

e-Aushadhi



User Manual For Manufacturing Fresh License (ASU)

=====

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➤ Fresh License Application

Submitting a New License Application

After logging in and receiving approval, the system will redirect you to the **User Dashboard**.

To begin a new license application:

All New Licensee Can Apply for Fresh License

◆ From the **left-hand menu**, click on **Fresh License > New Application**.

◆ The **New Application** screen will appear.

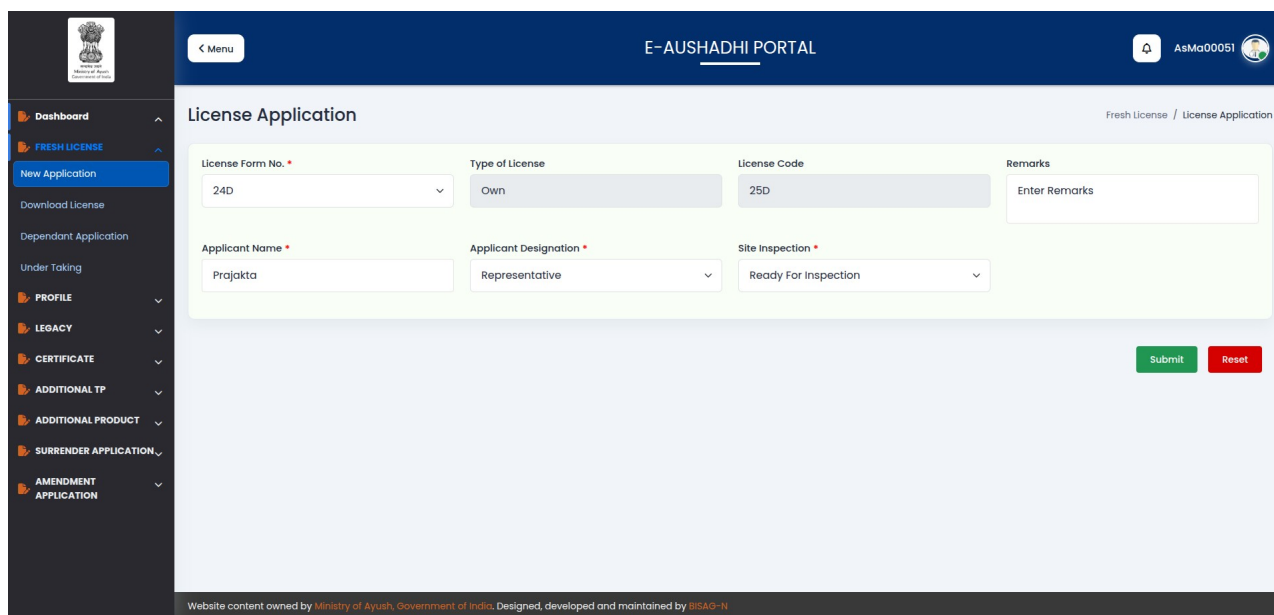
- Initially, the data table will be **empty**.

◆ Click on the **(plus) icon** to initiate a new application.

◆ You will be redirected to the **New Application Entry** page where you can begin entering the required details.

◆ If your license type is **Loan**, follow these steps to submit your application:

- Select your **Own Firm Name** and the relevant **License**.
- Enter the **OTP** sent to your own firm's registered email address.
- After verifying the OTP, you can submit the application.



The screenshot shows the 'License Application' form on the E-Aushadhi Portal. The form is titled 'License Application' and is part of the 'Fresh License / License Application' section. It contains the following fields:

- License Form No. ***: A dropdown menu with '24D' selected.
- Type of License**: A dropdown menu with 'Own' selected.
- License Code**: A text input field with '25D' entered.
- Remarks**: A text area with the placeholder 'Enter Remarks'.
- Applicant Name ***: A text input field with 'Prajakta' entered.
- Applicant Designation ***: A dropdown menu with 'Representative' selected.
- Site Inspection ***: A dropdown menu with 'Ready For Inspection' selected.

At the bottom right of the form, there are two buttons: 'Submit' (green) and 'Reset' (red). The footer of the page states: 'Website content owned by Ministry of Ayush, Government of India. Designed, developed and maintained by BISAG-N'.

Viewing Submitted Applications

After submitting your new license application:

- The application details will appear under the **New Application** tab within the **Fresh License** section on the left-hand menu.
- You can view the status, application number, and other submitted details in the table.
- If your application has any queries then Administration send back the application to you. You can see the reason by clicking on the Refer back button.
- When you resolved the query and then click on comply button to resubmit the application.
- If your application is rejected by the Administration, you must click on comply button to resubmit the application within **60 days**. If you fail to do so, your application will be **disabled**.
- If You enter incorrect details you can also delete the application.

