
Bharat Parenterals Limited
Offering a rich array of QUALITY formulations

Welcome to our
Journey towards Excellence!

"We endeavor to manufacture a rich array of quality formulations supported by our strong Formulation & Development department (DSIR Approved) and State-of-Art manufacturing facilities for our own market and for several reputed organizations domestically as well as internationally."

Get to know us!

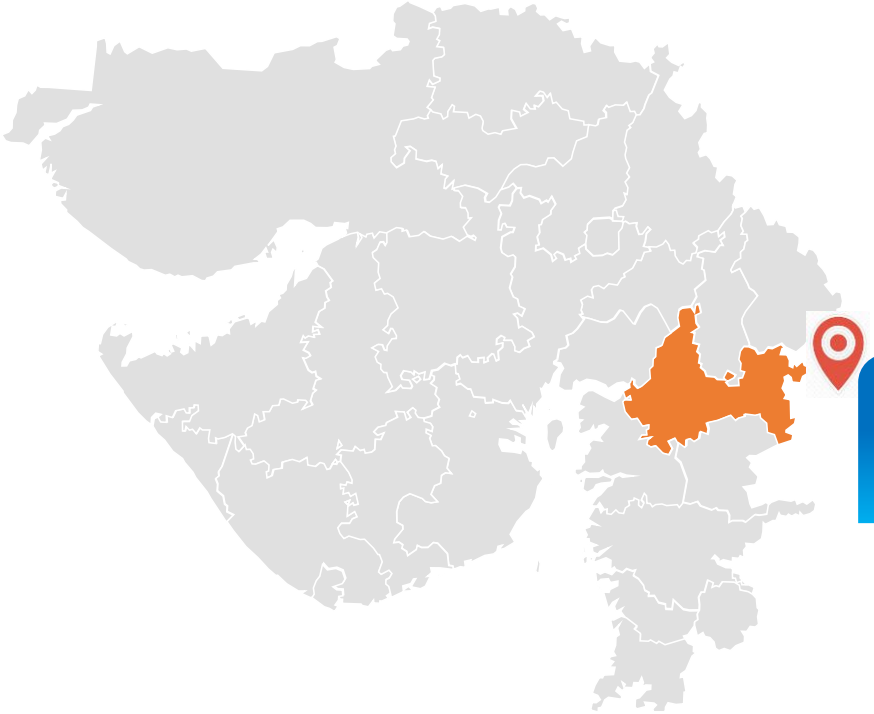
MANUFACTURED TO cGMP STANDARDS

Bharat Parenterals Limited is a pharmaceutical manufacturing company having its manufacturing facility at Haripura, Vadodara. It was formed under the chairmanship of Mr. Ramesh Desai and incorporated as limited company on 3rd September 1992.

Since its inception in 1992, Bharat Parenterals Ltd has traversed a long way in order to be recognized as a Trusted Global Player amongst the pharmaceutical companies in India today.

Growing at a fast yet steady pace, the second-generation contract development & manufacturing organization, led by Mr. Bharat R Desai, is well on its way to becoming a leader in New Age Formulations, both nationally and internationally.

Our Inception



Vadodara,Gujarat
Head Office + Factory

Our Vision

A visionary pharmacist's philosophy to enrich every human's life and an aspiration to have firm roots and make a mark in the Indian pharmaceutical industry were the foundation stones of Bharat Parenterals Limited.

" Providing access to affordable and quality medicines to the whole wide world and leading the way for medical innovation."

Our Mission

- To become a significant global player by providing high quality, affordable and innovative solutions in medicine and treatment.
- Aspire to acquire a major share in the Contract Research and Manufacturing Services (CRAMS).
- Provide uniform, high-quality medicines in the international and domestic market
- Build mutually beneficial relationships with customers by maintaining a world-class facility and high standard of medicines.
- Set up an environmentally & socially responsible organization

Our Core Strengths

- As a progressive manufacturing organization, we engage in efficient sourcing, formulation development, manufacturing, marketing and distribution, with an aim for the highest quality products. A fully integrated company - we have our in-house Formulation R&D, Business Development, Manufacturing, and Regulatory Compliance capabilities

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- Our state-of-the art Formulation & Development Department which has been recognized by the Department of Scientific & Industrial Research (DSIR) is well equipped with modern sophisticated equipment for New Formulation Development as well as scale-up of the formulations as per customer-specific requirement under the supervision of qualified and experienced technical staff.

