

Food & Drugs Control Administration

BLOCK NO. 8, 15+ FLOOR, Dr. JIVRAJ MEHTA BHAVAN, GANDHINAGAR, GUJARAT STATE, INDIA PIN: 382010



Certificate No.: S-GMP/20051984

G.M.P. CERTIFICATE

This is to certify that M/s. PRUDENCE PHARMA CHEM, PLOT NO:7407, B/H. LYKA LAB., G.I.D.C. IND. ESTATE, ANKLESHWAR- 393 002, Dist.- BHARUCH is holding valid drug manufacturing licenses in Form No. 25 bearing No. G/1656 issued by this administration under the provisions of Drugs & Cosmetics Act 1940 & Rules thereunder. Under the said licenses the firm is permitted to manufacture & sell drugs covered under the following categories.

	Dosage Form (s)	Category (ies)
Bulk Drugs (APIs)		General
Library		

The firm has employed competent technical staff to undertake manufacturing & testing of the permitted drugs. They are following GOOD MANUFACTURING PRACTICES in manufacturing and testing as laid down under the REVISED SCHEDULE - M of Drugs & Cosmetics Act 1940 & Rules There under.

The manufacturing plant is subjected to inspection at suitable intervals by competent authority.

DAUGS CO.

This certificate is valid from Dt: 19/05/2020 to 18/05/2022.

(Dr. H. G.KOSHIA)

Commissioner
Food & Drugs Control Administration
Gandhinagar, Gujarat State

Email : comfdca@gujarat.gov.in

Phone : 91-79-23253417, Fax: 91-79-232-53400

MANGESH SHIVTARKAR Accounts Assistant ORICON HOUSE.

ORICON HOUSE.

OTH FLOOR, 12, K.

DUBHASH MARG,
FORT, MUMBAI400 001.

ATTESTED

1 5 JUL 2020



H. CHOWE HARY
PUBLIC NOTARY
(GOVT OF INDIA)

1 F. SEP, 2020



भारत सरकार GOVERNMENT OF INDIA अपोस्टिल / APOSTILLE

(Convention de La Haye du 5 octobre 1961)

Country

REPUBLIC OF INDIA

This public document

COMMERCIAL DOCUMENT
has been signed by H G KO

has been signed by H G KOSHIA

acting in the capacity of COMMISSIONER

bears the seekstamp of ASIAN EXPORTERS CHAMBER OF COMMERCE & INDUSTRY, MUMBAI

Certified

at NEW DELHI, INDIA the 21-Sep-2020

by SO (OI/Attestation) MINISTRY OF EXTERNAL AFFAIRS

No MHMC0003960720

Seal / Stamp

Signature

is issued to PRUDENCE PHARMA CHEM



(अनाल बनाप) (SUNIL CHANAP) अनुभाग अधिकारी (ओ आई) Section Officer (OI) भी. पी. दी. प्रमाग / C.P.V. D.vie विवेश मंत्रालय, नई दिल्ली Ministry of External Affairs New



ATTESTED

SWARN LATA EXECUTIVE DIRECTOR

ASIAN EXPORTERS CHAMBER OF COMMERCE AND INDUSTRY
NAVI MUMBAI, INDIA

Ref No.

23782/20

FORM 25

(See Rule 70)

Licence to manufacture for sale (or for distribution) of drugs other than those specified in Schedule C, C(1) and X

Number of licence and date of issue G/25/1656 Date- 08/04/2015

1. <u>I. MR. R.D. CHODVADIYA, 2. MR. V. G. CHODVADIYA, 3. MR. N. P. CHODVADIYA, 4. MR. G. D. CHODVADIYA, 5. MR. V. N. CHODVADIYA, PARTNERS</u> of M/s. <u>PRUDENCE PHARMA CHEM</u> is hereby licensed to manufacture the following categories of drugs being drugs other than those specified in Scheduled C, C(1), and X to the Drugs & Cosmetics Rules, 1945, on the premises situated at <u>PLOT No. - 7407, B / h. LYKA LAB., G.I.D.C. IND. ESTATE, ANKLESHWAR, DIST. - BHARUCH - 393 002.</u>

under the direction and supervision of the following (competent technical staff):-

