



الإمارات العربية المتحدة وزارة الصحة ووقاية المجتمع

Import Export

Agent Manual

Information Technology Department - Development & E-services section

5/31/2017

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1. Copyright and Confidentiality

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2. Acronyms, Abbreviations and Definitions:

Abbreviations	Definitions
MoHAP	Ministry of Health and Prevention

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3. Purpose and Objectives

This document explains all the steps which are required for Site Administrator and agent user to Create

Application, View Application, Reports, and User Manual Website contents. This document will provide

Ease to the business users in understanding all the screens of the application.

3.1. Manage Agent Contents

It contains information related to contents manipulation such as insert, update or delete operations

4. Assumptions

You need to have administrator or agent privileges and be logged in with your account to perform administrative or agent User tasks

5. For All Users

5.1. Login

For using the service, All the applicants needs to logged in from User Management

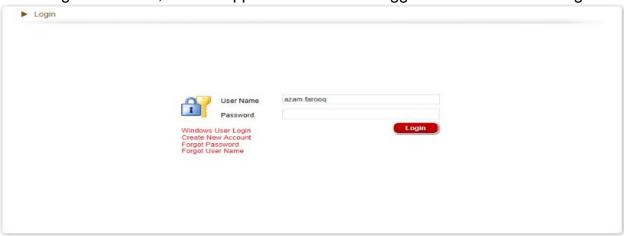


Figure 1 – Login (External Users)

For internal MOH users, following screen will be used:

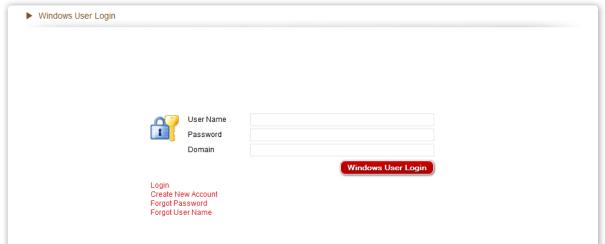


Figure 2 - Login (Internal Users)

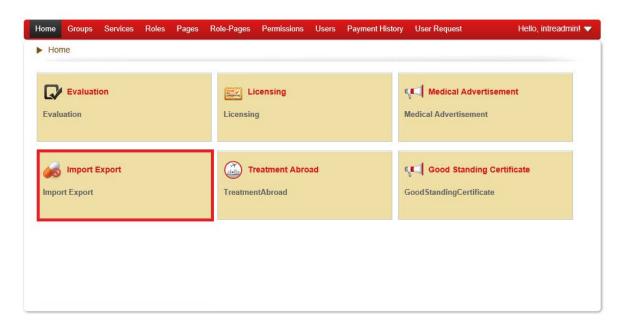


Figure 3 - Services Dashboard

6. Agent Menu

Agent will login application through User management and click on Drug Import Export service. The agent shall see the following options to create new or view existing applications:



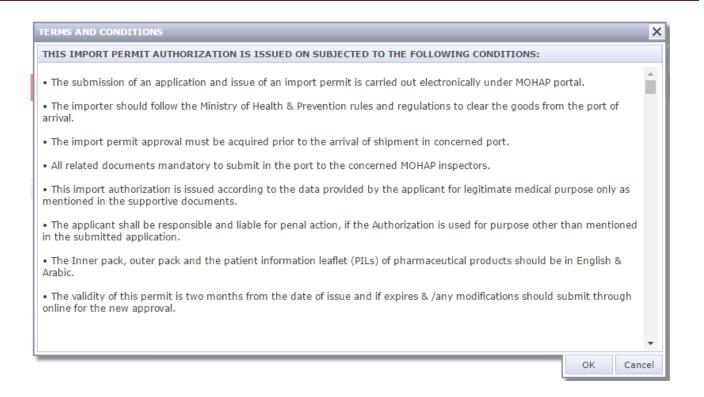
Figure 4 - Agent Menu

6.1. Create Application

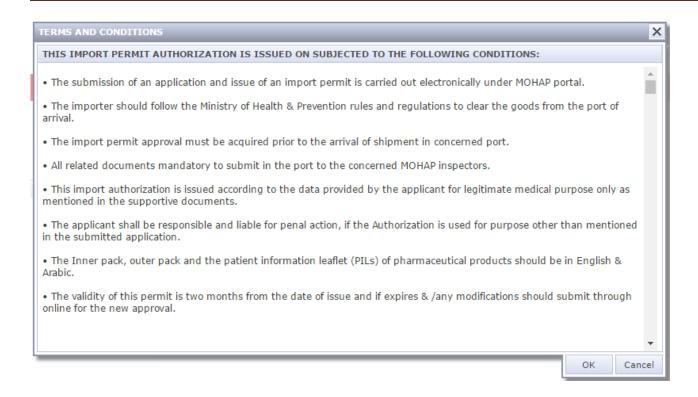


Figure 5 - Create Application

- The agent has the option to create or view an already created application from the above menu.
- Click on "Create Application" button of the relevant section and then proceed:
 - Authorization Applications: These are mandatory pre-approvals for importing the following types of products:
 - Biological Products
 - Controlled Application Chemical Precursors
 - Controlled Medicines (SCD/CD/NP))
 - o Import/Export Permits:
 - Pre-Authorization permits for importing or exporting products.
 - Accept the Terms and Conditions by clicking OK.



- Shipment Clearance Applications:
 - This is the final step in getting an import/export permit for as a shipment clearance.
 - Accept the Terms and Conditions by clicking OK.



6.2. Authorization Applications

6.2.1.Biological Product

On Module screen select Type Import/Export and select Module Biological Products



Figure 6 - Biological Product Authorization

 System will load the Biological Product Form for submitting an Authorization Request.