Here, you can:

- **Edit** your previously entered information.
- Fill in additional sections such as:
 - **Product Details**
 - **QC Register**
 - **Other Mandatory Fields as required by the licensing authority**

The screenshot displays the 'e-Aushadhi Portal' interface. On the left is a sidebar menu with options like Dashboard, FRESH LICENSE, New Application, Download License, Dependant Application, Under Taking, PROFILE, LEGACY, CERTIFICATE, ADDITIONAL TP, ADDITIONAL PRODUCT, SURRENDER APPLICATION, AMENDMENT APPLICATION, and RETENTION APPLICATION. The main content area is titled 'Fresh License (Manufacture) Application - Ayurveda, Unani, Siddha, Sowa Rigpa'. It features a table with columns for various application details and a legend explaining the status indicators (orange for refer-back, black for restricted action).

- Ensure all information is complete and accurate before final submission to avoid delays in processing.

Adding Product Details

Product Management

Adding Product Details

When you click on the **Add Product** button, a new tab or form will open where you can enter product-specific information.

You will be required to fill in the following fields:

- **Product Name**
- **Product Type**
- **Category**
- Any other applicable product details as shown in the form

After filling in the product information:

- Click **Save** to add the product.
- The entered product will be displayed in the **Product Details** data table below the form.

You can add multiple products by repeating this process.

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Add Product

Fresh License / Add Product

Product Name *

Type *

--Select--

Category/Dosage Form *

--Select--

Sub Type *

--Select--

Permission Purpose *

--Select--

Remarks

Composition Title *

e.g. Each Gram Contents

Drug Categories Based On Their pharmacological Actions *

-- Select --

Diseases and Conditions *

Contains (Shang/Alfne/ganja/charas/Other Narcotics Ingredients/ Schedule E-1 Ingredients/Shelf Generated Alcohol)? *

☐ Yes ☐ No

Shelf Life in Month *

(This data is used for official information, not for verification)

☐ Regular As Per Clause ☐ Apply for access of shelf life

Note : Drug Categories Based On Their Pharmacological Actions and Diseases and Conditions are used for Office use Only.

Submit Reset

Product Details

10 Entries Per Page

Search:

Sr No.	Product Name	Type	Category	Sub Type	Status	Edit	Composition	Attachments	AIF	Delete
1	vxv	Ayurvedic Classical Medicine	Anjana	Regular	Pending					

Showing 1 to 1 of 1 entry

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Completing Product Details:-

- After **adding and saving** a product entry, it will appear in the **Product Details** data table.
- In this table, you are required to complete additional fields for each product, such as:

Produc Composition

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Add Composition

Fresh License / Product Data / Add Composition

Ingredient Type *

--Select--

Ingredient Name *

Botanical Name *

Form of Ingredients *

--Select--

Reference *

--Select--

Part Used *

--Select--

Quantity *

Measuring Units *

--Select--

Submit Reset

Composition

10 Entries Per Page

Search:

Sr No.	Ingredient Name	Botanical Name	Ingredient Type	Ingredient Form	Reference	Part Used	Quantity	Measuring Units	Edit	Delete
No data available in table										

Showing 0 to 0 of 0 entries

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◆ Product Attachments



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Fresh License / Product Data / Add Attachment

Attachment List

(1)	Draft Lable Annexure 2 As per D and C Rule 161 *	Choose file	20240528150_F_sample.pdf
(2)	Annexure 3 Master Formula card with manufacturing SOP *	Choose file	20240528150_F_sample.pdf
(3)	Annexure 4 Quality control specification *	Choose file	20240528150_F_sample.pdf
(4)	Annexure 5 Proof of effectiveness With Reference Page As per Rule 158 B *	Choose file	20240528150_F_sample.pdf
(5)	D and C Act 1940 schedule I Book Reference of Ingredinetns Or Formulation As per D and C Rule 158 B *	Choose file	20240528150_F_sample.pdf
(6)	Safety study data As per D and C Rule 158 B	Choose file	No file chosen
(7)	Shelf life or date of expiry of medicines *	Choose file	20240528150_F_sample.pdf
(8)	Validated data As per D and C Rule 161 B *	Choose file	20240528150_F_sample.pdf
(9)	Extra Field	Choose file	No file chosen

Submit Reset

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◆ Product AIF (Additional Information Form)

e-Aushadhi Portal

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Fresh License / Product Data / Additional Information Form

Additional Information Form

(1)	Has the proposed product's category and dosage been approved? *	No
(2)	For this proprietary medicine, are there any commercially available products with a similar name or a similar qualitative and quantitative composition? *	No
(3)	Whether the unit has fulfilled all new requirement according to the schedule? *	No
(4)	Does the firm have an in-house laboratory facility? *	No
(5)	Are all the necessary facilities and equipment available for the complete quality control testing of the product? *	No
(6)	Has the manufacturing unit complied with all new requirements as specified in the applicable schedule? *	No
(7)	What is the therapeutic use of the additional product, or what disease is it intended to treat? *	please Specify
(8)	Does the parent firm hold a valid license for the product category specified in this application? *	No
(9)	Will the additional product be marketed through promotional media such as newspapers, television, radio, or print advertisements (e.g., bills)? If yes, please provide copies of the proposed materials. *	No
(10)	Is there any existing marketing agreement or contract with another firm for this product? If yes, a copy of the agreement must be submitted. *	No
(11)	Do all raw materials used in the manufacture of the product conform to the official standards specified in a recognized pharmacopoeia or official book? *	No
(12)	Has an application for this product, with the same name or formulation, been previously submitted? If so, was it ever rejected, and if yes, please state the reason for rejection. *	No
(13)	Is there any other product currently on the market with the same composition but a different brand name? *	No
(14)	Is there any other product currently available with a different composition but the same brand name? *	No

Submit Reset

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Each product row in the data table must have these fields filled in before final submission.

