



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

Date : **21 SEP 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/84495/2019/11/29495**

On the basis of the inspection carried out on **02/07/2019, 03/07/2019 and 21/08/2019**, 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.

1. Name of the Firm : **MEHTA API PVT. LTD.**
Address : **GUT NO. 546, 571, 519 & 520, VILLAGE KUMBHAVALI, TALUKA PALGHAR, THANE 401506 MAHARASHTRA STATE, INDIA**
2. Licence No. : **KD659 In Form 25, KD458 In Form 28**

Table 1

Sr.No.	Dosage Form(s)	Categor(ies)	Activity(ies)
1	Active Pharmaceutical Ingredients (Bulk Drugs)	General (Other than Cephalosporins, Penicillin, Cytotoxic, Hormones)	Synthesis, Purification, 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 18 Sep 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.S.

Bandra-kurla Complex,

Bandra (E), Mumbai – 400 051.

Maharashtra, INDIA.

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MEHTA API PVT. LTD. - NEW-WHO-GMP/CERT /KD/84495/2019/11/29495

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

Signature :

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

Food & Drug Administration, M.S.

Bandra (E), Mumbai.

Maharashtra State, India

Date: 19 Sep 2019



19 SEP 2019