

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/1392 Granted on the Date: 18/12/1996

To M/s. VAPI CARE PHARMA PVT. LTD.

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. - 225 / 3, Nr. MORARJI CIRCLE, G.I.D.C., VAPI, DIST. - VALSAD

has been retained from: 01/01/2022 To 31/12/2026

Name(s) of drugs as retained (each item to be separately specified) : As per list Approved & Annexed
Names of approved competent technical staff as retained : As per list Approved & Annexed



Signature :

Designation :

(Dr. H. G. KOSHIA)
Commissioner,

Food & Drugs Control Administration,
Gujarat State

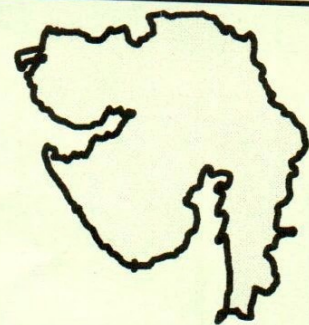
Date: 09/02/2022

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



Food & Drugs Control Administration

BLOCK NO. 8, 1ST FLOOR, Dr. JIVRAJ MEHTA BHAVAN,
GANDHINAGAR, GUJARAT STATE, INDIA. PIN : 382010



Certificate No. : **21102997**

Briyo Supplements LLP

On the basis of the inspection carried out on 04 - 05/10/2021 & 21/10/2021 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.

Proprietary Document | For Viewing Only

1 Name & Address of site: **VAPI CARE PHARMA PVT. LTD.**
PLOT NO. - 225 / 3, G.I.D.C., NEAR MORARJI CIRCLE,
CITY. - VAPI - 396 195, DIST. - VALSAD
GUJARAT STATE, INDIA

2 Manufacturer's Licence number : **G/1392** **G/1024**

3 Table : 1

Dosage Form (s)	Category (ies)	Activity (ies)
Tablet, Capsule, Dry Syrup, Oral Liquid, External Preparation (Ointment), ORS	General	Manufacturer
Tablet, Capsule, Dry Syrup	Beta-lactum	

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 **24/10/2024** 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

Format of this certificate is as per WHO TRS No. 908 of 2003.

Address of certifying authority

Food & Drugs Control Administration, Block
No. 8, 1ST floor, Dr. Jivraj Mehta Bhavan,
Gandhinagar, Gujarat State, India. - Pin :
382010

Name & function of : (Dr. H.G. KOSHIA)
responsible Person Commissioner

Email : comfdca@gujarat.gov.in

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

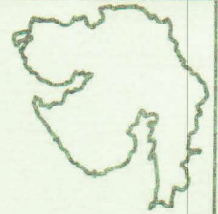
Date : 25/10/2021





Food & Drugs Control Administration

BLOCK No. 8, 1st FLOOR, Dr. JIVRAJ MEHTA BHAVAN,
GANDHINAGAR, GUJARAT STATE, INDIA. PIN : 382010



Certificate No. : **24075133**

On the basis of the inspection carried out on **01 - 02/07/2024 & 16/07/2024** 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 & Address of site : **VAPI CARE PHARMA PVT. LTD.**

PLOT NO. - 225 / 3, G.I.D.C., NEAR MORARJI CIRCLE,
CITY - VAPI - 396 195, DIST. - VALSAD,
GUJARAT STATE, INDIA

2 Manufacturer's Licence
number :

G/25/1392

G/28/1024

3 Table : 1

Dosage form (s)	Category (ies)	Activity (ies)
Tablet, Capsule, Dry Syrup, Oral Liquid, External Preparation (Ointment), ORS	General	Manufacturer
Tablet, Capsule, Dry Syrup	Beta-lactam	

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 **29/07/2027**. 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.

Format of this certificate is as per WHO TRS No. 908 of 2003.