Q.C. (Quality Control) Register

After completing the product details section, the next step is to fill out the **Quality Control (Q.C.) Register**.

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Fresh License / Add Q.C (Quality Control) Register

Q.C (Quality Control) Register

Product Name *	Testing Protocol *	Study Start Date *	Study End Date *
HCQS 200 Tablet	(3) Month Accelerated Stability	28/05/2025	04/06/2025
Shelf Life of Product in Month	Batch No *	Report Date *	Test Report *
12	HCQS 200MG	28/06/2025	Choose file 20240528150_F_sample.pdf
Quality Control * ☒ In ☐ Out			

Submit Reset

Qc (Quality Control) Register

10 Entries Per Page

Search:

Sr No.	Testing Protocol	Study Start Date	Batch No	Shelf Life of Product in Month	Study End Date	Report Date	Test Report	Quality Control	Lab Name	Product Name	Action
No data available in table											

Showing 0 to 0 of 0 entries

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This section captures critical testing and compliance-related information, including:

- **Product Name** -- Select the product for which quality control data is being entered.
- **Testing Protocol** -- Specify the method or standard used for testing.
- **Batch No.** -- Enter the batch number of the product tested.
- **Test Report** -- Upload or enter details of the corresponding test report.
- **Date of Testing, Result**, and other required fields (if applicable)

Ensure accuracy and completeness of all Q.C. records before proceeding.

The screenshot shows the E-AUSHADHI PORTAL interface. The left sidebar contains navigation options: Dashboard, FRESH LICENSE, New Application, Download License, Dependant Application, Under Taking, PROFILE, LEGACY, CERTIFICATE, ADDITIONAL TP, ADDITIONAL PRODUCT, SURRENDER APPLICATION, and AMENDMENT APPLICATION. The main content area is titled 'Q.C (Quality Control) Register' and includes a form for entering new records. The form fields are: Product Name (dropdown), Testing Protocol (dropdown), Study Start Date (date picker), Study End Date (date picker), Shelf Life of Product in Month (text input), Batch No. (text input), Report Date (date picker), and Test Report (file upload). There are also checkboxes for Quality Control (In/Out) and buttons for Submit and Reset. Below the form is a table titled 'Qc (Quality Control) Register' with columns: Sr No., Testing Protocol, Study Start Date, Batch No., Shelf Life of Product in Month, Study End Date, Report Date, Test Report, Quality Control, Lab Name, Product Name, and Action. The table contains two entries. Entry 1: Sr No. 1, Testing Protocol 3 Month Accelerated Stability, Study Start Date 2025-05-28, Batch No. HCQS 200MG, Shelf Life of Product in Month 12, Study End Date 2025-06-04, Report Date 2025-06-28, Test Report (upload icon), Quality Control In, Lab Name -, Product Name HCQS 200 Tablet, Action (edit, delete). Entry 2: Sr No. 2, Testing Protocol 3 Month Accelerated Stability, Study Start Date 2025-06-02, Batch No. para, Shelf Life of Product in Month 12, Study End Date 2025-06-08, Report Date 2025-06-13, Test Report (upload icon), Quality Control In, Lab Name -, Product Name Paracetamol, Action (edit, delete). The table shows 1 to 2 of 2 entries. At the bottom, it says 'Website content owned by Ministry of Ayush, Government of India. Designed, developed and maintained by BISAG-N'.

Q.C (Quality Control) Section

After Submitting the License details section, the next step is to fill out the **Quality Control (Q.C.) Section**.



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Q.C (Quality Control) Section

Legacy / Q.C (Quality Control) Section

Quality Control In ☒

 Quality Control Out ☐

Save

 Reset

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Q.C (Quality Control) Section

Legacy / Q.C (Quality Control) Section

Quality Control In ☐

 Quality Control Out ☒

 Approved Laboratory Under License Form 48 *

 KESHVAM LAB PHARMA

 Consent Letter *

 Choose file | file.pdf

Laboratory Address

 vsvsvsvsvsvsvsAhmedabadGujarat382418

Remarks

 QC Section

Save

 Reset

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Technical Person Management

Adding Technical Person (TP) Details

- Click on the **(plus) icon** to open the **Add Technical Person** page.

