

ONLINE NATIONAL DRUGS LICENSING SYSTEM (ONDLS)

(WWW.STATEDRUGS.GOV.IN)

सी.डैक
CDAC

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Sign in

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GOVERNMENT OF INDIA

Online National Drugs Licensing System
(Central Drugs Standard Control Organisation)

ABOUT ANALYTICS SERVICE REGULATOR CONTACT US

Search License, Verify License Verify Track Application Sale | Manufacturing | Blood Centre | Certificate Login/Register

स्वास्थ्य एवं परिवार कल्याण मंत्रालय
MINISTRY OF
HEALTH AND
FAMILY WELFARE

CENTRAL DRUGS STANDARD CONTROL ORGANISATION
MINISTRY OF HEALTH, GOVERNMENT OF INDIA

World Health Organization

Search Pharmacy

Search Blood Center

APPLY FOR CERTIFICATE

NO CONVICTION COPP MARKET STANDING NEUTRAL CODE PRODUCTION WHO-GMP

235438 Licences Issued

147511 Approved Application

74992 Registered Firm

89892 Registered Technical Person

Statistics →

↑

Login Homepage

Search License, Verify License

Verify

Track Application

Sale | Manufacturing | Blood Centre | Certificate

Login/Register

Online National Drugs Licensing System

"ONDLS portal is a single window platform for online processing of various applications submitted by the applicants for issuance of manufacturing and sales licenses including Blood Centers and other certificates."

Statistics → 235438 Licences Issued

147511 Approved Application

74992 Registered Firm

89892 Registered Technical Person

Click here to log in

REGISTER OR SIGN IN

An OTP will be sent to your mobile number for verification

Enter your mobile number

GET OTP

By Sign In/Registration, I agree to the Terms of Service and Privacy Policy

OR

SIGNIN WITH USER NAME AND PASSWORD

Enter Registered Mobile Number here and alternatively login through User ID and Password

Manufacturer Dashboard (Site/Firm)

The screenshot shows the 'Manufacturer of Drugs' dashboard. At the top left is the Government of India logo and the text 'GOVERNMENT OF INDIA ONLINE NATIONAL DRUGS LICENSING SYSTEM'. At the top right are 'Manufacturer of Drugs' and 'Dashboard' buttons. A red box highlights the 'Click here for Fresh Application' button, which points to the 'Application Processing' module. The dashboard features several blue tiles: 'Product Management' (with 'Licenses' and 'Old Products Under Review' sub-tiles), 'Application Processing' (with 'Certificates' and 'Extension Form for Schedule-M' sub-tiles), 'Technical Member', 'Authorized Person', 'Old Licence Management' (with 'Firm Approval Listing' sub-tile), and 'Firm Approval Listing'. A note at the bottom states: 'Note :- For WHO-GMP & CoPP, first you need to Submit your Existing License through Old License Management Tile and get it approved from respective State FDA.'

Click here for Fresh Application

Product Management

Licenses

Old Products Under Review

Application Processing

Certificates

Extension Form for Schedule-M

Technical Member

Authorized Person

Old Licence Management

Firm Approval Listing

Manufacturer of Drugs

Dashboard

Note :- For WHO-GMP & CoPP, first you need to Submit your Existing License through Old License Management Tile and get it approved from respective State FDA.

Designed, Developed & Maintained by CDAC

Application Page

GOVERNMENT OF INDIA
ONLINE NATIONAL DRUGS LICENSING SYSTEM

Manufacturer of
Drugs

Dashboard

Applications

**Click here
for Fresh
Application**

Fresh Application Submission

Save as Draft

Submitted Applications

Approved Applications

Query Raised Applications

Rejected Applications

Post Approval Change Request Applications

**Request for Surrender/
Withdrawn**

Apply Retention/Renewal

Designed, Developed & Maintained by CDAC

Application Draft Page

Click here and select Application Type

Application Draft Initial Page

Click here and select Form Name

Application Type*

Issue of Certificate

Select Application Type
(Drug) Manufacturing License
Test License
Issue of Certificate

I agree that I will provide accurate information and I will be solely responsible for any information provided.

Form Name*

Select Form

WHO-GMP
Certificate of Pharmaceutical Product(COPP)
Free Sale Certificate(FSC)
Performance Certificate
Neutral Code
Market Standing Certificate

Subm

Note:- You can apply for fresh application here ONLY.

1. In case you want to retain old license then kindly submit your old license data through old license management tile to state FDA for data verification.
2. After old license data verified by FDA you may file for retention/renewal or endorsement or post approval.
3. No details will be changed or updated after successful submission of the application or old license data.

