



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

Date : 02 MAR 2019

INSTITUTO DE SALUD PUBLICA

Certifico que la presente
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MINISTRO DE FE

Stgo. 25/07/19

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/75607/2019/11/27031**

On the basis of the inspection carried out on **04/10/2018 & 05/12/2018**, 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 : **PELL TECH HEALTH CARE PVT. LTD.**
Address : **PLOT NO. 20B, TANSA FARM ESTATE,
VILLAGE MET, GONSAI, BHIWANDI-WADA,
THANE 421312 MAHARASHTRA STATE,
INDIA**
- Licence No. : **KD622 In Form 25,
KD437 In Form 28**

Table 1

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

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

certificate remains valid until 26 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.

certifying authority :

Name of the Authorised person : **A. T. NIKHADE**

Food & Drugs Administration, M.S.
Bandra – Kurla Complex,
(E), Mumbai – 400 051.

Signature :

Stamp and Date : **Joint Commissioner (HQ) & Controlling
Authority**

Food & Drug Administration, M.S.

Bandra (E), Mumbai.

Maharashtra State, India

Date: 27 Feb 2019

Maharashtra, INDIA

Tel: +91-22-26592363/64

Fax: +91-22-26591999

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PELL TECH HEALTH CARE PVT. LTD.

NEW-WHO-GMP/CERT/KD/75607/2019/

727031

ATTESTED BY ME

J D RAWAL

RAJ GEORGE

ATTESTED

Prora

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 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.

Example -1

Pharmaceutical Product (s)	Category (ies)	Activities
Dosage form (s)		
Tablets		Packaging, Quality
Injectables		Labelling, Packaging,

भारत सरकार GOVERNMENT OF INDIA
अपोस्टिल / APOSTILLE
(Convention de La Haye du 5 octobre 1961)
INDIA
This public document of the type
COMMERCIAL DOCUMENT
is issued to PELL TECH HEALTH CARE PVT LTD
has been signed by BIJU GEORGE
with the seal / stamp of SR MANAGER, IMC CHAMBER OF
COMMERCE AND INDUSTRY, MUMBAI-INDIA
Certified by
Section Officer(OI) MINISTRY OF EXTERNAL AFFAIRS
on 15-Mar-2019 at NEW DELHI, INDIA
with reference no. MHMC0006393919

Seal / Stamp
 Signature
 (भावना शर्मा)
 अनुभाग अधिकारी (ओ.ओ.)
 Section Officer (OI)
 सी. पी. वी. प्रभाग / C.P.V. Division
 Ministry of External Affairs, New Delhi

6. The requirements for good practices in 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.