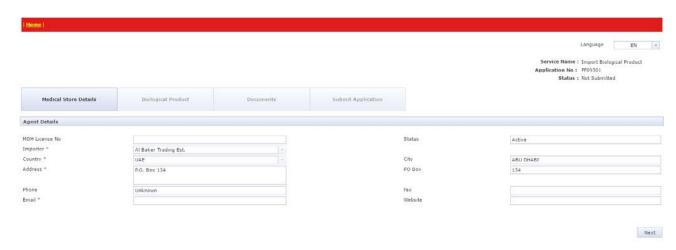


Figure 7 - Biological Product Authorization Request

Verify the Medical and Store Detail and click next.

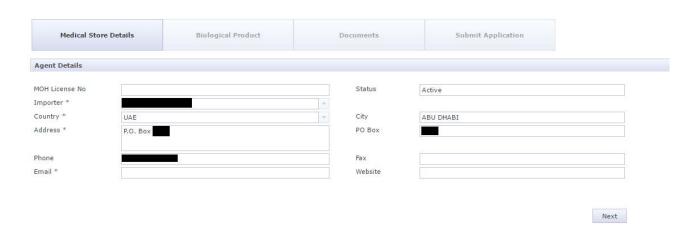


Figure 8 – Medical Store Details

On Biological Product tab and entre data in all the fields relevant fields.

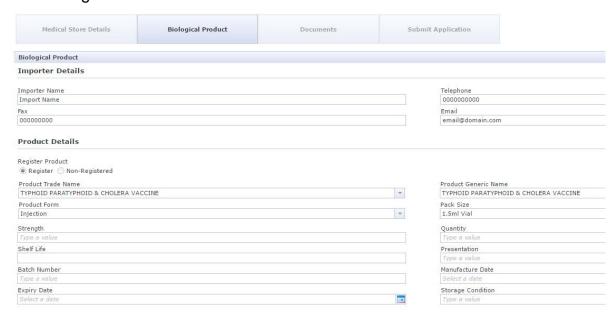


Figure 9 - Biological Product

 Select the Documents Tab and attach all the required documents. Required documents are mentioned in Required Document list.

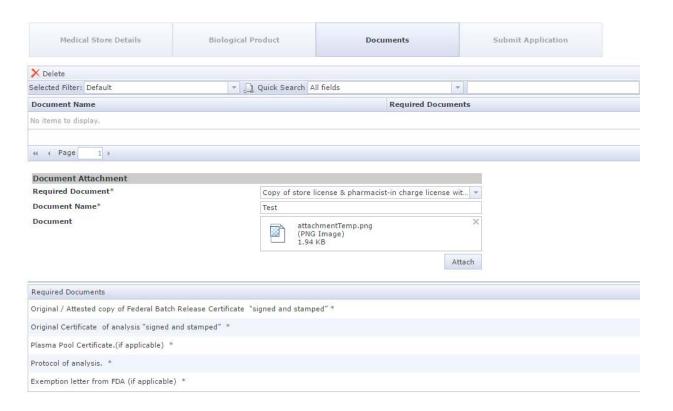


Figure 10 - Documents Tab

Click on Declaration tab and check the check box given and press Submit.



Figure 11 - Declaration Tab

	Page has following actions	
1.	Next	On clicking you will go on next tab

2.	Previous	On clicking you will go on Previous tab
3.	Clear	Clear all the un saved information
4.	Save and close	Saves the application and closes the form

6.2.2. Chemical Precursors

On Module screen select Type Import/Export and select Module Import Controlled Application Chemical Precursors



Figure 12 - Chemical Precursors

• System will load the tabs



Figure 13 - Import Controlled Application Chemical Precursors Tabs

Verify Importing Agent details and press next.

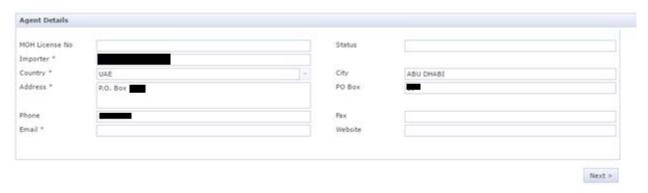


Figure 14 - Importer Details

Select chemical Precursors tab and enter data in all the fields and press
 Save

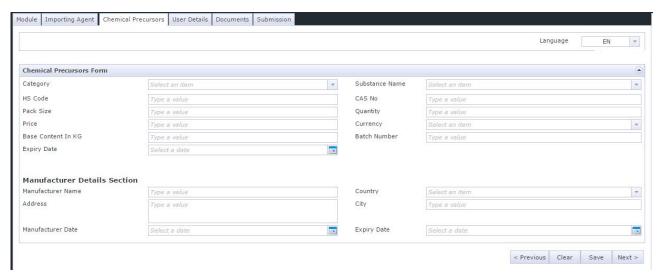


Figure 15 - Chemical Precursors Tab

 Click next or click on User Details and add all the details and click on Documents tab.

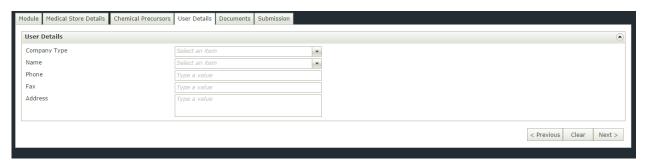


Figure 16 - Chemical Precursor User Tab

- Add all the required documents and click Submission tab. Required
 Document list is mentioned in required documents grid on screen.
- All the added documents will appear in Document grid.
- Enter Document name, select file and upload it.

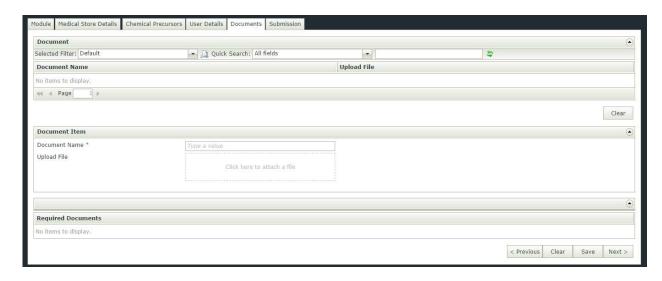


Figure 17 - Chemical Precursor Documents Tab

Click on Submission tab and check the check box given and press Submit.



