

e-Aushadhi



User Manual For Manufacturing Fresh License (Homeopathy)

=====

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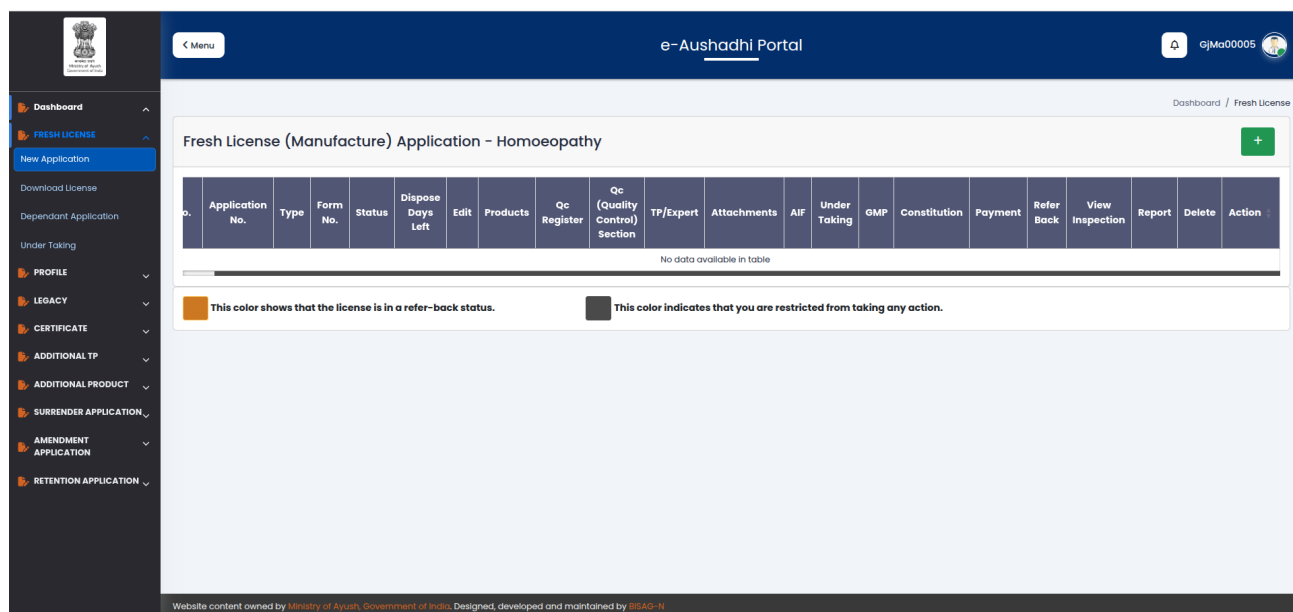
➤ Fresh License (Homoeopathy) Application

Submitting a New License Application

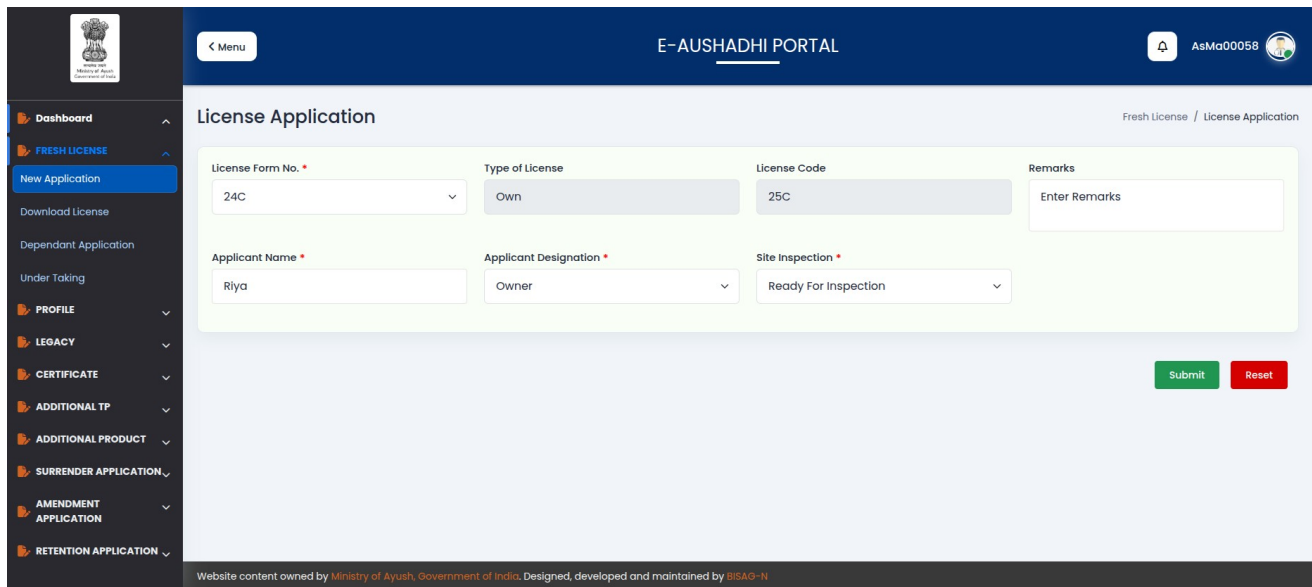
After logging in and receiving approval, the system will redirect you to the **User Dashboard**.