- As a backbone to these processes our Analytical Development Laboratory with ultra-modern instruments for Method Development and Analytical Validations ensures the highest standards of Quality Control.

Legacy of more than 25 years of expertise in:

- Custom Manufacturing
- Formulation & Development
- Analytical Method Validation
- Regulatory Services
- Technology & Process Transfer
- Stability Studies

Other BPL Services

- Contract Research
- Contract Manufacturing

Formulation Research & Development

- A strong Formulation & Development set up has been a critical part of our business strategy. It is the backbone which has fortified new product lines and novel formulations systems year after year.

- Our team of qualified scientists work dedicatedly day in and day out in order to make a wide range of products available in line with the needs of our Domestic and Exports markets.

- Scientists undertake novel molecules for dosage form development by determining its composition, correct dosage, safety and efficacy for manufacturing and commercialization. Besides developing formulations, our R&D team improves existing formulations to ensure maximum efficacy and product compliance by the patient.

Our Formulation & Development Department has the following Capabilities:

- To Formulate, Evaluate and Develop the generic Product which is similar as Reference Product with same Strength and dosage form.
- Target to meet all the tests as Reference product and respective Pharmacopoeial monograph.
- To Formulate & evaluate Robust formulation which is Stable as per storage requirements.
- Operate and calibrate all the Lab scale machines which is required to manufacturing of Tablets / Capsules / Injection / Lyophilize product / Cream / Suspension and etc.
- Reverse Engineering of Reference product.
- Pre-formulation study of API'S and Drug – Excipients Compatibility study.
- Design manufacturing Formula and optimize the quantity of Excipients based on QbD.
- Technology transfer to manufacturing site.
- Exhibit batch executions for commercialization of finished product.

Formulation Research & Development

Formulation Research & Development

Formulation Research & Development

Analytical Development Laboratory

- Our fully equipped GLP Analytical Laboratory is well suited for drug product Analytical Method Development, Validation and Analysis to support R&D, Exhibit, Clinical, Validation and Commercial activities.

- Our Microbiology Lab is equipped to carry out Method Development and Validations/Verifications of Preservative Efficacy, Sterility, Microbial Limit and Bacterial Endotoxins

Manufacturing Facilities

- The BPL production facilities are located on the outskirts of Vadodara city, about 24 kms away at village Haripura, Vadodara, Gujarat and is spread over 30,000 sq. mtrs.

  and house three separate production blocks:

- ❑ General Unit
- ❑ Beta-Lactam Unit
- ❑ Antiretroviral Products Unit

A new unit dedicated to Cephalosporins will soon be operational

- In 2015, a new building was inaugurated which houses the **DSIR approved** Formulation, Research and Development laboratory, ADL Laboratories & the administrative departments.

- The plant is well operational with the most contemporary and validated manufacturing and analytical equipment with detailed SOPs in place for Quality Management, Personnel Management, Premises, Equipment, Documentation, Production, Complaints, and Self Inspection/Audits.

Manufacturing Capacity

GENERAL BLOCK			
Sr. No.	**Finished Dosage Forms**		**Annual Capacity**
1	Tablets		1.35 Billion
2	Capsules		1.439 Billion
3	Injection	Ampoules	104 Million
		Vials	168 Million
		Dry Powders	52 Million vials
4	Oral Liquids		73 Million bottles
5	Ointments/Creams/Gels		26 Million tubes
6	Powders		52 Million pouches

Continue

BETA LACTAM BLOCK		
Sr. No.	Finished Dosage Forms	Annual Capacity
1	Tablets	315 Million
2	Capsules	360 Million
3	Dry Syrup	27 Million bottles
4	Dry Powder for Injection	36 Million vials