Figure 18 - Chemical Precursor Submission Tab

	Page has following actions	
	Next	On clicking you will go on next tab
2.	Previous	On clicking you will go on Previous tab
3.	Clear	Clear all the un saved information
4.	Submit	Submit the application

6.2.3. Controlled Medicines (SCD/CD/NP)

On Module screen select Type Import/Export and select Module Import Controlled Medicines (SCD/CD/NP)



Figure 19 - Import Controlled Medicines (SCD/CD/NP)

System will load the Controlled medicines and Narcotics form.

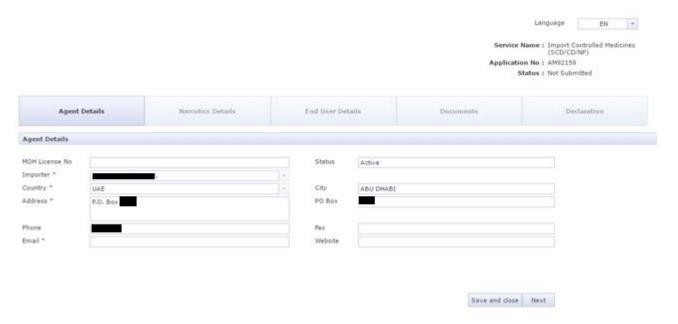


Figure 20 - Controlled medicines and Narcotics form

Verify the Agent details and click next.

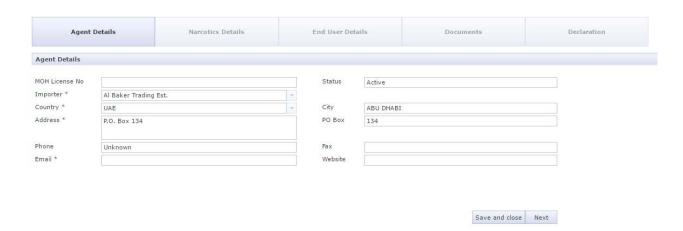


Figure 21 - Agent Details

Fill in the controlled or semi-controlled medicines details.

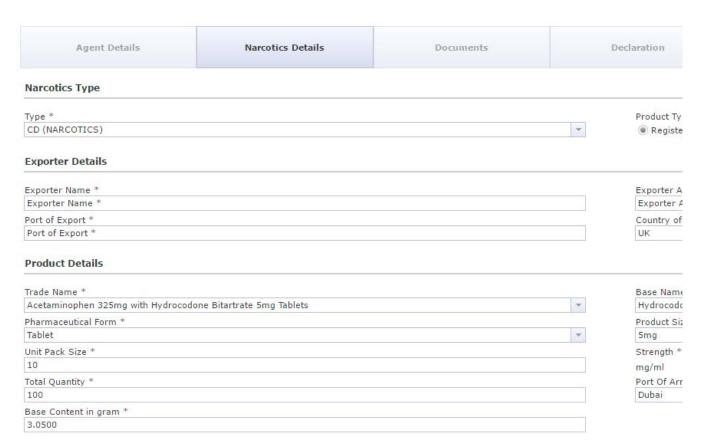


Figure 22 - Narcotic Details

- Attached the necessary documents
- Select Documents tab and add all the required documents and click Submission tab. Required Document list is mentioned in required documents grid on screen.
- All the added documents will appear in Document grid.
- Enter Document name, select file and upload it.

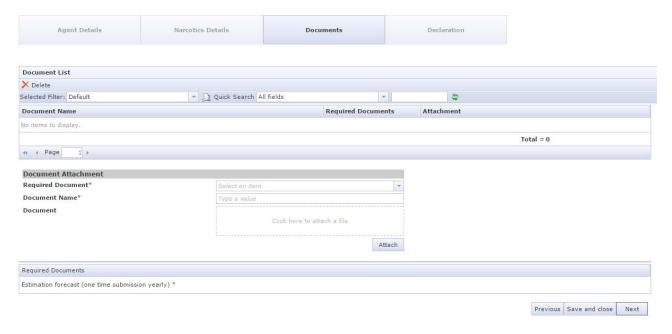


Figure 23 - Controlled Medicine Documents Tab

Click on Declaration tab and check the check box given and press Submit.

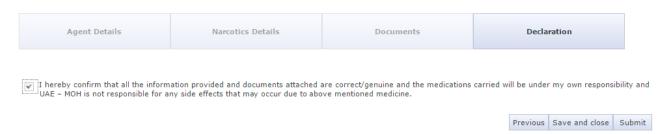


Figure 24 - Controlled Medicine Declaration

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	Page has following actions	
	Next	On clicking you will go on next tab
2.	Previous	On clicking you will go on Previous tab
3.	Save and Close	Saves the data and closes the form
4.	Submit	Submits the application.

6.3. Import/Export Permits Applications

Click Create Application from the following menu:



Figure 25 - Create Permit Application

Click OK button to agree with the Terms and Conditions

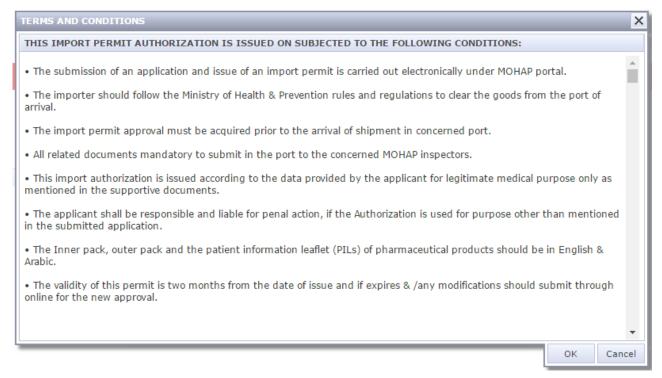


Figure 26 - Permit Terms and Conditions

• On Module screen select Type Import/Export and select desired Module type.