Address of certifying authority

Food & Drugs Control Administration, Block
No. 8, 1st floor, Dr. Jivraj Mehta Bhavan,
Gandhinagar, Gujarat State, India - Pin :
382010

Email : comfdca@gujarat.gov.in

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

Date : 30/07/2024

Name & function of : **(Dr. A. G. KOSHIA)**
responsible Person : **Commissioner**





Vapi Care Pharma Pvt Ltd
Plot No. 225/3, GIDC, near Morarji Circle
Vapi, Gujarat, 396195, INDIA

This certificate confirms that the above company is registered (as provided in registration details) with U.S Food and Drug Administration pursuant to The Public Health Security and Bioterrorism Preparedness and Response Act of 2002 (Bioterrorism Act), section 305, added section 415 to the Federal Food, Drug, and Cosmetic Act.

Registration Details

Registration Number: 10240331366
Certificate Validity: 5th April 2025

This certificate is NOT issued by the U.S Food and Drug Administration, and FDABasics is NOT affiliated with the U.S Food and Drug Administration. This certificate is issued as confirmation of FDA registration and U.S. Agent appointment. The purpose of the certificate is to provide documentary evidence of facility registration. This certificate does not represent any form of approval. It remains the responsibility of the Registrant (owner/operator/in charge of the facility) to comply with all applicable FDA requirements including renewal and compliance with GMP.



Jerry Doane

A handwritten signature in black ink, appearing to read 'Jerry Doane'.

Control Number: vapi10282024



Manufacturing Site Registration Certificate

It is certified that the following Manufacturing Site has been registered in the UAE Ministry of Health, in accordance with the Article 65 of the Federal Law No.3 of 1984

Certificate #	21262 -C 1	First Reg. Date	16/12/2020
Registration #	21262	Reg. Expiry Date	15/12/2025
Committee Meeting No.	43/2020	Meeting Date	16/12/2020
Payment Receipt No.	EPAY20200000023299	Payment Date	06/12/2020
Manufacturing Site Name	Vapi Care Pharma Pvt Ltd		
Address	Plot No: 225/3, Near Moraji Circle , Vapi, Gujarat, Post Code: 396195, Gujarat, India		
Activities Registered For			
Manufacturing/processing of starting material , Assembly , Manufacture of dosage forms , Packaging & Labeling , Storage & Handling , Laboratory Testing , Batch releaser (certification) , Secondary packaging			
Non Hazard Line(s) of Production Registered For			
<ul style="list-style-type: none">- Non-Sterile Products-Non Sterile Products (Dosage Forms) : Capsules, soft shell- Non-Sterile Products-Non Sterile Products (Dosage Forms) : Capsules, hard shell- Non-Sterile Products-Non Sterile Products (Dosage Forms) : Tablets- Non-Sterile Products-Non Sterile Products (Dosage Forms) : Suspension- Non-Sterile Products-Non Sterile Products (Dosage Forms) : Powders- Non-Sterile Products-Non Sterile Products (Dosage Forms) : Semi-Solids (Ointment)- Packaging and Labelling : Primary Containers- Secondary Packaging : Primary Containers- Non-Sterile Products-Batch certification only : Other non-sterile medicinal products			
Manufacturing Site for product Class(s)		GSL Products	
<div><div>1.</div><div>The evidence(s) of GMP for the above mentioned activity(s) is are acceptable.</div></div> <div><div>2.</div><div>Failure to provide current acceptable GMP or any equivalent evidence prior to the expiry / as & when requested could result in refusal to receive product registration dossier or removal of the affected company and its product(s) from the register, according to the case.</div></div> <div><div>3.</div><div>Registration as Manufacturing Site makes it eligible to involve in registered activities in respect to the products to be registered in the U.A.E.</div></div> <div><div>4.</div><div>The Manufacturing Site should apply for minor variation as and when an amendment is needed in registration status.</div></div> <div><div>5.</div><div>This registration applies only to the above site name and address.</div></div>			

Issued on: 16/12/2020

Pass Code: 132176



هذه الشهادة صادرها من وزارة الصحة ووقاية المجتمع وتعتبر من الوثائق الحكومية الرسمية ولا تحتاج إلى
توقيع، ويحظر تطبيقاً تقليدياً أو إدخال أي تعديلات عليها سواءً بالإضافة أو الحذف أو التغيير في بياناتها أو
غير ذلك من أنواع التعديل، وعند الشهادة لاغية إذا شابهها شيء من ذلك. للتأكد من صلاحية الشهادة يرجى
المسح الضوئي للرمز ثنائي الأبعاد.

