

Office of The Commissioner, Food & Drugs Administration M.S. Bandra - Kurla Complex, Bandra (E), Mumbai - 400 051

Date:

0 1 MAR 2019

CERTIFICATE OF GOOD MANUFACTURING PRACTICES

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

Certificate No.: NEW-WHO-GMP/CERT/KD/82148/2019/11/27049

On the basis of the inspection carried out on 06/12/2018, 07/12/2018 and 19/01/2019 ,we certify that the site indicated on this Certificate complies with Good Manufacturing Practices for the dosage forms, categories and activities listed in Table 1.

Name of the Firm

EMIL PHARMACEUTICAL INDUSTRIES PVT.

LTD.

Address

101, MANGALUM, KULUPWADI, BORIVALI (EAST) MUMBAI 400066

Manufacturing At

PLOT NO. J-76, M.I.D.C., TARAPUR, BOISAR THANE 401506 MAHARASHTRA STATE,

INDIA

Licence No.

KD669A In Form 25A, KD780A In

Table 1

Sr.No.	Dosage Form(s)	Categor(ies)	Activity(les)	
1	Capsules	General (Other than Cephalosporins, Penicillin, Cytotoxic, Hormones)	Production, Filling, Packing, labelling, Quality Control, Quality Assurance	
	Oral Powders / Granules / Pellets	General (Other than Cephalosporins, Penicillin, Cytotoxic, Hormones)	Production, Filling, Packing labelling, Quality Control, Quality Assurance	
3	Tablets	General (Other than Cephalosporins, Penicillin, Cytotoxic, Hormones)	Production, Filling, Packing, labelling, Quality Control, Quality Assurance	

The responsibility for the quality of the individual batches of the pharmaceutical products manufactured through this process lies with the manufacturer.

This certificate remains valid until 12 Feb 2022. It becomes invalid if the activities and / or categories certified herewith are changed or if the site is no longer considered to be in compliance with GMP

Address of certifying authority

Food & Drug Administration, M

Bandra-kurla Complex. Bandra (E). Mumbai - 400

Maharashtra, INDIA.

Tel +91-22-26592363/

Fax. +91-22-2659195

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MIL PHARMACEUTICAL

NEW-WHO GM

Name of the Authorised person : A. T. NIKHADE

Signature

Stamp and Date Joint Commissioner (HQ) **Authority**

Food & Drug Administration, M.S.

Bandra (E), Mumbai

Maharashtra State, India

Date: 27 Feb 2019

ATTESTED

ED SIGNATORY

ICC

JIGNA KOTHARI Asst. Director

IMC CHAMLER OF COMMERCE AND INDUSTRY

MUMBAHINDIA

MAR 2019

Explanatory notes

- This certificate which is in the format recommended by WHO, certifies the status of the site listed in point 1 of the certificate.
- 2. The certification number should be traceable within the regulatory authority issuing the certificate.
- 3. Where the regulatory authority issues a licence for the site, this number should be specified record "not applicable" in cases where there is no legal framework for the issuing of a licence.
- Table 1
 List the dosage forms, starting materials, categories and activities. Examples are given below.

Example -1

Pharmaceutical Product (s)1	Category (ies)	Activity (ies)	
Dosage form (s)			
Tablets	Cytotoxic	Packaging	
	Hormone	Production, Packaging, Quality control.	
Injectables	Penicillin	Repackaging & Labelling.	
	Cefalosporin	Aseptic preparation, Packaging, Labelling.	

Example - 2.

	utical Product (s)1	Category (ies)	Activity (ies)
-	aterial (s)2		
Paracetamol		Analgesic	Synthesis, Purification,
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Market St.	2.2.1.		Kandivali (E), Mumbai-400 101
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