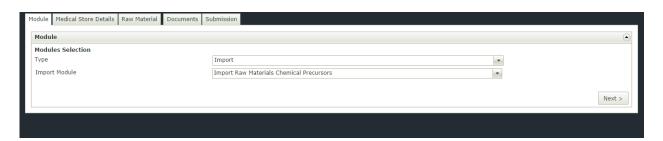


Figure 27 - Import Permit Modules

6.3.1. Import Medicinal Product

On Module screen select Type Import/Export and select desired Module type.



Figure 28 - Import Permit Modules

System will load the tabs



Figure 29 - Medicinal Product Tabs

· Verify Medical and Store Detail and press clear or next.

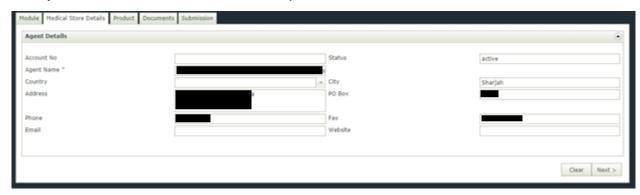


Figure 30 - Medicinal Product Agent Details

- Select Product tab and enter data in all the fields and press save.
- Saved product will be visible in Product list top on the page.
- If you are using already registered Product then select registered product and enter details in MAH and Invoice fields.

- If you are using unregistered product then select unregistered product and enter data in other fields and save the product with entering data in MAH and Invoice section.
- You can add a maximum of 10 (ten) Products in one permit request.

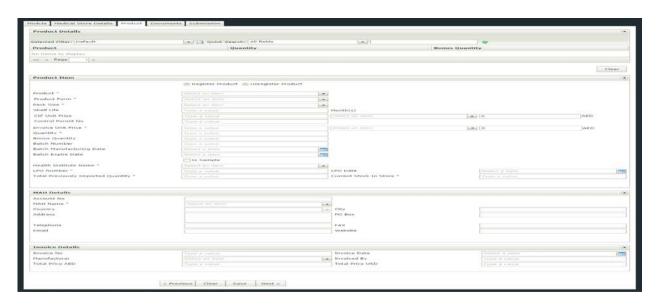


Figure 31 - Medicinal Product Form

- Select Documents tab and add all the required documents and click Submission tab. Required Document list is mentioned in required documents grid on screen.
- All the added documents will appear in Document grid.
- Enter Document name, select file and upload it.

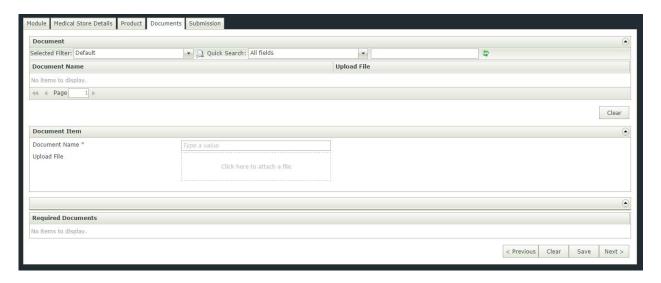


Figure 32 - Medicinal Product Document

Click on Submission tab and check the check box given and press Submit.

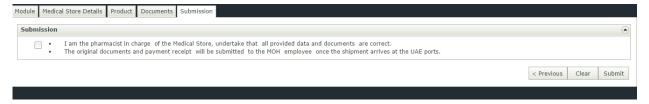


Figure 33 - Import Medicinal Products Permit Submission

System will now ask for a payment. Please refer to the Payment section (6.5.
 Payments) for more details.

	Page has following actions	
	Next	On clicking you will go on next tab
2.	Previous	On clicking you will go on Previous tab
3.	Clear	Clear all the un saved information

4.	Re-Submit	This button will appear if the Drug department sends the
		application back to the agent in case if any requirement is
		missing.

6.3.2.Import Medicinal Devices and Diagnostic Kit

On Module screen select Type Import/Export and select Medicinal Devices and Diagnostic Kit



Verify Medical and Store Detail and press clear or next.

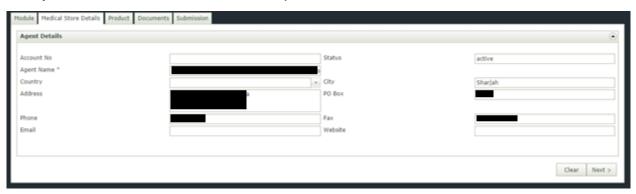


Figure 34 - Medicinal Devices Agent Details

- Select Product tab and enter data in all the fields and press save.
- Saved product will be visible in Product list top on the page.
- You can add a maximum of 10 (ten) products in one permit.

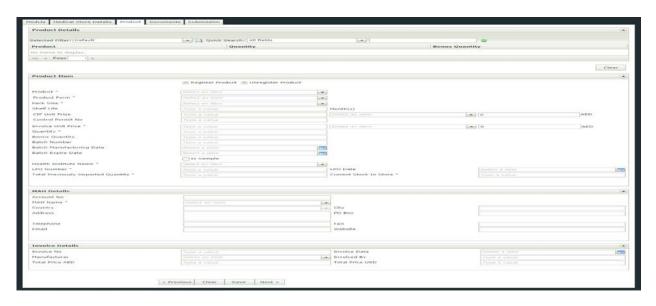


Figure 35 - Medicinal Devices Product Form

- Select Documents tab and add all the required documents and click Submission tab. Required Document list is mentioned in required documents grid on screen.
- All the added documents will appear in Document grid.
- Enter Document name, select file and upload it.

