Certifico que la presente fotocopia es fiel al original que he tenido, a la vista

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

1 3 AUG 2018

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/72365/2018/11/24339

On the basis of the inspection carried out on 07/06/18, 08/07/18 and 18/07/2018, 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

CIRON DRUGS & PHARMACEUTICALS PVT.

LTD.

Address

N-118,118/1, 119,119/1,119/2,113 MIDC, TARAPUR, BOISAR, DIST. THANE 401506

MAHARASHTRA STATE, INDIA

Licence No.

KD80 in Form 25, KD74 in Form 28, KD/3 In Form 28B

| Sr.No. | Dosage Form(s) | Categor(ies) | Activity(ies) |
|--------|---|--|---|
| 1 | External Preparation (Ointments / Creams / Lotion/Gel/Ear drop/Nasal drop/Spray) | General (Other than Cephalosporins, Penicillin, Cytotoxic, Hormones) | Production, Filling, Packing, labelling, Quality Control, Quality Assurance |
| 2 | Eye / Ear Drops | General (Other than Cephalosporins, Penicillin, Cytotoxic, Hormones) | Production, Filling, Packing, labelling, Quality Control, Quality Assurance |
| 3 | Eye Drops / Ophthalmic Preparations | General (Other than Cephalosporins, Penicillin, Cytotoxic, Hormones) | Production, Filling, Packing, labelling, Quality Control, Quality Assurance |
| 4 | Inhalation | General (Other than Cephalosporins, Penicillin, Cytotoxic, Hormones) | Production, Filling, Packing, labelling, Quality Control, Quality Assurance |
| 5 | Liquid Injection (SVP) | General (Other than Cephalosporins, Penicillin, Cytotoxic, Hormones) | Production, Filling, Packing, labelling, Quality Control, Quality Assurance |
| 6 | Liquid Orals | 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 01 Aug 2021 . It becomes invalid if the activities and / or categories certified shanged or if the site is no longer considered to be in compliance with and

Address of certifying Food & Drug Admi Bandra-kurla Cor

rised person : A. T. NIKHADE

Signature:

and Date : Joint Commissioner (HQ) & Controllin Authority Food & Drug Administration, M.S. Bandra (E), Mumbai.

Maharashtra State, India Date:09 Aug 2018

n.9 Alig 2018

Goorded with the Chamber SAMIR PAUL PINTO

A copy of this document / CERTIFICATE

Authorised Signatory

Bombay Chamber of Commerces





भारत सरकार GOVERNMENT OF INDIA अमोस्टिल / APOSTILLE

Convention de La Haye du 5 octobre 1961)

This public document of the type COMMERCIAL DOCUMENT

ississued to CIRON DRUGS & PHARMACEUTICAL

PVT, LTD

SAMIR PAUL PINTO

with the seal / stamp of ASSTT MANAGER, BOMBAY CHAMBER OF COMMERCE AND INDUSTRY

Certified by

Section Officer(OI) MINISTRY OF EXTERNAL AFFAIRS
on 27-Nov-2018 at NEW DELHI, INDIA

WITH THE PROPERTY MHMC0035552818

Seatusta



(OCSABBAN)
(OCSABBAN)
Section Officer (O.I.)
জারা, সল্যাৎCP.V. Division
সিংগ্রামিক স্থান্য সেরা বিশ্ব

Y Assis



Office of The Commissioner, Food & Drugs Administration M.S. Bandra - Kurla Complex, Bandra (E), Mumbai - 400 051 Date: 1 3 AUG 2018

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TARAPUR, BOISAR, DIST. THANE 401506 MAHARASHTRA STATE, INDIA

2. Licence No.

KD80 in Form 25,

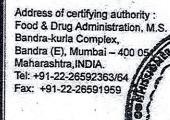
KD74 In Form 28, KD/3 In Form 28B

Table 1

| Sr.No. | Dosage Form(s) | Categor(ies) | Activity(ies) |
|--------|---------------------------------|--|---|
| 7 | Lyophilised / Powder injectable | 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 01 Aug 2021 . 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.



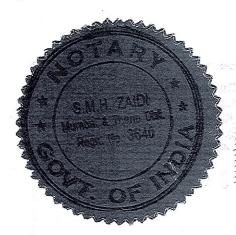
Name of the Authorised person : A. T. NIKHADE

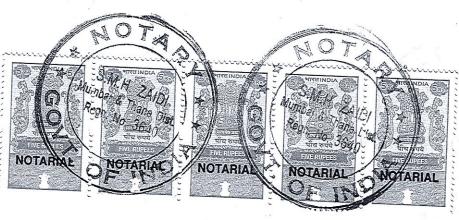
Signature:

imp and Date : Joint Commissioner (HQ) & Controlling Authority

> Food & Drug Administration, M.S. Bandra (E), Mumbai.

Maharashtra State, India Date:09 Aug 2018





S. M.H. ZAIDI
NOTARY
Government of India
Mumbal & Thane Dist

95 NOV 2018