



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/ND/89374/2019/11/29520**

On the basis of the inspection carried out on **20/06/2019, 21/06/2019 and 01/08/2019**, 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.

- Name of the Firm : **LIVEALTH BIOPHARMA PVT LTD**
Address : **77, RATNAJYOT IND PREMISES IRLA
GAOTHAN, VILE PARLE WEST MUMBAI
400056**
Manufacturing At : **D-82, MIDC AREA, CROSS ROAD NO.4-A,
HINGNA, NAGPUR 440028 MAHARASHTRA
STATE, INDIA**
- Licence No. : **MH102695A In
Form 28A**

Table 1

Sr.No.	Dosage Form(s)	Categor(ies)	Activity(ies)
1	Liquid Injection (SVP)	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 08 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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LIVEALTH BIOPHARMA PVT LTD - NEW-WHO-
GMP/CERT/ND/89374/2019/11/29520

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: 21 Sep 2019



21 SEP 2019