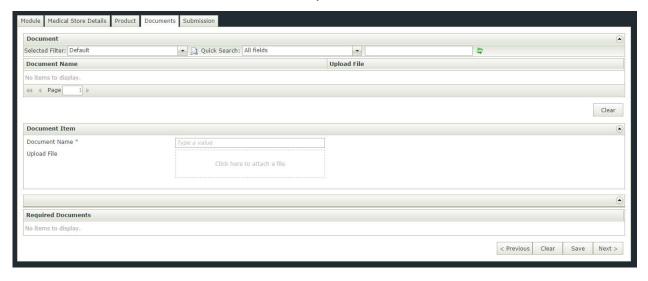


Figure 36 - Medicinal Devices Documents Tab

Click on Submission tab and check the check box given and press Submit.

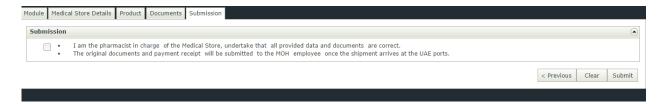


Figure 37 - Import Medicinal Devices and Diagnostic Kit Submission

	Page has following actions	
	Next	On clicking you will go on next tab
2.	Previous	On clicking you will go on Previous tab
3.	Clear	Clear all the un saved information
4.	Submit	Submits the application to the Drug Department

6.3.3. Import Raw Material Permit

On Module screen select Type Import/Export and select Module Import Controlled Application Controlled Medicines

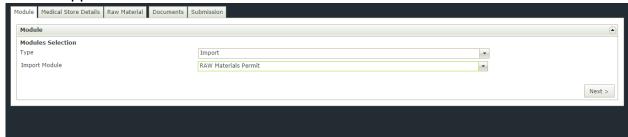


Figure 38 - Import Raw Material Permit

• System will load the tabs



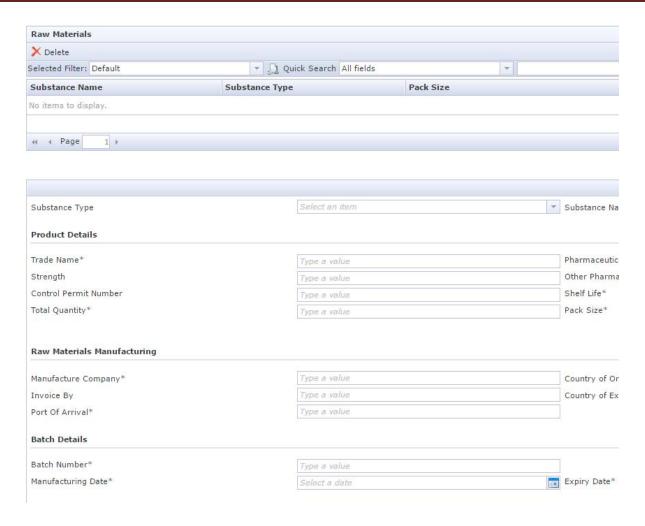
Figure 39 - Raw Material Tabs

Verify the Local Manufacturing details and click next.



Figure 40 - Raw Material Agent Details

- Select Raw Materials Tab. Fill the form and press Save.
- All the saved raw material items will save in Raw material grid.
- You can add a maximum of 10 (ten) substances in one permit.



- Select Documents tab and add all the required documents and click Submission tab. Required Document list is mentioned in required documents grid on screen.
- All the added documents will appear in Document grid.
- Enter Document name, select file and upload it.



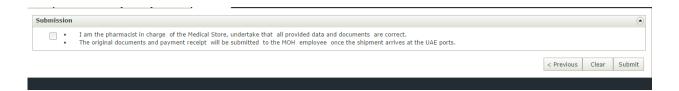


Figure 41 - Import Raw Material Permit Submission

	Page has following actions	
	Next	On clicking you will go on next tab
2.	Previous	On clicking you will go on Previous tab
3.	Clear	Clear all the un saved information
4.	Submit	Submits the application to the Drug Department

6.3.4.Import Chemical Precursors

On Module screen select Type Import/Export and select Module Import Controlled Application Chemical Precursors



Figure 42 - Import Chemical Precursors

System will load the forms

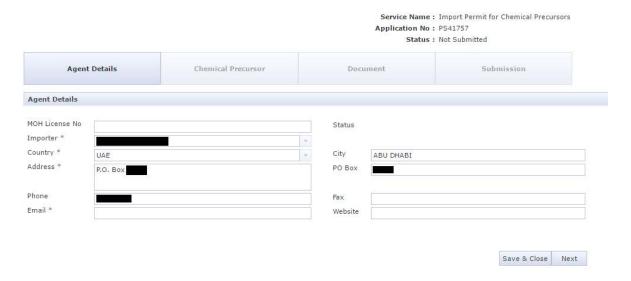


Figure 43 - Import Chemical Precursors Form

Verify the agent details and click next.

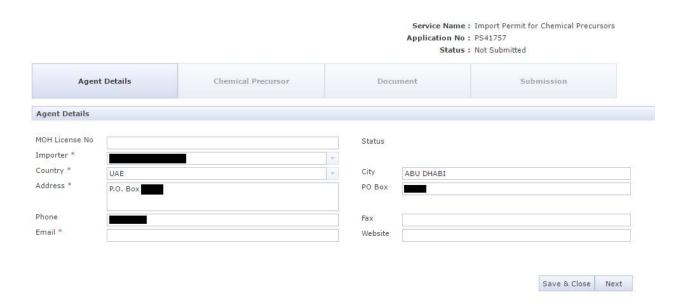


Figure 44 - Medical Store Details Tab

Select chemical Precursors tab and enter data in all the fields and press
 Save

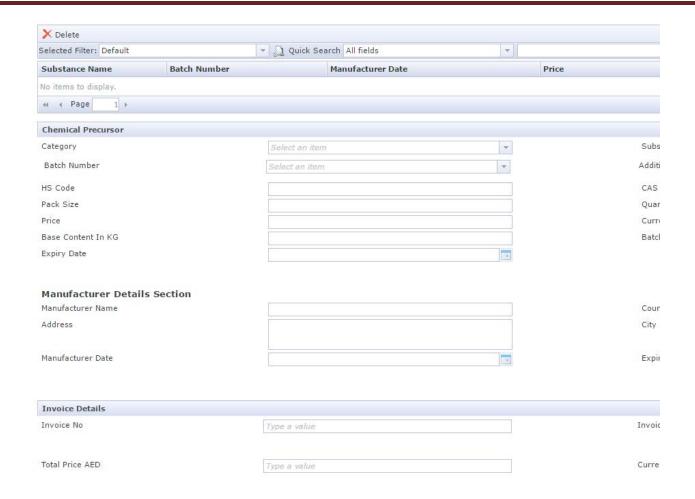


Figure 45 - Chemical Precursor Tab

- System will save the data in Chemical precursor's grid.
- You can add a maximum of 10 (ten) items (pre-approval or authorization of each item will be required prior to adding the items).
- Add all the required documents and click Submission tab. Required
 Document list is mentioned in required documents grid on screen.
- All the added documents will appear in Document grid.
- Enter Document name, select file and upload it.