On this page, fill in all **mandatory fields** related to the technical person, such as:

- Name
- Qualification
- Experience
- Contact Information
- Other required personal and professional details

Appointment Information Section

In the **TP (Technical Person) / Expert** section:

- TP (Technical Person)** Depends on **QC (Quality Control) Section** .

if you fill **QC Section In** Then 4 TP is Mandatory

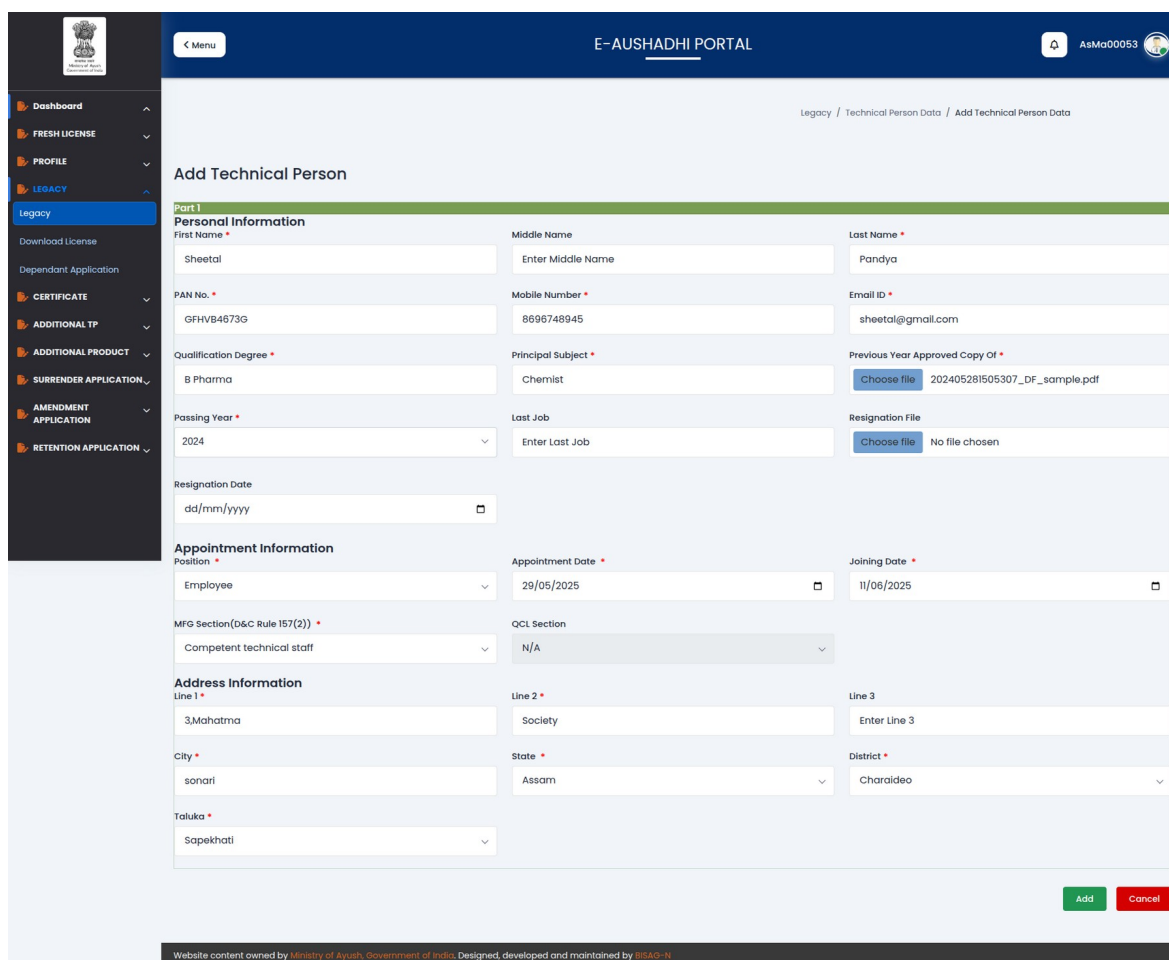
1 MFG - Component Technical Staff

3 QCL - Chemist, Botanist, Expert in ASU

In the **Appointment Information** section:

1 **First**, under the **MFG (Manufacturing)** section:

- You must select **Competent Technical Staff** from the dropdown.
- Fill in all required appointment details and click **Save**.



The screenshot shows the 'Add Technical Person' form in the E-Aushadhi Portal. The form is divided into several sections: Personal Information, Appointment Information, and Address Information. The Personal Information section includes fields for First Name, Middle Name, Last Name, PAN No., Mobile Number, Email ID, Qualification Degree, Principal Subject, Previous Year Approved Copy Of, Passing Year, Last Job, Resignation Date, and Resignation File. The Appointment Information section includes fields for Position, Appointment Date, Joining Date, MFG Section (D&C Rule 157(2)), and QCL Section. The Address Information section includes fields for Line 1, Line 2, Line 3, City, State, District, and Taluka. The form is currently filled with sample data, and the 'Add' button is visible at the bottom right.

to add additional technical person:

- Click **Add New TP**.
- At this stage, the **MFG section will be marked as N/A**.
- Under the **QCL (Quality Control Lab)** section, select the relevant **expertise** from the dropdown:

- Expert in ASU



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Legacy / Technical Person Data / Add Technical Person Data