(a) Competent technical staff (Names)

As Per List Approved & Annexed As Per List Approved & Annexed

(b) Names of Drugs (each item to be separately specified)

- 2. The licence authorizes the sale by way of wholesale dealing and storage for sale by the licensee of the drugs manufactured under the licence, subject to the conditions applicable to licence for sale.
- 3. The licence shall be in force From: 08/04/2015 To 07/04/2020

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4. The licence is subject to the conditions stated below and to such other conditions as may be specified in the rules for the time being in force under the Drugs and Cosmetics Act, 1940.

ORICON HOUSE, 6TH FLOOR, 12, K. DUBHASH MARG, FORT, MUMBAI-400 001.

* Date: 08/05/2015

Commissioner
Food & Drugs Control Admini

Food & Drugs Control Administration Gujarat State.

Conditions of Licence

- This licence and any certificate of renewal in force shall be kept on the approved premises and shall be produced at the request of an Inspector appointed under the Drugs and Cosmetics Act, 1940.
- Any change in the (competent technical staff) named in the licence shall be forthwith reported to the Licensing Authority.
- If the licensee wants to manufacture for sale additional items of drugs not included above he should apply to the Licensing Authority for the necessary endorsement as provided in Rule 69(5). This licence will be deemed to extend to the categories so endorsed.
- 4. The licensee shall inform the Licensing Authority in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence shall be deemed to be valid for a maximum period of three months from the date on which the change takes place unless, in the meantime, a fresh licence has been taken from the Licensing Authority in the name of the firm with the changed constitution.

FDCA/GUJARAT/25/899

ATTESTED

MANGESH SHIVTARKAR Accounts Assistant

KAR 1 5 JUL 2020



ATTESTED

M. H. CHOWDHARY PUBLIC NOTARY (GOVT OF INDIA) 1 8 SEP 2020



भारत सरकार GOVERNMENT OF INDIA अपोस्टिल / APOSTILLE

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COMMERCIAL DOCUMENT

has been signed by H G KOSHIA

acting in the capacity of COMMISSIONER

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COMMERCE & INDUSTRY, MUMBAI

Certified

NEW DELHI, INDIA the 21-Sep-2020

by SO (Ol/Attestation) MINISTRY OF EXTERNAL AFFAIRS

No. MHMC0003960620

Seal / Stamp

is issued to PRUDENCE PHARMA CHEM



(सुनील चनाप) (SUNIL CHANAP) अनुभाग अधिकारी (ओ आई) Section Officer (OI) सी. पी. वी. प्रभाग / C.P.V. D.VI विदेश मंजालय, नई दिल्ली Ministry of External Affairs New



ATTESTED

SWARN LATA EXECUTIVE DIRECTOR ASIAN EXPORTERS CHAMBER OF COMMERCE AND INDUSTRY NAVI MUMBAI, INDIA

Ref No.

RETENTION OF LICENCE

Retention of Licence to manufacture for sale (or for distribution) of drugs other than those specified in Schedule C, C(1) and X

Certified that Licence in Form 25, No: G/25/1656, Granted on the Date: 08/04/2015

To M/s. PRUDENCE PHARMA CHEM

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 at the premises situated at PLOT No. - 7407, B / h. LYKA LAB., G.I.D.C. IND. ESTATE, ANKLESHWAR - 393 002, DIST. - BHARUCH.

has been retained from: 08/04/2020 To 07/04/2025

Name(s) of drugs as retained (each

item to be separately specified)

Names of approved competent

technical staff as retained

As per list Approved & Annexed

ALTOCAL AND

As per list Approved & Annexed

ORICON HOUSE.

ORICON HOUSE.

OTH FLOOR, 12, K.

DUBHASH MARG.
FORT, MUMBA!