Figure 46 - Chemical Precursor Documents Tab



Figure 47 - Chemical Precursor Submission Tab

	Page has following actions	
	Next	On clicking you will go on next tab
	Previous	On clicking you will go on Previous tab
2.	Clear	Clear all the un saved information
3.	Submit	Submits the application to the Drug Department

6.3.5. Export Controlled Application Chemical Precursors

 On Module screen select Type Export and select Module Export Controlled Application Chemical Precursors

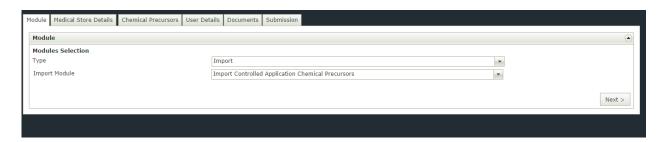
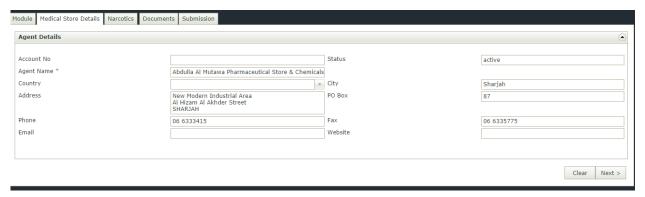


Figure 48 - Export Controlled Application Chemical Precursors

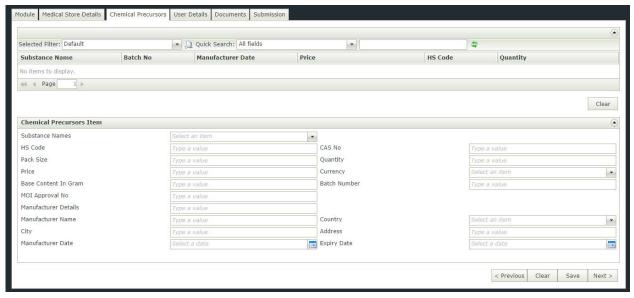
System will load the tabs



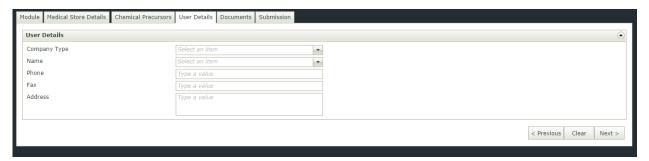
Select Medical and Store Detail and press clear or next.



Select chemical Precursors tab and enter data in all the fields and press
 Save



- System will save the data in Chemical precursor's grid.
- Save as many items here as you want.
- Now click on User Details and add all the details and click on Documents tab.



- Add all the required documents and click Submission tab. Required
 Document list is mentioned in required documents grid on screen.
- All the added documents will appear in Document grid.
- Enter Document name, select file and upload it.

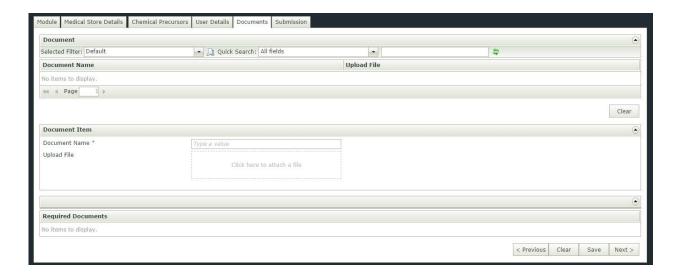




Figure 49 - Export Controlled Application Chemical Precursors Submission

	Page has following actions	
	Next	On clicking you will go on next tab
2.	Previous	On clicking you will go on Previous tab
3.	Clear	Clear all the un saved information

6.3.6. Export Medicinal Devices and Diagnostic Kit

On Module screen select Type Export and select Medicinal Devices and Diagnostic Kit

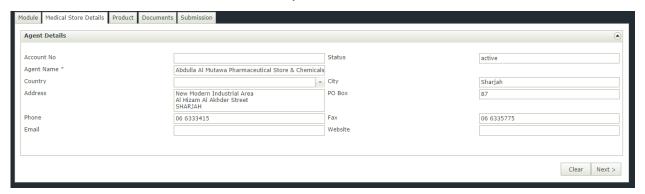


Figure 50 - Export Medicinal Devices and Diagnostic Kit

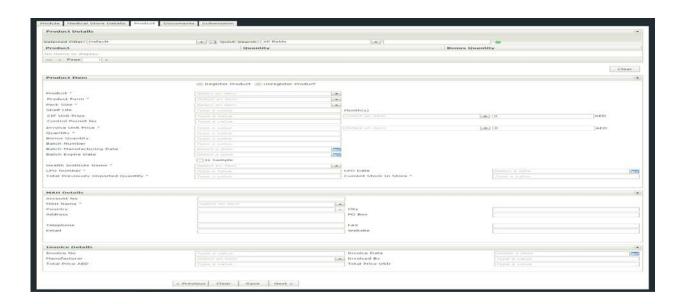
System will load the tabs



Select Medical and Store Detail and press clear or next.