Add Technical Person

Part 1

Personal Information

First Name *

Nitya

Middle Name

Enter Middle Name

Last Name *

Sharma

PAN No. *

JFBJR5473H

Mobile Number *

9765383743

Email ID *

nitya@gmail.com

Qualification Degree *

B pharma

Principal Subject *

Botanist

Previous Year Approved Copy Of *

Choose file

202405281505307_DF_sample.pdf

Passing Year *

2024

Last Job

Enter Last Job

Resignation File

Choose file

No file chosen

Resignation Date

dd/mm/yyyy

Appointment Information

Position *

Employee

Appointment Date *

30/05/2025

Joining Date *

04/06/2025

MFG Section (D&C Rule 157(2))

N/A

QCL Section *

Expert in ASU

Address Information

Line 1 *

A-34,Nilkanth

Line 2 *

Apartment

Line 3

Enter Line 3

City *

algapur

State *

Assam

District *

Hailakandi

Taluka *

Hailakandi

Add

Cancel

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• Chemist



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Legacy / Technical Person Data / Add Technical Person Data

Add Technical Person

Part I

Personal Information

First Name *

Naina

Middle Name

Enter Middle Name

Last Name *

Rathore

PAN No. *

GHKTR5784G

Mobile Number *

8658584576

Email ID *

naina@gmail.com

Qualification Degree *

B Pharma

Principal Subject *

Chemistry

Previous Year Approved Copy Of *

Choose file

202405281505307_DF_sample.pdf

Passing Year *

2025

Last Job

Enter Last Job

Resignation File

Choose file

No file chosen

Resignation Date

dd/mm/yyyy

Appointment Information

Position *

Partner

Appointment Date *

05/06/2025

Joining Date *

11/06/2025

MFG Section(D&C Rule 157(2))

N/A

QCL Section *

Chemist

Address Information

Line 1 *

4,Shashtri nagar

Line 2 *

bungalows

Line 3

Enter Line 3

City *

athani

State *

Assam

District *

Dhubri

Taluka *

Dhubri

Add

Cancel

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- Botanist



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Legacy / Technical Person Data / Add Technical Person Data

Add Technical Person

Part 1
Personal Information
 First Name *
 Nandini

Middle Name
 Enter Middle Name

Last Name *
 Chauhan

PAN No. *
 JKFNT6458H

Mobile Number *
 9759747585

Email ID *
 nandini@gmail.com

Qualification Degree *
 B Pharma

Principal Subject *
 Biology

Previous Year Approved Copy Of *
 Choose file 202405281505307_DF_sample.pdf

Passing Year *
 2023

Last Job
 Enter Last Job

Resignation File
 Choose file No file chosen

Resignation Date
 dd/mm/yyyy

Appointment Information
 Position *
 Proprietor

Appointment Date *
 29/05/2025

Joining Date *
 05/06/2025

MFG Section(D&C Rule 157(2))
 N/A

QCL Section *
 Botanist

Address Information
 Line 1 *
 3,Akash

Line 2 *
 Apartment

Line 3
 Enter Line 3

City *
 salmara

State *
 Assam

District *
 South Salmara Mancachar

Taluka *
 South Salmara

Add Cancel

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if you fill **QC Section Out** Then 1 TP is Mandatory
1 MFG - Component Technical Staff

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Legacy / Technical Person Data / Add Technical Person Data

Add Technical Person

Part 1

Personal Information

First Name *	Middle Name	Last Name *
Sheetal	Enter Middle Name	Pandya
PAN No. *	Mobile Number *	Email ID *
GFHVB4673G	8696748945	sheetal@gmail.com
Qualification Degree *	Principal Subject *	Previous Year Approved Copy Of *
B Pharma	Chemist	Choose file 202405281505307_DF_sample.pdf
Passing Year *	Last Job	Resignation File
2024	Enter Last Job	Choose file No file chosen
Resignation Date	dd/mm/yyyy	

Appointment Information

Position *	Appointment Date *	Joining Date *
Employee	29/05/2025	11/06/2025
MFG Section(D&C Rule 157(2)) *	QCL Section	
Competent technical staff	N/A	

Address Information

Line 1 *	Line 2 *	Line 3
3,Mahatma	Society	Enter Line 3
City *	State *	District *
sonari	Assam	Charaideo
Taluka *		
Sapekhati		

Add Cancel

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Application Attachment

After entering all required details—including Product, Q.C. Register, and Technical Personnel—the next step is to upload documents in the **Application Attachment List** section.

In this section:

- You will see a list of **mandatory and optional documents** required for your application type.

- For each document listed, you must:
 - **Attach the file** in the required format (PDF, JPEG, etc.).
 - Ensure that the **file size** complies with portal guidelines.

Mandatory documents must be uploaded to proceed with final submission.

Ensure each uploaded document is clear, legible, and matches the description provided.