To begin a new license application:

- ◆ 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:**
 1. Select your **Own Firm Name** and the relevant **License**.
 2. Enter the **OTP** sent to your own firm's registered email address.
 3. 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 has a breadcrumb trail 'Fresh License / License Application'. The form fields are as follows:

License Form No. *	Type of License	License Code	Remarks
24C	Own	25C	Enter Remarks

Applicant Name *	Applicant Designation *	Site Inspection *
Riya	Owner	Ready For Inspection

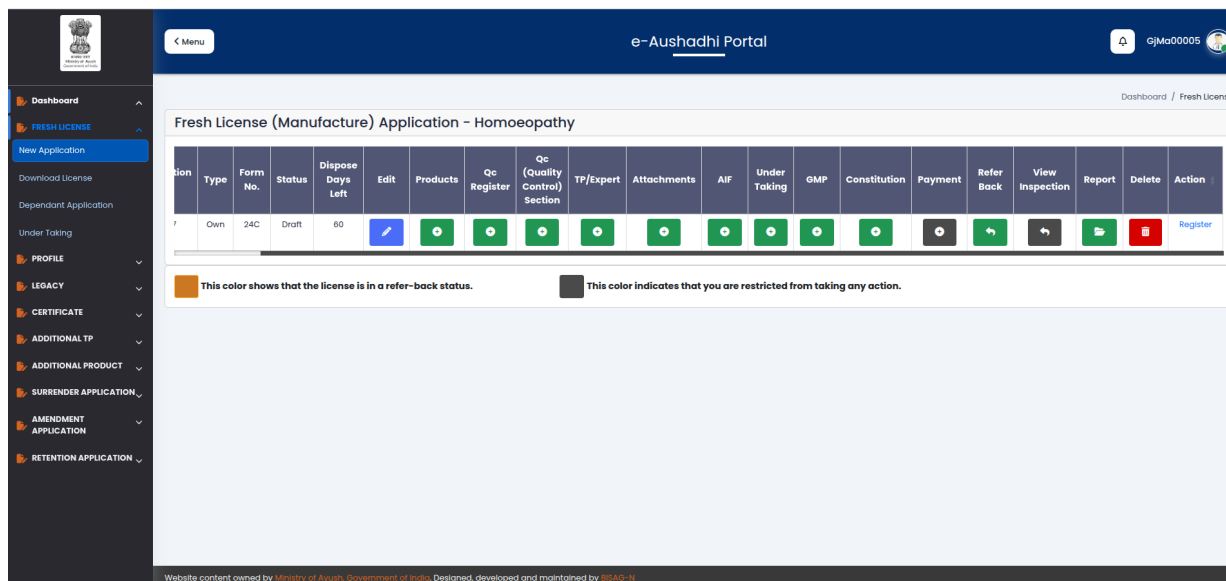
At the bottom right of the form, there are two buttons: 'Submit' (green) and 'Reset' (red).