General Liquid and Ointment

General Liquid and Ointment

General Tablet

General Tablet

General Injection

General Injection

General Injection

Quality Control

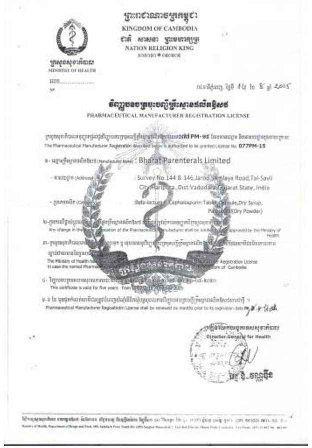
Quality Control

Quality Control

Accreditations

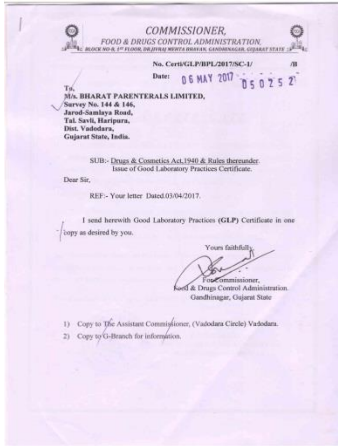
Certificates


Republic of Yemen
Ministry of Public Health and Population
Supreme Board of Drugs & Medical Appliances
Director General
وزارة الصحة العامة والسكان
الهيئة العليا للأدوية والمستلزمات الطبية
المدير العام
Bharat parenteral
BHARAT PARENTERALS LIMITED.
SURVEY NO.144 & 146, VILL. HARIPURA, JAROD SAMLAYA ROAD, TAL-SAVLI, INDIA
ACCEPTED LINES:
I) GENERAL PRODUCTION LINES:
1- SOLID DOSAGE FORM: TABLET & CAPSULE.
2- LIQUID DOSAGE FORM FOR ORAL USE.
3- EXTERNAL PREPARATIONS: CREAM, OINTMENT, JELL AND LIQUID FOR EXTERNAL USE.
4- STERILE DOSAGE FORM: POWDER FOR INJECTION ONLY.
II) B- LACTAM PRODUCTION LINES (PENCILLIN AND CEPHALOSPORIN):
1- NON- STERILE DOSAGE FORMS: TABLET, CAPSULE AND DRY SYRUP.
2- STERILE DOSAGE FORM: POWDER FOR INJECTION ONLY.
BHARAT PARENTERALS LIMITED.
ASIANA PHARMA.
10/10/2022
P.O.Box 265 Sana'a - Tel: +967-1-619174 - Fax: +967-1-619173
E-mail: sbdma@y.net.ye www.sbd-ye.org

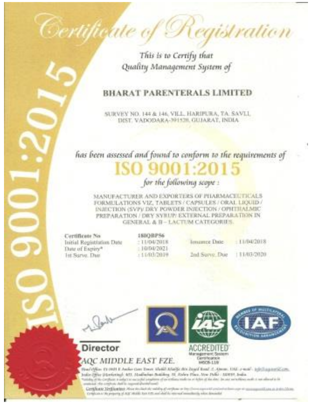

KINGDOM OF CAMBODIA
NATION RELIGION KING
MINISTRY OF HEALTH
PHARMACEUTICAL MANUFACTURER REGISTRATION LICENSE
Bharat Parenterals Limited
Survey No.144 & 146, Jarod Samlaya Road, Tal-Savli, Vill. Haripura, Dist. Vadodara, Gujarat State, India
Director General for Health

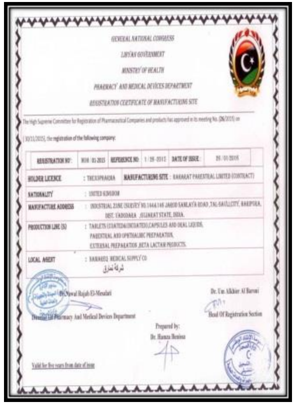

FORM I. R.
CERTIFICATE OF INCORPORATION
निगमन का प्रमाण-पत्र
I HEREBY CERTIFY THAT BHARAT PARENTERALS LIMITED
IS THIS DAY INCORPORATED UNDER THE COMPANIES ACT, 1956 (NO. 1 OF 1956) AND THAT THE COMPANY IS LIMITED.
GIVEN UNDER MY HAND AT AHMEDABAD THIS THIRD DAY OF DECEMBER ONE THOUSAND NINE HUNDRED NINETY TWO.
Registrar of Companies, Gujarat, Dadra & Nagar Haveli


FORM 26
(See Rules 73 and 83)
Certificate of Renewal of Licence to manufacture for sale of Drugs Other than those specified in Schedule X
Certified that Licence in Form 25 No:G/1269 Granted on the Date: 30/03/1995
Form 28 No:G/925 Granted on the Date: 30/03/1995
To M/s. BHARAT PARENTERALS LIMITED
for the manufacture of the following categories of drugs being drugs other than those specified in Schedule C and C (1) and "X" to the Drugs and Cosmetics Rules, 1945 drugs Specified in Schedule C and C (1) excluding those specified in Schedule "X" to the Drugs and Cosmetics Rules, 1945,
at the premises situated at SURVEY No. - 144 & 146, JAROD SAMLAYA ROAD, VILLAGE - HARIPURA, TAL. - SAVLI, DIST - VADODARA - 391 520.
has been renewed from : 01/01/2018 To 31/12/2022
Name(s) of drugs (each item to be separately specified) : As per list Approved & Annexed
Names of approved competent technical staff : As per list Approved & Annexed.
Signature
Designation
(Dr. H. G. KOSHIA)
Commissioner
Food & Drugs Control Administration
Gujarat State.
Date:- 16/05/2018