Firm Details Confirmation

Firm Detail Confirmation

Applicant Firm Details

Firm Name :- Anphar Organics Private Limited	Firm Constitution :- Private Limited
Firm Address :- Sidco Epip Kartholi Bari Brahma Jammu, Jammu, District-Jammu, State-Jammu And Kashmir, India -181133	
Site Type :- Own Site	Site Id :- JK0000553

Authorized Person

Select



Save Details

Select the Authorised Person and click on Save Details button

Application for COPP

Application for Certificate of Pharmaceutical Product (COPP)

Select From Below to Proceed with COPP application*

select

select

New Products For Which Inspection is Required

Old Products For Which Inspection is Not Required

Click here and select option as required

- **New Products For Which Inspection is Required – Products in which inspection was not carried out during the WHO-GMP inspection.**
- **Old Products For Which Inspection is Not Required – Products inspected during the WHO-GMP inspection but not applied for CoPP.**

Application for COPP

Application for Certificate of Pharmaceutical Product (COPP)

(Additional Products For Which COPP is Required)

Select Licence*

MLF25202

Select Product*

Gamma benzene hexachloride+Cetrimide In House Specification, 1.01% w,

Select Brand

DEMO(brand)



Add Brand

Select Country *

Afghanistan Albania American Samoa Angola |

Brand Will Reflect After Successfully Uploading of Form-51

Save

- Select License – All active licenses will be available here.
- Select Products – All products will be available for selected License.
- Select Brand – Registered Brand name will be shown here, if not registered you may register through clicking on Add Brand Button.
- Select Country – You can choose multiple countries, and the CoPP certificate will be generated separately for each product-country combination.

Add Brand Name

Application for Certificate of Pharmaceutical Product (COPP)

(Additional Products For Which COPP is Required)

Select Licence*

select

Select Product*

select

Select Brand

select

Select Country *



Brand Will Reflect After Successfully Uploading of Form-51

Click here for add Brand Name

Save

Product Brand Details

Product Brand Detail

Product Basic Details

Product number :- **10120240400056** Product Type :- **Single Ingredient**

Product Name :- **Clootrimazole -1.0% w/w I.P.** Dosage Name :- **Powder**

ONDLS Licence Number :- **MLF252024JK000005** Old Licence Number :- **JK/01/07-08/131**

Brand Form 51

Brand Name* **Brand Detail**

Save Brand Detail

Show 10 entries Search:

Select	Brand Name	Brand Detail	Action
No data available in table			

Generate Form 51 **upload Form**

Fill your Brand Name and Brand Details Here and Save

Generate Form 51

Product Basic Details

Product number :- **10120240400056** Product Type :- **Single Ingredient**

Product Name :- **Clotrimazole -1.0% w/w I.P.** Dosage Name :- **Powder**

ONDLS Licence Number :- **MLF252024JK000005** Old Licence Number :- **JK/01/07-08/131**

Brand Form 51

Brand Name* **Brand Detail**

Save Brand Detail

Show 10 entries Search:

Select	Brand Name	Brand Detail	Action
<input type="checkbox"/>	Test	Test	<input type="checkbox"/>

After generated Form 51 Selected Brand cannot Delete or Select

Select your Brand and Click here to Generate Form 51

Generate Form 51 **DownLoad Form 51** **upload Form**

Click here to Download Form 51

After download Form, Signed and sealed on it and upload Form from here

Upload Form 51

Upload Form 51

Upload Cover Letter

[Browse...](#) No file selected.

**Click here to upload
signed form 51 and Submit**

Submit

Application for COPP

Application for Certificate of Pharmaceutical Product (COPP)

(Additional Products For Which COPP is Required)

Select Licence*

MLF252024JK000005

Select Product*

Gamma benzene hexachloride+Cetrimide In House Specification, 1.01% w,

Select Brand

Select

 Add Brand

Select

DEMO(brand)

Save

Select Country *

After added Brand, Brand Name is showing against Selected Product

Application for COPP

Application for Certificate of Pharmaceutical Product (COPP)

(Additional Products For Which COPP is Required)

Select Licence*

select

Select Product*

select

Select Brand

select

 Add Brand

Select Country *

select

Brand Will Reflect After Successfully Uploading of Form-51

Save

Show 10 entries

Search:

Licence Number	Product Details	Brand Name	Country Name	Action
MLF252024JK000005	Gamma benzene hexachloride+Cetrimide-I.P. 1.0% w/v, I.P. 0.1% w/v(Lotion)	demo	Angola,American Samoa,Albania,Afghanistan	

Showing 1 to 1 of 1 entries

Previous 1 Next

← Back

Confirm and Proceed

Note :- You can also add Multiple products from multiple Licenses at a time.

After saved data is showing in below table. You can also delete any drafted Product

Click here Confirm and Proceed

Checklist for WHO-GMP

Checklist Document Validation and Upload Page

Show 10 entries

Search:

S.No	CheckList Item	Document Upload Status
1	* Application with covering letter on company's letter head duly signed and stamped by Authorized signatory indicating the name and designation of the authorized signatory along with the name and address of the firm	☒
2	* Site Master file (as specified under WHO TRS 961) including Manufacturing layout Schematic diagram of water system specifying circulation loop and MOC Schematic diagram of HVAC system specifying terminal filter configuration List of equipment and Instrument, List of major changes after last inspection	☒
3	* List of major changes after last inspection	☒
4	* Section applied for COPP as per WHO Certification Scheme	☒
5	* List of Products applied for issuance of Certificate of Pharmaceutical Product	☒
6	* Copy of Product permission for product(s) applied for COPP	☒
7	* List of Not of Standard Quality (NSQ) sample declared by State or Central Drugs Testing Laboratory	☒
8	* Export Data of last 3 years in case of revalidation	☒
9	* Application Form For Certificate of Pharmaceutical Product	☒
11	* Name of the Drug Substances: (INN, Chemical Name, Company Code, CAS) Chemical Structure, Molecular formula, Molecular Weight(For Bulk Drug)	☒

1 to 10

Previous 1 2 3 Next

Proceed

Upload all required checklist documents and Procced.