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➤ 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 complybutton 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**

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

➤ **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



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e-Aushadhi Portal

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Dashboard

FRESH LICENSE

New Application

Download License

Dependant Application

Under Taking

PROFILE

LEGACY

CERTIFICATE

ADDITIONAL TP

ADDITIONAL PRODUCT

SURRENDER APPLICATION

AMENDMENT APPLICATION

RETENTION APPLICATION

Product Name *

Paracetomal

Type *

Homoeopathic Single Ingredient Medicine

Category/Dosage Form *

Globule

Potency and Dilution *

24X

Permission Purpose *

Regular

Remarks

DDDDDDDDDD

Composition Title

e.g. Each Gram Contents

7 ml

Shelf Life in Month *

4

Drug Categories Based On Their pharmacological Actions *

Respiratory System (Lungs & Breathings)

Diseases and Conditions *

Tests

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
No data available in table										

Showing 0 to 0 of 0 entries

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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RETENTION APPLICATION

Product Name *

Enter Product Name

Type *

--Select--

Category/Dosage Form *

--Select--

Potency and Dilution *

--Select--

Permission Purpose *

--Select--

Remarks

Enter Remarks

Composition Title

e.g. Each Gram Contents

Each Gram Contents

Shelf Life in Month *

Enter Month

Drug Categories Based On Their pharmacological Actions *

-- Select --

Diseases and Conditions *

Enter Disease

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	Paracetomal	Homoeopathic Single Ingredient Medicine	Globule	24X	Pending					

Showing 1 to 1 of 1 entry

➤ 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:

1. Product Composition



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

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 GuMa00009
 

Add Composition

Fresh License / Product Data / Add Composition

Ingredient Type *

INERT

Ingredient Name *

vvvvv

Botanical Name *

vbnbnvbn

Reference *

French Homoeopathic Pharmacopoeia

Part Used *

LATEX

Quantity *

66

Measuring Units *

QS

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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2. Product Attachments



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 Government of India

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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..._sample.pdf
(2)	Annexure 3 Master Formula card with manufacturing SOP *	Choose file	20240528150..._sample.pdf
(3)	Annexure 4 Quality control specification *	Choose file	20240528150..._sample.pdf
(4)	Annexure 5 Proof of effectiveness With Reference Page As per Rule 158 B *	Choose file	20240528150..._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..._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..._sample.pdf
(8)	Validated data As per D and C Rule 161 B *	Choose file	20240528150..._sample.pdf
(9)	Extra Field	Choose file	No file chosen

Submit
 Reset

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3. AIF (Additional Information Form)

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



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e-Aushadhi Portal

GuMa00001

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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Q.C. (Quality Control) Register

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

Q.C. (Quality Control) Register

Product Name: HCQS 200 Tablet | Testing Protocol: Real Time Stability | Study Start Date: 03/06/2025 | Study End Date: 20/06/2025

Shelf Life of Product In Month: 0 | Batch No.: HCQS | Report Date: 23/06/2025 | Test Report: Choose file 2024052815_F_sample.pdf

Quality Control: ☒ In ☐ Out

Qc (Quality Control) Register

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

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

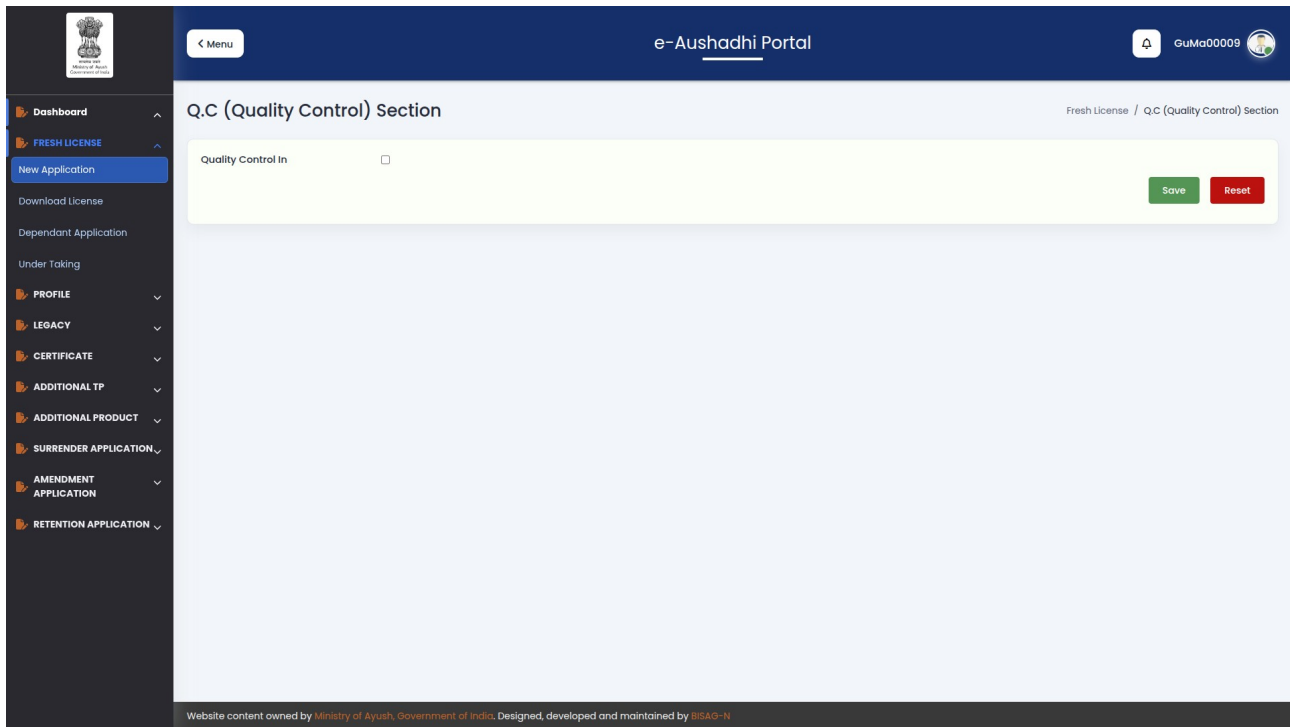
- **Product Name** -- Select the product for which quality control data is being entered.
- **Testing Protocol** -- Specify the method or standard used for testing.
- **Batch No.** -- Enter the batch number of the product tested.
- **Test Report** -- Upload or enter details of the corresponding test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

