/>	
4.	Does the information submitted by the applicant satisfy the certifying authority on all aspects of the manufacture of the product? <sup>16</sup>	Yes <input type="checkbox"/> No <input checked="" type="checkbox"/> NA <input checked="" type="checkbox"/>	

**WHO-GMP-CERT.NO. 26/1/Drug/20/2019/8456, Valid up to: 14/05/2023**

Address of certifying authority:

**Drug Licensing & Controlling Authority**

**Directorate General of Medical Health & Family Welfare,  
Sahastradhara Road, Dehradun-248001 (Uttarakhand) (INDIA)**

**Directorate General of Medical Health & Family Welfare,  
Sahastradhara Road, Dehradun-248001 (Uttarakhand) (INDIA)**



Name of the authorized person:

Signature:

Stamp & Date

Stamp & Date

*Tajber Singh*  
19/6/2020  
**Tajber Singh**  
Drug Controlling & Licensing Authority (Mfg.)  
(Uttarakhand)

**Attested**

*Sushmita*  
**SUSHMITA**  
Secretary



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1. This certificate which is in the form recommended by WHO, establishes the status of the pharmaceuticals 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.
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 be provided only with the consent of the product-licence holder or, in the case of non-availability, the applicant must indicate that the party concerned has not agreed inclusion of this information. It should be noted that information concerning the site of production is part of the product licence. If the product licence is not available, the product licence must be updated or it will be invalid.
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 a Summary of Product Characteristics (SPC).
12. In this circumstance, permission for issuing the certificate is required from the product-licence holder. This permission must 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 of improving its stability under tropical conditions.
  - (c) The product has been reformulated to exclude excipients not approved for use in pharmaceuticals products in the country on 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 that 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 Export Committee on specifications of Pharmaceutical Preparations (WHO Technical Report Series No. 833, 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.
17. The layout for this Model Certificate is available on diskette in WordPerfect from the Division of Drug Management and Policies, World Health Organization, 1211 Geneva 27, Switzerland.



39935

सं. No. 39935  
वर्गिण्य मंडल में सहायक सचिव के हस्ताक्षर सत्यापित किए जा रहे हैं।  
The Signature of Asstt. Secretary of Commerce/Secretary of External Affairs  
को विमोचनी नहीं है।  
विदेश मंत्रालय इन दस्तावेजों के लिए कोई भी ज़िम्मेदारी नहीं लेता है।  
Ministry of External Affairs has no responsibility for the contents of this document.

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