Checklist for WHO-GMP

Upload Checklist

Note: Please write NA, in case you do not have any remarks to enter.

Application with covering letter on company's letter head duly signed and stamped by Authorized signatory indicating the name and designation of the authorized signatory along with the name and address of the firm

No file selected.

Remarks:^{*}

Enter Remarks

Click here, Upload the Related document, also put your remarks and Submit

Checklist for WHO-GMP

20	* Analytical Method Validation / Verification Protocol of the each applied product	
21	* Analytical Method Validation / Verification Report of the each applied product	
22	* List of Reference/ working/ primary standards or materials	
23	* Stability study data report as per requirements mentioning batch size. (should be presented in tabular form with details of Batch No, Batch size, Date of Manufacturing, Date of initiation, Packaging details) Accelerated (6 months, Real Time (12 months), Forced degradation/stress studies (if applicable), Post approval Stability Protocol and Stability Commitment as per WHO TRS 1010 Annex 10	
24	* List of the major changes after last inspection conducted	
25	* Proposed storage statement and shelf-life	
26	* Product summary Sheet	

1 to 25

Previous 1 Next

Proceed

After uploading all required
Documents Click here to Proceed

Payment Challan upload

Payment Details

Note:

1. Fill inspection fee wherever necessary otherwise fill zero.

Purpose*

Challan Details

Challan No.*

Challan Date.*

Application Fee*

Inspection Fee*

Any Other Fee

Total Amount of Uploaded Challans*

Bank Name*

Branch Code *

Challan of Fees Paid To Be Upload*

Affidavit.pdf

Save & Proceed

**Upload Challan Details and click
on Here for Save & Proceed**

Application Preview

Application Preview

Corporate & Site Details

Firm Name :	ANPHAR ORGANICS PRIVATE LIMITED	Firm Address :	SIDCO EPIP KARTHOLI BARI BRAHMANA JAMMU, JAMMU, DISTRICT-JAMMU, STATE-JAMMU AND KASHMIR, INDIA -181133
CIN/PAN No :	AAECA5483G	Contact No :	1923220064
Firm Address Proof:	GSTN	Firm Address Proof Document:	View Document
Site Type :	OWN SITE	Site Id :	JK0000553
Fax No :	1923-220113		

Director Details

Name	Designation	Email Id	PAN NO	Joining Date	Status
SHAILENDER GUPTA	DIRECTOR	SHAILENDER@ANPHARORG.IN	AKAPG1996P	10/03/2015	ATTACHED
K NAGAPPAN	DIRECTOR	DRNAGAPPAN@YAHOO.COM	AABPN0303Q	07/08/2014	ATTACHED
K BALAKUMAR	DIRECTOR	VIJAY.BALAKUMAR@STERIL-GENE.COM	AACPB2388N	07/08/2014	ATTACHED
S MANOHAR	DIRECTOR	DRS.MANOHAR@HOTMAIL.COM	AATPM8720C	07/08/2014	ATTACHED

Application Preview

Authorised Person Details

Name	Designation	Email	Mobile No
Mr. Sandeep Mugutrao Jagtap	Plant Head		9622052465

Payment Details

Bank Name :- ABC	Amount :- 1000
Challan Date :- 2025-07-08 00:00:00.0	

WHO-Dosage Details

Show	10	entries	Search:
Licence Number	↑↓	Product Details	Brand Name
MLF252024JK000005	↑↓	Clotrimazole -I.P. 1.0% w/w(Powder)	Country Name
Showing 1 to 1 of 1 entries		Test	NA
Previous			1 Next

Final Submit

Click here for Final Submission

Final Application Submission

Your Application has been submitted successfully. Kindly note your
file no. **JK/WHOGMP/2025/00002** for future correspondence.

After Final Submission, You have received an Application Number

Submitted Application Status

Submitted Application

Show 10 entries Search:

File No.	Form Name	Applicant Address	State FDA Status	CDSCO HQ Status	Action
JK/COPP [REDACTED]	WHO-GMP	anphar organics private limited SIDCO EPIP Kartholi bari brahmana jammu, jammu, (India) - 181133	Submitted	Submitted	
JK/WHO-GMP/2025/00002 [REDACTED]					

Showing 1 to 2 of 2 entries

Action Button

- Generate Form
- Withdraw Application
- View Post Submission Change Request
- Apply Endorsement
- Post Submission Change Request

You can also track your Submitted Application.

Go to Application Processing Tile → Submitted Application and You can show your submitted application status as showing above image and also can take any step after click on action Button.

THANK YOU