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

➤ Q.C. (Quality Control) Section

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

Here **QC (Quality Control) Section In** Is Mandatory.



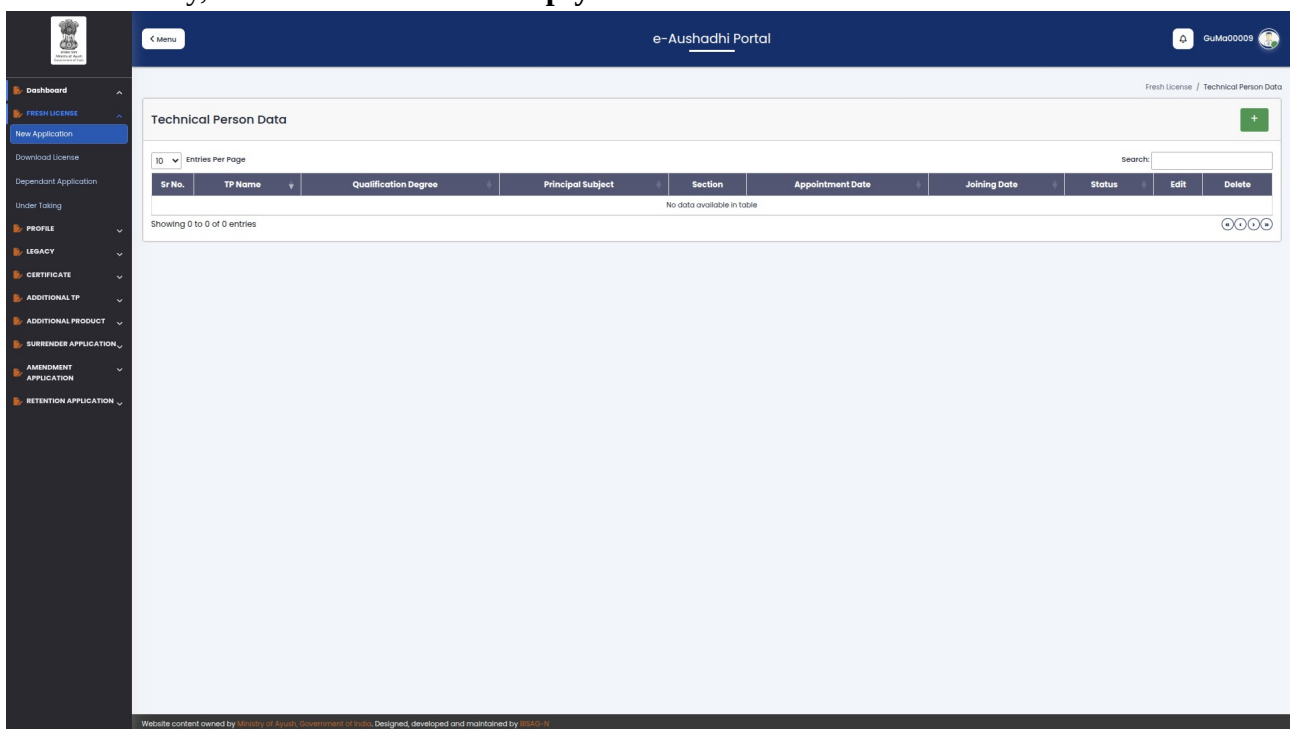
The screenshot shows the 'Q.C (Quality Control) Section' of the e-Aushadhi Portal. The left sidebar contains a menu with options: Dashboard, FRESH LICENSE (highlighted), New Application, Download License, Dependant Application, Under Taking, PROFILE, LEGACY, CERTIFICATE, ADDITIONAL TP, ADDITIONAL PRODUCT, SURRENDER APPLICATION, AMENDMENT APPLICATION, and RETENTION APPLICATION. The main content area has a header 'Q.C (Quality Control) Section' and a sub-header 'Fresh License / Q.C (Quality Control) Section'. Below this is a form titled 'Quality Control In' with a single input field and two buttons: 'Save' and 'Reset'.