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Fresh License / License Attachment

Application Attachment List

(1)	Owners ID proof *	Choose file	20240528150_F_sample.pdf
(2)	Company-Firm ownership proofs-eg, patnership, company registration certificate, memorandum of artical, etc *	Choose file	20240528150_F_sample.pdf
(3)	Premises address proof Any government approved proof *	Choose file	20240528150_F_sample.pdf
(4)	Annexure A Site master plan *	Choose file	20240528150_F_sample.pdf
(5)	Annexure B Category wise list of manufacturing equipment or machineriles *	Choose file	20240528150_F_sample.pdf
(6)	Annexure C Category wise list of quality control instrument-equipment or glass wares	Choose file	No file chosen
(7)	Appointment letter of TP-Expert *	Choose file	20240528150_F_sample.pdf
(8)	Consent letter of government approved laboratory in case of outside laboratory testing of raw and finished product *	Choose file	20240528150_F_sample.pdf
(9)	Annexure D TP aggrement letter *	Choose file	20240528150_F_sample.pdf
(10)	Experience letter of TP-Expert	Choose file	No file chosen
(11)	Extra Field	Choose file	No file chosen

Submit
 Reset

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AIF (Additional Information Form)

After completing the **Application Attachment List**, the next step is to fill out the **AIF -- Additional Information Form**.



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Additional Information Form

(1) What business was conducted by the applicant(s) within the last three years? *

(2) Has the applicant or any other person on their behalf ever been involved in the manufacture or sale of drugs prior to this application? If so, please provide the relevant dates. *

No

(3) Is the applicant or owner/proprietor a registered/approved pharmacist or expert or compitant technical staff? *

No

(4) Is the applicant or owner currently operating any other business? *

No

(5) Has the applicant, owner, or any person currently employed at these premises ever been convicted and sentenced under any of the following acts? *

5.1) Drugs and Cosmetics Act, 1940 *

No

5.2) Dangerous Drugs Act, 1930 *

No

5.3) The poisons Act, 1919 *

No

5.4) The Pharmacy Act, 1948 *

No

5.5) The Drugs and Magic Remedies (Objectionable Advertisements) Act, 1954 *

No

(6) Does the applicant manufacture drugs for sale at any premises other than the one for which this license is being sought? If so, please list the addresses of such other premises. *

No

(7) What categories of drugs are currently manufactured, or are intended to be manufactured, by the firm? *

(8) Have any of the firm's licenses ever been suspended or canceled in the past? *

No

(9) Please state the business hours, working days, and holidays. Any change in the holiday schedule must be intimated to the licensing authority. *

(10) Please provide the name of the trade or professional association of which the applicant is a member, and the date when the membership commenced. *

Submit

Reset

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Undertaking for License

After completing the **AIF (Additional Information Form)**, the next step before submission is to acknowledge and accept the **Undertaking for License**.

In this step:

- You must **read and review** the declaration/undertaking terms provided on the screen.
- Confirm your acceptance by **checking the acknowledgment box**.

Undertaking License

Select All

I/We, the undersigned, hereby solemnly affirm and declare as follows:

- ☒ (1) I/We acknowledge and accept full and complete responsibility for the accurate and truthful submission of all applications and their accompanying documents and related data, including but not limited to information and particulars, through the official e-aushadhi.gov.in portal.
- ☒ (2) I/We understand and agree that any document or data found to be fabricated, falsified, erroneous, or misleading, submitted under my/our login credentials on the e-aushadhi.gov.in portal, shall be my/our sole responsibility, and I/we shall be held accountable for any and all consequences arising therefrom, including legal liabilities.
- ☒ (3) I/We further acknowledge and agree that any document uploaded or submitted through my/our designated user login on the e-aushadhi.gov.in portal, irrespective of the absence of a physical signature thereon, shall be deemed to have been duly signed and authenticated by me/us, thereby bearing my/our full endorsement and legal validity.
- ☒ (4) I/We confirm my/our complete awareness and understanding of the importance and security implications associated with the usage of my/our unique user ID and Password for accessing the e-aushadhi.gov.in account. Furthermore, I/we acknowledge full cognizance of the mobile number and email address linked to this e-aushadhi account and accept responsibility for their security and proper maintenance.
- ☒ (5) I/We hereby affirm that I/we possess comprehensive knowledge of the provisions of the Drugs and Cosmetics Act, 1940, and the Rules framed thereunder, and I/we undertake to strictly adhere to and comply with all such provisions. I/we further acknowledge that compliance with the aforementioned Act and Rules is my/our sole responsibility.
- ☒ (6) I/we are fully cognizant of and shall adhere to all Good Manufacturing Practices (GMP) norms and standards applicable to our operations. I/we affirm that all documents related to GMP, submitted by me/us, are valid, accurate, and complete. Furthermore, I/we undertake to provide any and all additional details or information as and when required by legal provisions or by the concerned authority.
- ☒ (7) I/We hereby commit to full adherence to all stipulated license conditions, product-specific conditions, and any other conditions of approval issued by the competent authority. I/we further acknowledge and agree that in the event of any violation or non-compliance with these conditions, I/we shall bear full and sole responsibility for all consequences, liabilities, and penalties arising therefrom.
- ☒ (8) I/we affirm that all documents pertaining to the safety, efficacy, and quality of drugs, submitted by me/us, are genuine, accurate, and in strict compliance with the Drugs and Cosmetics Act, 1940, and all other applicable acts and provisions. I/we further acknowledge and agree that I/we shall be solely responsible for any consequences arising from the submission or use of any fabricated, falsified, or misleading documents in this regard.
- ☒ (9) I/We declare that I/we have never been involved in the violation of any criminal law or any other law, and there is no adverse history or record pertaining to such violations against me/us.
- ☒ (10) I/we are fully aware of the provisions of the Drugs and Magic Remedies (Objectionable Advertisements) Act, 1954, and the rules framed thereunder. I/we solemnly undertake to comply with all regulations and provisions concerning advertisements and claims as set forth in the aforementioned Act and its rules. Furthermore, I/we affirm that I/we shall not publish any advertisement that is in violation of the said Act or any other applicable laws and rules.

