

**INNOVATE  
IMPROVE  
INSPIRE**





## Company Profile



Since 1980, Octavius Pharma has been empowering lives through its high quality technologically driven diverse range of products. Over the years, Octavius has expanded its product portfolio & operations in different market segments worldwide.

As a **global leader in direct compressible granules**, we develop products at our state-of-the-art R&D facility as per latest Pharmacopoeia. Our global portfolio includes direct compressible granules, finished dosage formulations, herbal / ayurvedic / food supplements & herbal extracts.



## Key Strengths

- Formulation Development & Technology
- Direct Compressible Granules
- cGMP Manufacturing Facility
- Quality & Regulatory Systems

WHO-GMP Certified

**Food & Drugs Control Administration**  
BLOCK NO. 4, 1<sup>ST</sup> FLOOR, D. J. RAJ MEHTA BLDG,  
GANDHINAGAR, GUJARAT STATE, INDIA PIN - 382010

Certificate No: **L/19021258**

On the basis of the inspection carried out on **06/07/2018 & 26/12/2018** 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: **OCTAVIUS PHARMA PVT. LTD.**  
**MPGD AT - 507-8 TO 512, G.I.D.C. ESTATE,,**  
**City : WADHWAN CITY - 363 035, Dist. SUREDRANAGAR**  
**GUJARAT STATE, INDIA**

2. Manufacturer's Licence number: **G/25A/4401-A G/26A/5328-A**

3. Table 1

Dosage Form(s)	Category (s)	Activity (s)
Tablet, Capsule, Oral Liquid, Oral Dry Powder, Nasal Spray, Sterile Preparation	General	Manufacturer

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 **28/01/2022**. 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.

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

Address of certifying authority:  
Food & Drugs Control Administration,  
Block No. 4, 1<sup>ST</sup> Floor, Dr. Jeejeeb Mehta Bldg., Gandhinagar, Gujarat State, India - Pin - 382010

Name & function of responsible Person: **Dr. H.G. ANANDIA**  
Commissioner

Signature: 

Phone: 0179-2323417, Fax: 0179-232-4349

Date: **18/03/2019**



**“Quality is not an act, it is a habit”**

➤ **Direct Compressible Granules**

➤ **Herbal & Food Supplements**

➤ **Finished Dosage Forms**

(Tablets, Capsules, Cream, Ointment, Oral Powders,  
Dry Powder Inhalation, Nasal Spray/Drops)



## Facilities & Infrastructure

We have separate dedicated units for Pharmaceutical & Herbal/Ayurvedic products. The units are having global accreditation with huge capacities to ensure timely delivery of the products & are operated by skilled persons having vast experience.

Our state-of-the-art pharmaceutical manufacturing facility with stringent quality controls has been **WHO-GMP certified**. Our well developed infrastructure facilities with latest technology & experienced personnel caters to the upcoming needs of the industry.

### Facility 1 :

DC Granules &  
Finished Dosage  
Formulation

### Facility 2 :

Herbal, Ayurvedic &  
Food Supplements

## Quality

We have a modern & well-equipped Quality Control laboratory which has all necessary instruments like **HPLC Auto samplers, GC Auto samplers**, Stability chambers, IR Spectroscopy, UV Spectroscopy & many more for analysis of API, finished products, packaging & related materials used.

Our quality assurance team ensures that our products are pure, effective, safe & stable and are released only after thorough analysis.





## Research & Development

We always look for something new & innovative. We strive to create new market spaces through innovation & as a result have created a R&D centre which conforms to international quality standards.

It primarily focuses on differentiated portfolio to strengthen our competitive position. We use our insights to develop innovative products that offer **substantial benefits over existing products** for below dosage forms :

Oral  
Solids

Oral  
Liquids



## Regulatory

We have a dedicated team of professionals for Regulatory affairs with vast experience. We provide full support from the beginning of cooperation to the marketing authorization. We assist customers throughout entire registration lifecycle process & provide complete export documentation including **COPP, FSC & CTD dossier**.



## Global Presence



## Collaborate with us

- Product development
- In-licensing
- Contract manufacturing
- Customized products
- Technology transfer
- Marketing tie-ups

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