This certificate is generated by MOHAP Drug Department, it is official government document and does not require signature, it is strictly prohibited to imitate or amend to this certificate. Any attempt to change, delete, erase or add will cause this certificate to be rendered invalid. To verify this certificate validity, please scan the QR Code.



Registration Expiration Date

2026-12-31

Last Updated

2025-02-25

Registration Status

VALID

Registration Status Reason

US Agent Confirmed Facility Assignment

Is this facility engaged in the manufacturing/processing, packing, or holding of food for hun
United States?

Yes

No

Are you a fishing vessel engaged in processing (21 CFR 1.226(f))?

Yes

No

Section 1: Type of Registration

Facility Location :

Foreign Registration

UPDATE OF REGISTRATION INFORMATION:

Registration Number: 10240331366

Are you the new owner of a previously registered facility?

Yes

No

Previous Owner's Title:

Previous Owner's Name :

Previous Owner's Registration Number :

Section 2: Facility Name/Address Information

Facility Name

Vapi Care Pharma Pvt Ltd

Facility Name Suffix

Other

Telephone Number

091 997 8210829

Fax Number

RETENTION OF LICENCE

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

Certified that Licence in Form **28**, No: **G/1024** Granted on the Date: **18/12/1996**

To M/s. **VAPI CARE PHARMA PVT. LTD.**

for the manufacture of the 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 **PLOT No. - 225 / 3, Nr. MORARJI CIRCLE, G.I.D.C., VAPI, DIST. - VALSAD**

has been retained from: **01/01/2022 To 31/12/2026**

Name(s) of drugs as retained (each item to be separately specified) : **As per list Approved & Annexed**
Names of approved competent technical staff as retained : **As per list Approved & Annexed**



Signature :

Designation :

(Dr. H. G. KOSHIA)
Commissioner,

Food & Drugs Control Administration,
Gujarat State

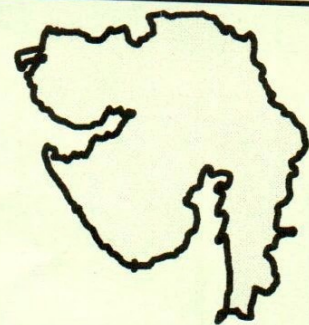
Date: **09/02/2022**

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



Food & Drugs Control Administration

BLOCK NO. 8, 1ST FLOOR, Dr. JIVRAJ MEHTA BHAVAN,
GANDHINAGAR, GUJARAT STATE, INDIA. PIN : 382010



Certificate No. : **21102997**

Briyo Supplements LLP

On the basis of the inspection carried out on 04 - 05/10/2021 & 21/10/2021 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 & Address of site: **VAPI CARE PHARMA PVT. LTD.**

**PLOT NO. - 225 / 3, G.I.D.C., NEAR MORARJI CIRCLE,
CITY. - VAPI - 396 195, DIST. - VALSAD
GUJARAT STATE, INDIA**

2 Manufacturer's Licence
number :

G/1392

G/1024

3 Table : 1

Dosage Form (s)	Category (ies)	Activity (ies)
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This certificate remains valid until **24/10/2024** 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

Format of this certificate is as per WHO TRS No. 908 of 2003.