- Select Product tab and enter data in all the fields and press save.
- If you are using already registered Product then select registered product and enter details in MAH and Invoice fields.
- If you are using unregistered product then select unregistered product and enter data in other fields and save the product with entering data in MAH and Invoice section.
- Saved product will be visible in Product list top on the page.
- Add as many products and save then as you want.



- Select Documents tab and add all the required documents and click Submission tab. Required Document list is mentioned in Required documents grid on screen.
- All the added documents will appear in Document grid.
- Enter Document name, select file and upload it.

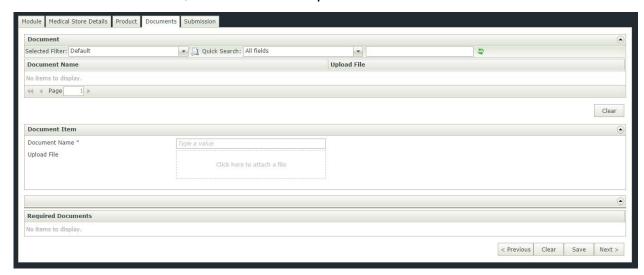




Figure 51 - Export Medicinal Devices and Diagnostic Kit Submission

	Page has following actions	
	Next	On clicking you will go on next tab
2.	Previous	On clicking you will go on Previous tab
3.	Clear	Clear all the un saved information

4.	Submit	Submits the application for payment and approval
5.	Re-Submit	This button will appear if the Drug department sends the application back to the agent in case if any requirement is missing.

6.3.7. Export Medicinal Products Permit

Module screen select Type Export and select Medicinal Devices and Diagnostic Kit

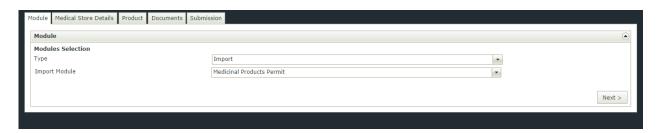
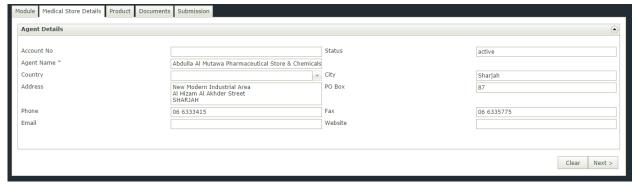


Figure 52 - Export Medicinal Products Permit

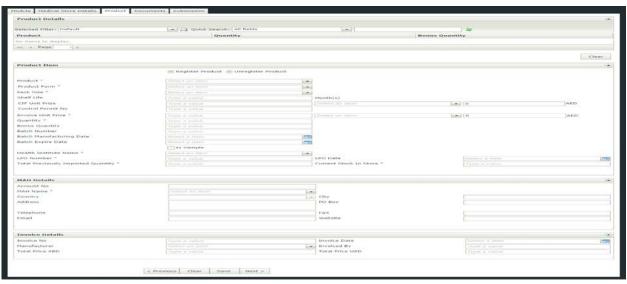
System will load the tabs



Select Medical and Store Detail and press clear or next.



- Select Product tab and enter data in all the fields and press save.
- If you are using already registered Product then select registered product and enter details in MAH and Invoice fields.
- If you are using unregistered product then select unregistered product and enter data in other fields and save the product with entering data in MAH and Invoice section.
- Saved product will be visible in Product list top on the page.
- Add as many products and save then as you want.



- Select Documents tab and Add all the required documents and click Submission tab. Required Document list is mentioned in Required documents grid on screen.
- All the added documents will appear in Document grid.
- Enter Document name, select file and upload it.

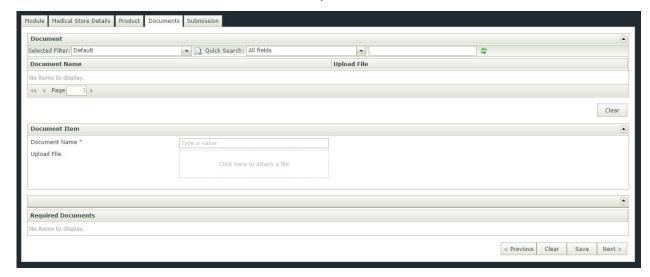




Figure 53 - Export Medicinal Products Permit Submission

	Page has following actions	
	Next	On clicking you will go on next tab
2.	Previous	On clicking you will go on Previous tab
3.	Clear	Clear all the un saved information
4.	Submit	Submits the application for payment and approval
5.	Re-Submit	This button will appear if the Drug department sends the application back to the agent in case if any requirement is missing.

6.3.8. Export Local Manufactured Products

 Module screen select Type Export and select Export Local Manufacturing Product.

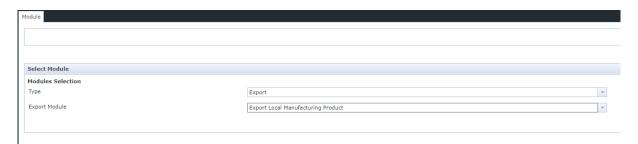


Figure 54 - Export Local Manufacturing Product

• System will load the tabs



Figure 55 - Local Manufactured Product Application Form

Review the agent details and click Next.

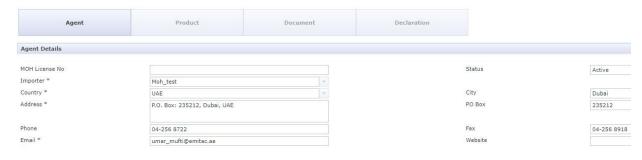


Figure 56 - LMP Agents Details

- Select Product tab and enter data in all the fields and press save.
- If you are using already registered Product then select registered product and enter details in MAH and Invoice fields.

- If you are using unregistered product then select unregistered product and enter data in other fields and save the product with entering data in MAH and Invoice section.
- Saved product will be visible in Product list top on the page.
- Add as many products and save then as you want.