➤ Technical Person Management

Adding Technical Person (TP) Details

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

- Initially, the data table will be empty.



The screenshot shows the 'Technical Person Data' section of the e-Aushadhi Portal. The left sidebar is the same as the previous screenshot. The main content area has a header 'Technical Person Data' and a sub-header 'Fresh License / Technical Person Data'. Below this is a table with columns: Sr No., TP Name, Qualification Degree, Principal Subject, Section, Appointment Date, Joining Date, Status, Edit, and Delete. The table is currently empty, showing 'No data available in table'. There is a search bar and a dropdown for 'Entries Per Page' (set to 10) above the table. A green '+' button is in the top right corner of the table area.

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

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

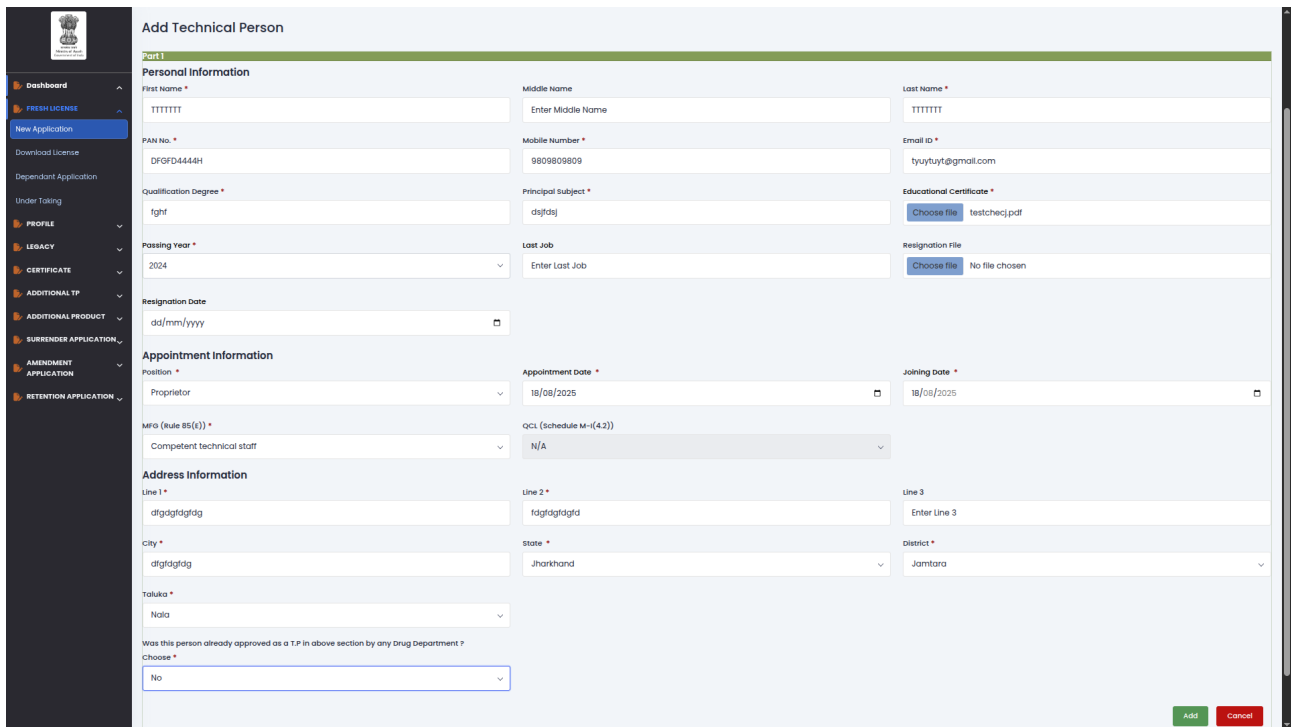
In the **Appointment Information** section:

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

In Fresh License Homoeopathy Total 2 TP is Mandatory

- 1 MFG Section
- 1 QCL Section

- MFG Section



Add Technical Person

Part 1

Personal Information

First Name *

Middle Name

Last Name *

PAN No. *

Mobile Number *

Email id *

Qualification Degree *

Principal Subject *

Educational Certificate *

Passing Year *

Last Job

Resignation File

Resignation Date

Appointment Information

Position *

Appointment Date *

Joining Date *

MFG (Rule 85(1)) *

QCL (Schedule M-II(4.2))

Address Information

Line 1 *

Line 2 *

Line 3

City *

State *

District *

Taluka *

Was this person already approved as a T.P in above section by any Drug Department ?

- QCL Section

➤ 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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SURRENDER APPLICATION

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RETENTION APPLICATION

Fresh License / License Attachment

Application Attachment List

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

GuMa00001

Fresh License / Additional Information Form

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. *

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

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

(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 *

5.2) Dangerous Drugs Act, 1930 *

5.3) The poisons Act, 1919 *

5.4) The Pharmacy Act, 1948 *

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

(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. *

(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? *

(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

In this step:

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

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Dependant Application

Under Taking

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ADDITIONAL TP

ADDITIONAL PRODUCT

SURRENDER APPLICATION

AMENDMENT APPLICATION

RETENTION APPLICATION

Undertaking License

Fresh License / 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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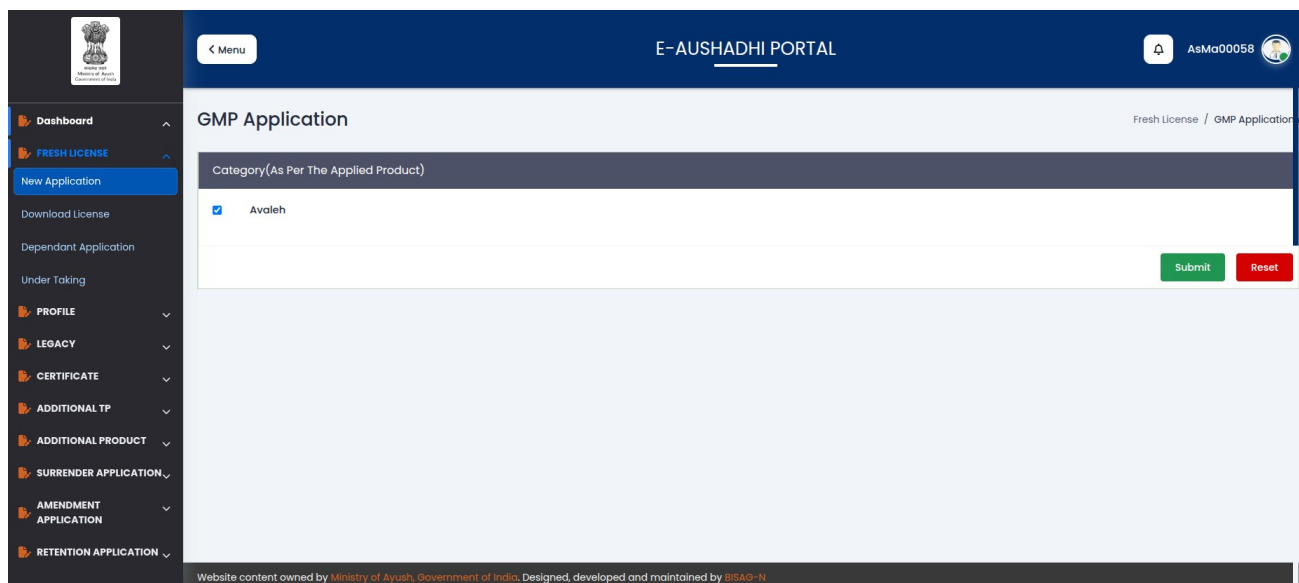
➤ GMP Application

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.



