FDA MAHARASHTRA

Office of The Commissioner,
Food & Drugs Administration M.S.
Bandra – Kurla Complex,
Bandra (E),
Mumbai – 400 051.
Date : 18 SEP 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/ND/76242/2018/11/24998

On the basis of the inspection carried out on 06/04/2017, 07/04/2017 and 06/06/2017 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 : 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**
2. 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 12 Jul 2019 . 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: +91-22-26592366/64
Fax: +91-22-2659[illegible]
1VIL10376242201805

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:18 Sep 2018

FOOD AND DRUG ADMINISTRATION MAHARASHTRA STATE

18 SEP 2018

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Reason: MoldSign Signature
Location: Moldova

LIST OF PRODUCT APPROVED UNDER WHO GMP

No. of certificate : NEW-WHO-GMP/CERT/ND/76242/2018/11 VALID UP TO :12 Jul 2019
/24998

Name of Manufactring Firm : LIVEALTH BIOPHARMA PVT LTD
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

Drug License No : MH002698A In Form 28A

Sr.No.	Name of the Product	Composition
1	BRAINIMAGE Gadopentetate Dimeglumine Injection USP 469mg/ml	Each ml Contains Gadopentetate Dimeglumine USP 469 mg Water for Injection USP qs
2	LIVEALTHIMAGE - 300 Iopamidol Injection USP 300mg I/ml	Each ml Contains Iopamidol USP 612.34 mg Water for Injection USP qs Iodine Content 300 mg/ml
3	LIVEALTHIMAGE - 370 Iopamidol Injection USP 370mg I/ml	Each ml Contains Iopamidol USP 755.22 mg Water for Injection USP qs Iodine Content 370 mg/ml
4	RADICALSCAN - 60 Diatrizoate Meglumine & Diatrizoate Sodium Injection 60%	Each ml Contains Diatrizoate Sodium USP 79 mg Diatrizoate Meglumine USP 521 mg Water for Injection USP qs Iodine Content 292 mg/ml
5	RADICALSCAN - 76 Diatrizoate Meglumine & Diatrizoate Sodium Injection 76%	Each ml Contains Diatrizoate Sodium USP 100 mg Diatrizoate Meglumine USP 660 mg Water for Injection USP qs Iodine Content 370 mg/ml
6	XREYIMAGE - 300 Iohexol Injection USP 300mg I/ml	Each ml Contains Iohexol USP 647 mg Water for Injection USP qs Iodine Content 300 mg/ml
7	XREYIMAGE - 350 Iohexol Injection USP 350mg I/ml	Each ml Contains Iohexol USP 755 mg Water for Injection USP qs Iodine Content 350 mg/ml

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Fax: +91-22-26591959
1VIL1037624220180918

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 18 Sep 2018

18 SEP 2018