COMMISSIONER,
FOOD & DRUGS CONTROL ADMINISTRATION,
BLOCK NO-8, 1st FLOOR, DR.JIVRAJ MEHTA BHAVAN, GANDHINAGAR, GUJARAT STATE
No. Cert/GLP/BPL/2017/SC-1/ 05025 2 /B
Date: 06 MAY 2017
To,
M/s. BHARAT PARENTERALS LIMITED,
Survey No. 144 & 146,
Jarod-Samlaya Road,
Tal. Savli, Haripura,
Dist. Vadodara,
Gujarat State, India.
SUB:- Drugs & Cosmetics Act,1940 & Rules thereunder. Issue of Good Laboratory Practices Certificate.
Dear Sir,
REF:- Your letter Dated.03/04/2017.
I send herewith Good Laboratory Practices (GLP) Certificate in one copy as desired by you.
Yours faithfully,
For Commissioner,
Food & Drugs Control Administration.
Gandhinagar, Gujarat State
1) Copy to The Assistant Commissioner, (Vadodara Circle) Vadodara.
2) Copy to G-Branch for information.


Food & Drugs Control Administration
BLOCK NO. 8, 1st FLOOR, Dr. JIVRAJ MEHTA BHAVAN
GANDHINAGAR, GUJARAT STATE, INDIA PIN: 382010
No.Cert/GLP/BPL/2017/SC-1/ 050251 /B, Date. 06 MAY 2017
G.L.P. CERTIFICATE
This is to certify that M/s. BHARAT PARENTERALS LIMITED, Survey No. 144 & 146, Jarod-Samlaya Road, Tal. Savli, Haripura, Dist. Vadodara is holding drugs manufacturing license in form No. 25 & 28, bearing No. G/25/1269 & G/28/925 under Drugs & Cosmetics Act, 1940 & Rules thereunder.
The firm is having its own testing laboratory as per Drugs & Cosmetics Act, 1940. The firm is complying with the Good Laboratory Practices (GLP) requirements of Schedule L1 of the Drugs and Cosmetics Rules 1945.
This certificate is issued on the basis of the inspection report and documents available in this office as on today and issued for Export & participation in tender for State and Central Governments.
This Certificate is valid up to: 06/05/2017 to 05/05/2019.
For Commissioner,
Food & Drugs Control Administration.
Gandhinagar, Gujarat State

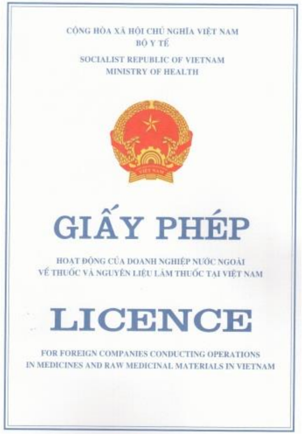

Certificate of Registration
This is to Certify that
Quality Management System of
BHARAT PARENTERALS LIMITED
SURVEY NO. 144 & 146, VILL. HARIPURA, TA. SAVLI, DIST. VADODARA-391520, GUJARAT, INDIA
has been assessed and found to conform to the requirements of
ISO 9001:2015
for the following scope :
MANUFACTURER AND EXPORTERS OF PHARMACEUTICALS FORMULATIONS VIZ. TABLETS / CAPSULES / ORAL LIQUID / INJECTION (SVP)/ DRY POWDER INJECTION / OPHTHALMIC PREPARATION / DRY SYRUP/ EXTERNAL PREPARATION IN GENERAL & B - LACTUM CATEGORIES.
Certificate No : 18IQRP56
Initial Registration Date : 11/04/2018
Date of Expiry* : 10/04/2021
1st Surv. Due : 11/03/2019
Issuance Date : 11/04/2018
2nd Surv. Due : 11/03/2020
ISO 9001:2015
Director
AQC MIDDLE EAST FZE.
IAS ACCREDITED
IAF