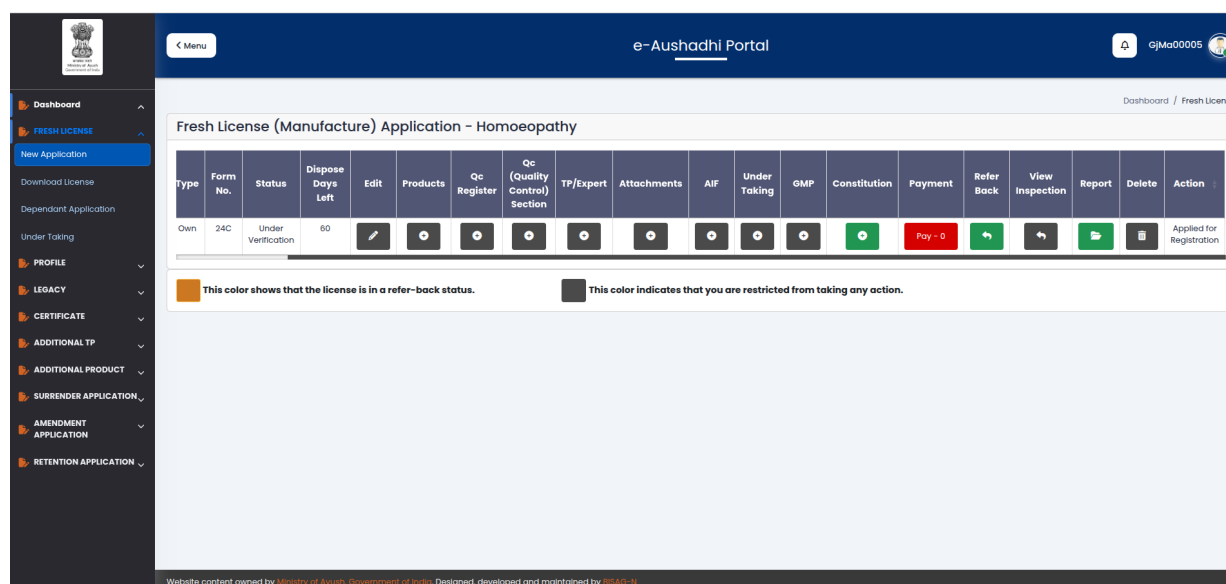
The screenshot shows the 'GMP Application' form on the E-Aushadhi Portal. The left sidebar contains a menu with options like Dashboard, FRESH LICENSE, New Application, Download License, etc. The main content area has a header 'GMP Application' and a sub-header 'Category(As Per The Applied Product)'. Below this, there is a checkbox labeled 'Avaleh' which is checked. At the bottom right of the form, there 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.



The screenshot shows the 'Fresh License (Manufacture) Application - Homoeopathy' page on the E-Aushadhi Portal. The left sidebar is the same as in the previous screenshot. The main content area displays a table with the following columns: Type, 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, and Action. The table contains one row with the following data: Type: Own, Form No.: 24C, Status: Under Verification, Dispose Days Left: 60, Edit: (pencil icon), Products: (plus icon), Qc Register: (plus icon), Qc (Quality Control) Section: (plus icon), TP/Expert: (plus icon), Attachments: (plus icon), AIF: (plus icon), Under Taking: (plus icon), GMP: (plus icon), Constitution: (plus icon), Payment: (plus icon), Refer Back: (plus icon), View Inspection: (plus icon), Report: (plus icon), Delete: (trash icon), and Action: Applied for Registration. Below the table, there are two legends: 'This color shows that the license is in a refer-back status.' (orange square) and 'This color indicates that you are restricted from taking any action.' (grey square).

Proceeding with Payment:

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



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Fresh License / Add Payment

 Payment for Application No - 00055

Application Information

Form No	24C	Firm Name	Riya	Application No.	00055
Reg Date	24-06-2025	Mobile No.	8694574793	Email ID	pooja@gmail.com
Total Amount	5200	Payment Date	24-06-2025	Apl Type	Own
Payment Type *	Offline	Challan File *	Choose file 2024052_ample.pdf	GAR 6 No. *	Trid32

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

Confirm
 Reset

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

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

- 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.