Note: This undertaking is made with full knowledge of its contents and implications, and by its electronic submission through the e-aushadhi.gov.in portal, I/we hereby bind myself/ourselves to its terms.

Save Reset

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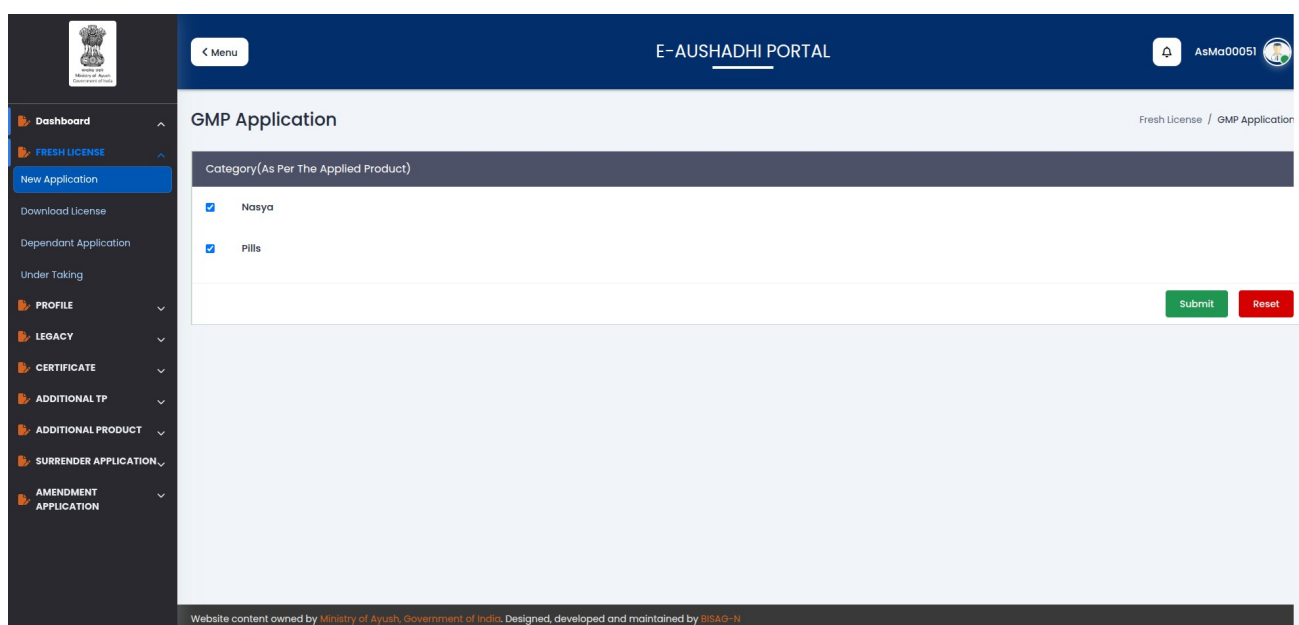
GMP Application

After completing the **Undertaking for License**, the next required step is to confirm and complete the **GMP Application** section.

In this step:

- You must check or confirm the **GMP Application** checkbox to indicate your intention to apply for a **GMP certificate**.
- Fill in any additional required fields if prompted, such as:
 - GMP application type or category

Checking this option is mandatory for issuing or renewing a manufacturing license under ASU guidelines.



The screenshot shows the E-Aushadhi Portal interface. The top navigation bar includes the Ministry of Ayush logo, a menu button, the text 'E-AUSHADHI PORTAL', a user profile icon with the ID 'AsMa00051', and a globe icon. The left sidebar contains a list of menu items: Dashboard, FRESH LICENSE, New Application (highlighted), Download License, Dependant Application, Under Taking, PROFILE, LEGACY, CERTIFICATE, ADDITIONAL TP, ADDITIONAL PRODUCT, SURRENDER APPLICATION, and AMENDMENT APPLICATION. The main content area is titled 'GMP Application' and shows a section for 'Category(As Per The Applied Product)' with two checked options: 'Nasya' and 'Pills'. At the bottom right of this section are 'Submit' and 'Reset' buttons. The footer of the page states: 'Website content owned by Ministry of Ayush, Government of India. Designed, developed and maintained by BISAG-N'.

Final Registration and Payment

Once **all required details** have been filled and verified — including product information, QC register, technical person, attachments, AIF, undertaking, and GMP application — you are ready to **register the application**.