400 001.

Signature:

Designation:

(Dr. H. G. KOSHIA) Commissioner,

Food & Drugs Control Administration, Gujarat State

Date: 22/04/2020

Note: You are requested to apply for the Retention of the above licence in 3 months before its Validity Expires.

EXECUTIVE DIRECTO
ASIAN EXPORTERS CHAMBER OF TO VIN

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ATTESTED

Margesh

Accounts Assistant

15 JUL 2020







SWARN LATA
EXECUTIVE DIRECTOR
ASIAN EXPORTERS CHAMBER OF COMMERCE AND INDUSTRY
NAVI MUMBAI, INDIA

Ref No.

3784/20

ATTESTED



MFG. & EXP. : Bulk Drugs & Drugs Intermediates

List of products specified in other than C,C1,& X to be manufactured under license in Form No. 25, G/1656 Renewal for period: <u>08.04.2020 to 07.04.2025</u>.

C	T				A 1 - 4 - 1
Sr. No	No.	Name of Products	Pharmacopeias/ Specification	For DOMESTIC /EXPORT	
1	A	GLIBENCLAMIDE	BP	EXPORT	4
	В	GLIBENCLAMIDE	EP	EXPORT	. 4
	С	GLIBENCLAMIDE [GLYBURIDE]	USP	EXPORT	
	D	GLIBENCLAMIDE	IP	DOMESTIC	192 F
2	А	AMLODIPINE BESILATE	BP	EXPORT	
	В	AMLODIPINE BESILATE	EP	EXPORT	
	C	AMLODIPINE BESYLATE	USP	EXPORT	
	D	AMLODIPINE BESYLATE	IP	DOMESTIC	
3	Α	GLIMEPIRIDE	T		
	В	GLIMEPIRIDE	BP	EXPORT	
	C	GLIMEPIRIDE	EP	EXPORT	
7	D	GLIMEPIRIDE	USP	EXPORT	
		GLIMEPIRIDE	IP	DOMESTIC	
ı	Α	S-AMLODIPINE BESYLATE [=LEVAMLODIPINE BESYLATE]	-	EXPORT	DRUGS CONTROL
	В	S-AMLODIPINE BESYLATE [=LEVAMLODIPINE BESYLATE]	IP	DOMESTIC	I SIMILIMON
	Α ,	CELECOXIB		1/3/200	PAT STATE
	B	CELECOXIB	BP	EXPORT / DOMESTIC	MAGARO
	С			EXPORT	lete the
	C	CELECOXIB	USP	EXPORT	
	А	ETORICOXIB	-	FYPORT	ON HOUSE, DOR, 12, K.
	В	ETORICOXIB		DOMESTIC 6TH FL	OOR, 12, K.
		7000		FORT	MUMBAI-
	A	SILODOSIN	JP		2001
	В	SILODOSIN		DOMESTIC	The state of the s
der	taking	9	1		~

The above is the only product to be manufactured by us at present, any addition there to or deletion there from shall not be done without the prior permission of the Commissioner, Food & Drugs Control Administration, Gujarat State, Gandhinagar-India

We further declare that this products are safe for use.

Accounts Assistant

Commissioner

Food & Drugs Control Administration Guiarat State, Gandhinagac

區 8 APR 2020 本 至 7 APR 2025



PUBLIC NOTARY (GOVT OF INDIA)

1 8 SEP 2020

भारत सरकार GOVERNMENT OF INDIA अपोस्टिल / APOSTILLE

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No. MHMC0003960820

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(SUNIL CHANAP) अनुभाग अधिकारी (ओ आई) Section Officer (OI) सी. ची. ची. प्रभाग / C.P.V. D.VI® विदेश मंत्रालय, नई दिल्ली Ministry of External Affairs NAW



ATTESTED

EXECUTIVE DIRECTOR

ASIAN EXPORTERS CHAMBER OF COMMERCE AND INDUSTRY NAVI MUMBAI, INDIA