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AsMa00058

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AMENDMENT APPLICATION

RETENTION APPLICATION

Payment for Application No - 00055

Application Information

Form No	24C	Firm Name	Riya	Application No.	00055
Reg Date	24-06-2025	Mobile No.	8694574793	Email ID	pooja@gmail.com
Total Amount	5200	Payment Date	24-06-2025	Apl Type	Own

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."

Previous Transaction Details

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.

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Entries Per Page

Search:

Sr No.	Transaction Id	Transaction Date and Time	Payment amount	Payment Gateway Date	Payment Status	Reconciliation
1	Trid32	2025-06-24 11:55:31.325	5200	2025-06-24	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.

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Fresh License (Manufacture) Application - Homoeopathy

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

This color shows that the license is in a refer-back status.

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

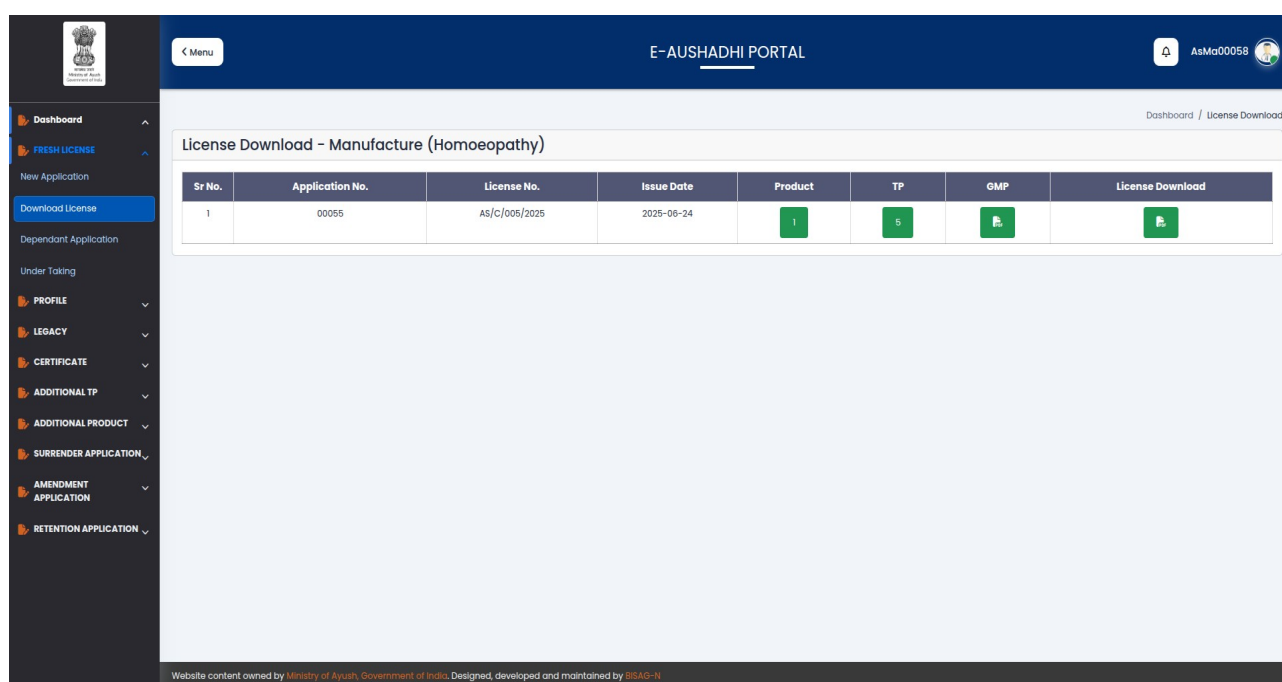
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After successful payment, a receipt will be generated and your application will be submitted for further processing.

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License Download:

- Once the 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.



The screenshot shows the E-Aushadhi Portal interface. The top navigation bar includes the Ministry of Ayurveda logo, a menu button, the portal name 'E-AUSHADHI PORTAL', and a user profile section with a notification bell and the username 'AsMa00058'. The left sidebar contains various application categories 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 'License Download - Manufacture (Homoeopathy)' and contains a table with the following data:

Sr No.	Application No.	License No.	Issue Date	Product	TP	GMP	License Download
1	00055	AS/C/005/2025	2025-06-24				

At the bottom of the page, a footer note states: 'Website content owned by Ministry of Ayurveda, Government of India. Designed, developed and maintained by BISAG-N'.

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