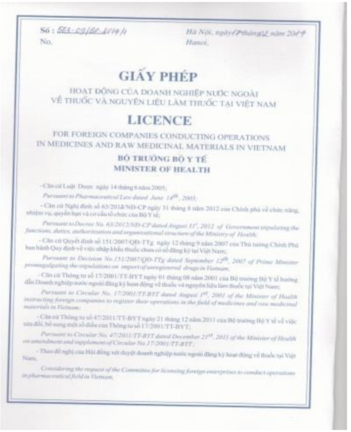

GENERAL NATIONAL CONGRESS
LIBYAN GOVERNMENT
MINISTRY OF HEALTH
PHARMACY AND MEDICAL DEVICES DEPARTMENT
REGISTRATION CERTIFICATE OF MANUFACTURING SITE
HOLDER LICENCE : THESPIAPHARMA
MANUFACTURING SITE : BHARAT PARENTERAL LIMITED (CONTRACT)
NATIONALITY : UNITED KINGDOM
MANUFACTURE ADDRESS : INDUSTRIAL ZONE (SURVEY NO.144&146 JAROD SAMLAYA ROAD, TAL SAVLI,CITY, HARIPURA, DIST. VADODARA, GUJARAT STATE, INDIA.
PRODUCTION LINE (S) : TABLETS (COATED&UNCOATED), CAPSULES AND ORAL LIQUID, PARENTERAL AND OPTHALMIC PREPARATION, EXTERNAL PREPARATION, BETA LACTAM PRODUCTS.
LOCAL AGENT : NAMAREQ MEDICAL SUPPLY CO.
Head Of Registration Section
Prepared by:
Valid for five years from date of issue

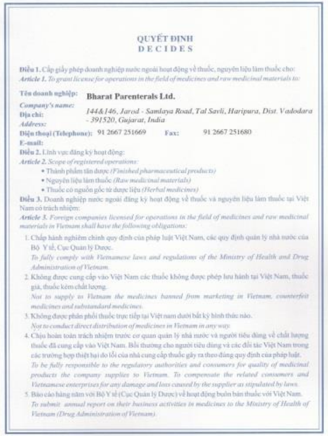

NATIONAL AGENCY FOR FOOD AND DRUG ADMINISTRATION AND CONTROL
NAFDAC/DER/HQ/VOL.2/66
18th February, 2016
The Chief Executive Officer
Bharat Paranterals Limited
Vill. Haripura, Ta. Savli,
Dist Vadodara
Gujarat
India
GOOD MANUFACTURING PRACTICE (GMP)
This is to certify that the GMP assessment of your Factory by Inspectors from National Agency for Food and Drug Administration and Control (NAFDAC) Nigeria on 4th December, 2015 was Satisfactory.
The product line inspected was:
- Oral Solids Dosage Forms (Tablets, Capsules)
- Liquid Dosage Form
- Semi Solid Dosage Forms
- Injectables (Parenterals)
The Products are:
1. Amovin 1.2g Injection
2. Biomedar 525 Tablet
3. Biomoxil
4. Ascorbal Suspension
5. Biomestatin Tablet
6. Nexthor Injection
You are to ensure that this standard is consistently maintained.
Mrs. Titilope O. Owolabi
For: Director-General (NAFDAC)


PHARMACY, MEDICINES & POISONS BOARD
ALL CORRESPONDENCE SHOULD BE ADDRESSED TO THE REGISTRAR
Telephone: (265) 01 755 165/01 755 166
Fax: (265) 01 755 204
P.O. Box 30241
Capital City
Lilongwe 3
Malawi
Ref: PMPB/CGMP/VOL IV/75
3rd May, 2013
The Managing Director,
Bharat Parenterals Ltd,
Shree Avenue, First Floor,
VIP Road, Karelibaug,
Vadodara-390018
INDIA.
Dear Sir,
CONFIRMATION OF REGISTRATION FOR BHARAT PARENTERALS LIMITED.
This letter serves to confirm that Bharat Parenterals Limited at Vadodara in India is registered as a pharmaceutical manufacturer with the Pharmacy Medicines and Poisons Board (PMPB).
The registration is effective on 1st June, 2012 and is valid for three (3) years.
Yours Sincerely,
G. Kadewele
REGISTRAR

Certificates







Bharat Parenterals Limited

Jarod Samlaya Road,
Vill.Haripura,Vadodara-391520
Gujarat, India

02667-251680

Send us your queries at -
Info@bplindia.in