Registering the Application:

- Click on the **Register** button at the bottom of the form.
- Upon successful registration, the **Payment** option will become enabled.

e-Aushadhi Portal

Dashboard / Fresh License

Fresh License (Manufacture) Application - Ayurveda, Unani, Siddha, Sowa Rigpa

Form No.	Status	Dispose Days Left	Edit	Products	Qc Register	Qc (Quality Control) Section	TP/Expert	Attachments	AIF	Under Taking	GMP	Constitution	Payment	Refer Back	View Inspection	Report	Delete	Action
24D	Under Verification	60																Applied for Registration

Legend:

- This color shows that the license is in a refer-back status.
- This color indicates that you are restricted from taking any action.

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Proceeding with Payment:

- Click on the **Payment** button to proceed.

E-AUSHADHI PORTAL

Fresh License / Add Payment

Payment for Application No - 00044

Application Information

Form No.	24D	Firm Name	Prajakta	Application No.	00044
Reg Date	19-06-2025	Mobile No.	9886965458	Email ID	himanshi@gmail.com
Total Amount	7200	Payment Date	19-06-2025	Api Type	Own
Payment Type *	Offline	Challan File *	Choose file	GAR 6 No. *	Transactionid234

Charge Back Policy

Terms & Conditions :-

- The transaction once done cannot be cancelled.
- Fees once paid is not refundable in any case.
- Refund claims (if found eligible and admissible) will be entertained ONLY OFFLINE and NO ONLINE REFUNDS will be done.
- Convenience fees / Bank charges, Commission of Bank plus service tax would be charged (if applicable) in addition to the fee Amount and can't be claim back as refund.
- This transaction is payment towards Government dues, fees hence would not be covered under charge back policy.
- I/We agree and accept all the terms and conditions mention above and I/We hereby certify that no online charge back claim will be made in this regards.

Please check all details before clicking on confirm button

Payment Status Description:

- SUCCESS: Payment Done Successfully.
- FAIL or FAILURE: The transaction FAILED from gateway.
- PENDING: The transaction is pending from gateway. Kindly Click on Reconciliation button to know latest status of the transaction.
- ABORT: The Transaction FAILED or CANCELLED from gateway.
- IN PROCESS: The transaction payment is initiated but response is awaited from payment gateway. Kindly Click on Reconciliation button to know latest status of the transaction.

Previous Transaction Details

10 Entries Per Page

Search:

Sr No.	Transaction Id	Transaction Date and Time	Payment amount	Payment Gateway Date	Payment Status	Reconciliation
No data available in table						

Showing 0 to 0 of 0 entries

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- You will be **redirected to the Payment Gateway page**, where:
 - Currently, only Offline Payment is available.**

- You must generate a **Challan** and make the payment through the designated offline mode (e.g., treasury or bank deposit).

Upload Challan:

- Upload a scanned copy of the **Challan** as proof of payment.
- Uploading the challan is **mandatory** to complete the payment process.

Ensure that the uploaded challan is clear, legible, and in the prescribed format.

After successful upload, your application will move forward for official processing.

Ministry of Ayush
Government of India

- Dashboard
- FRESH LICENSE
- New Application
- Download License
- Dependant Application
- Under Taking
- PROFILE
- LEGACY
- CERTIFICATE
- ADDITIONAL TP
- ADDITIONAL PRODUCT
- SURRENDER APPLICATION
- AMENDMENT APPLICATION

< Menu

E-AUSHADHI PORTAL

AsMa00051

Fresh License / Add Payment

Payment for Application No - 00044

Application Information

Form No	24D	Firm Name	Projakta	Application No.	00044
Reg Date	19-06-2025	Mobile No.	9886965458	Email ID	himanshi@gmail.com
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Previous Transaction Details

10 Entries Per Page

Search:

Sr No.	Transaction Id	Transaction Date and Time	Payment amount	Payment Gateway Date	Payment Status	Reconciliation
1	TransactionId234	2025-06-19 14:53:41.869	7200	2025-06-19	Success	true

Showing 1 to 1 of 1 entry

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Ensure you do not close the browser or refresh the page during payment processing.

After successful payment, a receipt will be generated and your application will be submitted for further processing.

e-Aushadhi Portal

Dashboard / Fresh License

Fresh License (Manufacture) Application - Ayurveda, Unani, Siddha, Sowa Rigpa

Form No.	Status	Dispose Days Left	Edit	Products	Qc Register	Qc (Quality Control) Section	TP/Expert	Attachments	AIF	Under Taking	GMP	Constitution	Payment	Refer Back	View Inspection	Report	Delete	Action
1	24D Under Verification	60											Paid - 5200					Applied for Registration

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 This color indicates that you are restricted from taking any action.

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License Download:

- Once the **Technical Scrutiny is successfully completed** and your application is approved:
 - You can **download your Manufacturing License** directly from the portal.
 - Navigate to the application dashboard.
 - Click on the **"Download License"** button available at the **top-right corner** of the application view page.

Make sure to keep a digital and printed copy of your license for future reference and compliance.

E-AUSHADHI PORTAL

Dashboard / License Download

License Download - Manufacture (Ayurveda, Unani, Siddha, Sowa Rigpa)

Sr No.	Application No.	License No.	Issue Date	Product	TP	GMP	License Download
1	00044	As/D/014/2025	2025-06-19				

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