Address of certifying authority

Food & Drugs Control Administration, Block
No. 8, 1ST floor, Dr. Jivraj Mehta Bhavan,
Gandhinagar, Gujarat State, India. - Pin :
382010

Name & function of : (Dr. H.G. KOSHIA)
responsible Person Commissioner

Email : comfdca@gujarat.gov.in

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

Date : 25/10/2021



Vapi care licenses Food & Drugs Control Administration

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Office of the Commissioner

Block No. 8, 1st Floor, Dr. Jivraj Mehta Bhavan, Gandhinagar, Gujarat State

ADDITIONAL PRODUCT PERMISSION

To,

VAPI CARE PHARMA PVT LTD**PLOT NO 225/3, NR. MORARJI CIRCLE, G.I.D.C., VAPI, DIST.- VALSAD - 396 195**Reference : Your Application Inward ID: **650768** Dated : **17-Oct-2022** (Reg ID : **625717**)

With reference to your Inward application, this is to inform you that your said application is considered & following **PRODUCT PERMISSION** has been granted, under the **LICENSE NO. G/28/1024 IN FORM NO. 28**.

Product Section : Tablets, Product Sub Section : General

Sr. No.	Name of Drugs																														
1	Prod ID : 825577 Permission Date : 17-Oct-2022 Type : Normal Permission Purpose: G-General Generic Name : Cholecalciferol Chewable Tablets 2000IU Brand Name : - Composition Title : Each Uncoated Chewable Tablet Contains:- <table><tr><th>Composition</th><th>Ingredients</th><th>Standards</th><th>Strength</th><th>UOM</th><th>Equivalent to</th></tr><tr><td>API</td><td>Cholecalciferol (as Stabilized)</td><td>IP</td><td>2000</td><td>International Unit (IU)</td><td></td></tr><tr><td>Color</td><td>Sunset yellow FCF</td><td>--</td><td>0</td><td>-----</td><td>-----</td></tr><tr><td>Excipients</td><td>-</td><td>--</td><td>0</td><td>QS</td><td></td></tr><tr><td>Contents</td><td>Appropriate overage of Vitamins added</td><td>-----</td><td>0</td><td>-----</td><td>-----</td></tr></table>	Composition	Ingredients	Standards	Strength	UOM	Equivalent to	API	Cholecalciferol (as Stabilized)	IP	2000	International Unit (IU)		Color	Sunset yellow FCF	--	0	-----	-----	Excipients	-	--	0	QS		Contents	Appropriate overage of Vitamins added	-----	0	-----	-----
Composition	Ingredients	Standards	Strength	UOM	Equivalent to																										
API	Cholecalciferol (as Stabilized)	IP	2000	International Unit (IU)																											
Color	Sunset yellow FCF	--	0	-----	-----																										
Excipients	-	--	0	QS																											
Contents	Appropriate overage of Vitamins added	-----	0	-----	-----																										

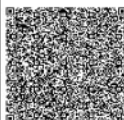
Terms and Conditions

- 1) Licensee should comply with all the provisions of Drugs & Cosmetics Act, 1940 & Rules 1945 as amended time to time.
- 2) Licensee should comply with all the provisions of Drugs (Price Control) Order 2013 as amended time to time (wherever applicable).
- 3) Licensee should abide by all the provision of Drugs & Magic Remedies (Objectionable Advertisement) Act, 1954 & Rules 1955 as amended up to date.
- 4) Licensee should not manufacture any drug by a name belonging to another manufacturer.
- 5) Licensee should not manufacture or sell drugs even if it is included in the approved list of product, if it is or as and when banned by Licensing Authority or Drugs Controller General of India or Government of India.
- 6) The permission is granted subject to the condition that, the product is safe for use in context of pharmaceutical Aids additives and excipients used in the formulation.
- 7) Any addition thereto or any deletion therefore will not be carried out without permission of Licensing Authority.
- 8) Above Product Permission is granted based on undertaking submitted by you with application.

(This Document is Digitally Signed)

Y G DARJI
(Deputy Commissioner)
For Commissioner

Food & Drugs Control Administration
Gujarat State, Gandhinagar



Reg ID : 625717

Doc ID: PP884200825577

VAPI CARE PHARMA PVT LTDLicense No : **G/28/1024**Print Date : **18/10/2022 11:48 AM**

Page 1 of 1

Signature valid

Digitally signed by
Date: 2022.10.18
11:52:09 +05:30
Reason: Sign
Location: AHD