



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

Date: 27/11/2014

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/23005/2014/11/8718**

On the basis of the inspection carried out on **11/08/2014 AND 12/08/2014**, 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.

1. Name of the Firm : **EMIL PHARMACEUTICAL INDUSTRIES PVT. LTD.**
Address : **101, MANGALUM, KULUPWADI, BORIVALI (EAST) MUMBAI 400066**
Manufacturing At : **NO. J-76, M.I.D.C., TARAPUR, BOISAR, THANE 401506 MAHARASHTRA STATE, INDIA**
2. Licence No. : **KD669A In Form 25A, KD780A In Form 28A**

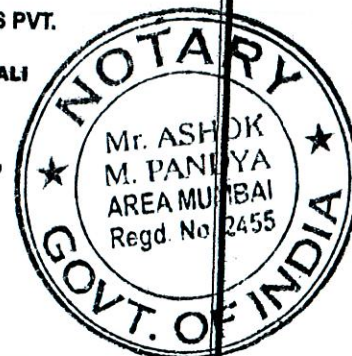


Table 1

Sr.No.	Dosage Form(s)	Categor(ies)	Activity(ies)
1	Capsules	General (Other than Cephalosporins, Penicillin, Cytotoxic, Hormones)	Production, Filling, Packing, labelling, Quality Control, Quality Assurance
2	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

No 3 2 6 0 0

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 28 Sep 2016. 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.S.
Bandra Kurla Complex,
Bandra (E), Mumbai - 400 051.
Maharashtra, INDIA.
Tel: +91-22-26592363/64
+22-26591959
0141121162

Name of the Authorised person : **O.S. SADHWANI**

Signature:

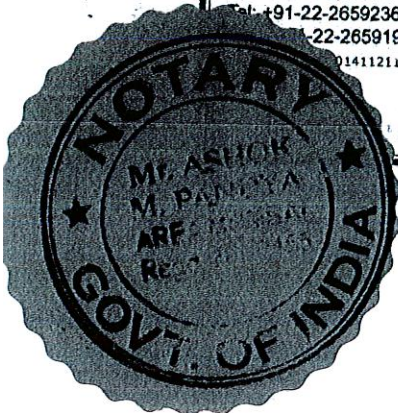
Stamp and Date: **Joint Commissioner (HQ) & Controlling Authority**
Food & Drug Administration, M.S.
Bandra (E), Mumbai.
Maharashtra State, India
Date: 21 Nov 2014

INSTITUTO DE SAUDE PUBLICA

Certifico que o presente fotocopia es fiel al original que he tenido a la vista y consta de 4 páginas

MINISTRO DE F.E.

Stgo. 19.11.2014



21 NOV 2014

ATTESTED

TRUSHA SHAH
Manager

AUTHORISED SIGNATORY
INDIAN MERCHANTS' CHAMBER
MUMBAI

Explanatory notes

1. 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 in the record "not applicable" in cases where there is no legal framework for the issuing of a licence.
4. Table 1
List the dosage forms, starting materials, categories and activities. Examples are given below.

NUEVA DELHI, INDIA

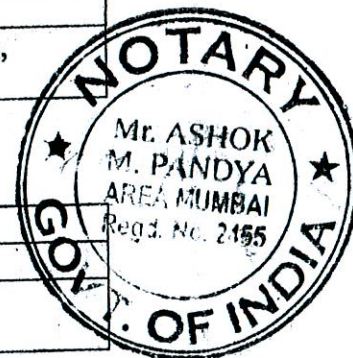
El Consul de Chile en Nueva Delhi, certifica la autenticidad de la firma de don Manuel J. Saldias Kuhlmann funcionario para asuntos consulares del Ministerio de Relaciones Exteriores de India.
Derechos percibidos: US \$ 12
NUEVA DELHI 04 de Dic de 2014

Example - 1

Pharmaceutical Product (s)1	Category (ies)	Activity (ies)
Dosage form (s)	COMPROB. DE PAGO No. <u>348</u>	MANUEL J. SALDIAS KUHLMANN Cónsul de Chile
Tablets	ACT. No. <u>10</u> Cytotoxic Hormone	Packaging Production, Packaging, Quality control.
Injectables	Penicillin Cefalosporin	Repackaging & Labelling. Aseptic preparation, Packaging, Labelling.

Example - 2.

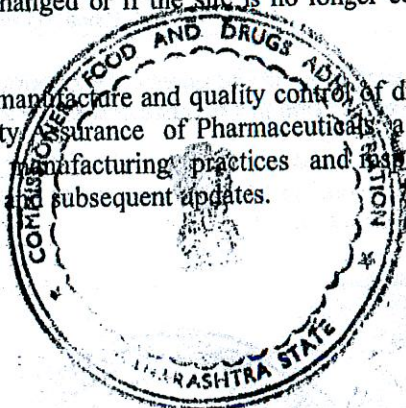
Pharmaceutical Product (s)1	Category (ies)	Activity (ies)
Starting material (s)2		
Paracetamol	Analgesic	Synthesis, Purification, Packing, Labelling.



155157 Use, whenever available. International Nonproprietary Names (INNs) or otherwise national nonproprietary names.
The certificate remains valid until the specified date. The certificate becomes invalid if the activities and/or categories certified are changed or if the site is no longer considered to be in compliance with GMP.
The Signature/Secretary of Commerce
The requirements for good practices the manufacture and quality control of drugs referred to in the certificate are those included in Quality Assurance of Pharmaceuticals, a compendium of guidelines and related materials. Good manufacturing practices and inspection. Volume 2, 1999. World Health Organization, Geneva and subsequent updates.

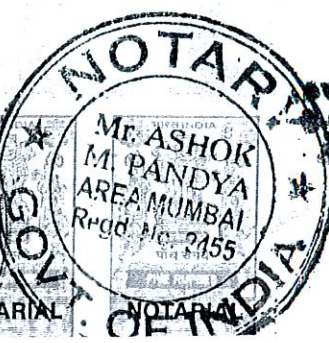
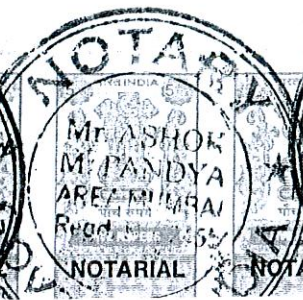
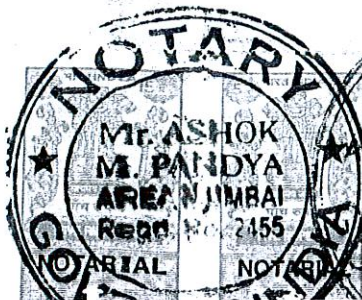


(त्रिलोक चन्द)
(TRILOK CHAND)
अनुभाग अधिकारी (अतिरिक्त)
Section Officer (Additional)
प्रदेश मंत्रालय, नई दिल्ली
Ministry of External Affairs
New Delhi



CERTIFIED TRUE COPY

ASHOK M. PANDYA
ADVOCATE & NOTARY (GOVT. OF INDIA)
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Mans, L. T. Road, Berivali (W)
MUMBAI-400 092



28 NOV 2014