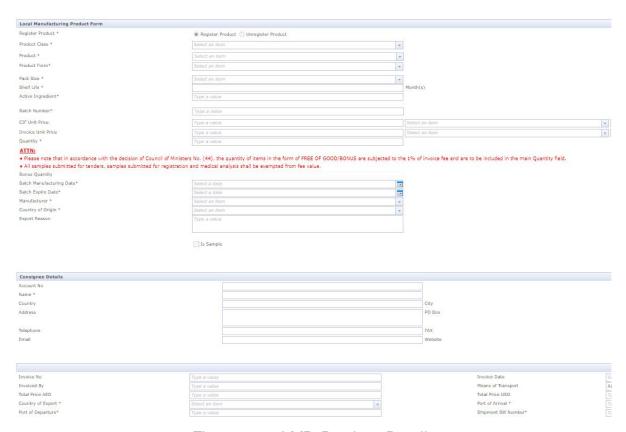


Figure 57 - LMP Product Details

- On Documents tab and add all required document. Required Document list is mentioned in required documents grid on screen.
- All the added documents will appear in Document grid.
- Enter Document name, select file and upload it.

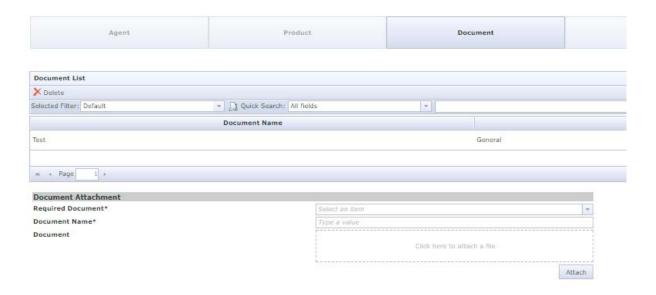


Figure 58 - LMP Documents



Figure 59 - LMP Declaration and Submission

Page has folk	Page has following actions	
Next	On clicking you will go on next tab	
Previous	On clicking you will go on Previous tab	
Clear	Clear all the un saved information	
Submit	Submits the application for payment and approval	
Re-Submit	This button will appear if the Drug department sends the application back to the agent in case if any requirement is missing.	

6.3.9. Export Raw Material Permit

On Module screen select Type Export and select Module Export Controlled Application Controlled Medicines

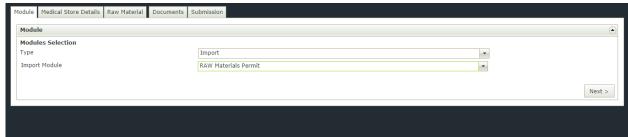
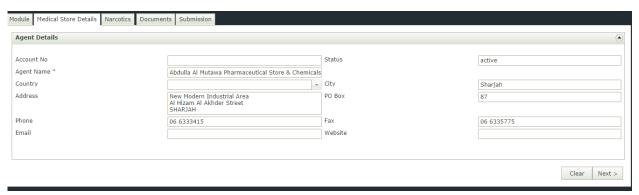


Figure 60 - Export Raw Material Permit

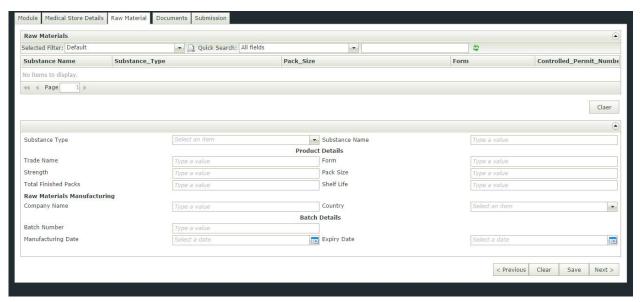
System will load the tabs



Select Medical and Store Detail and press clear or next.



- Select Raw Materials Tab. Fill the form and press Save.
- All the saved raw material items will save in Raw material grid.
- You can save as many items as you want.



- Select Documents tab and Add all the required documents and click Submission tab. Required Document list is mentioned in Required documents grid on screen.
- All the added documents will appear in Document grid.
- Enter Document name, select file and upload it.

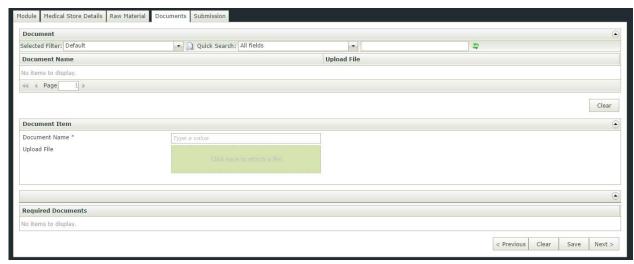




Figure 61 - Export Raw Material Permit Submission

Page has following actions		
	Next	On clicking you will go on next tab
2.	Previous	On clicking you will go on Previous tab
3.	Clear	Clear all the un saved information

6.4. Shipment Clearance Applications

Click Create Application from the following menu:



Figure 62 - Create Shipment Clearance Application

Click OK button to agree with the Terms and Conditions

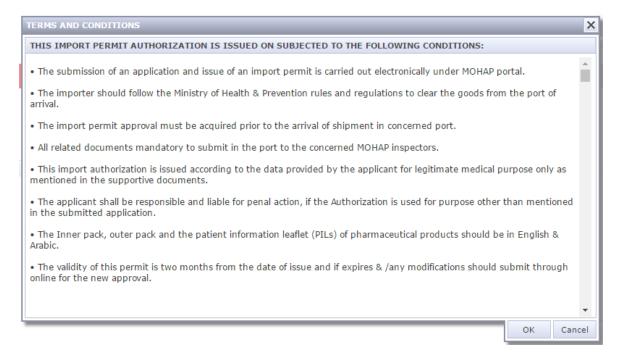


Figure 63 - Terms and Conditions

 On the Shipment Clearance menu, select the desired module/product type for which the shipment clearance is required.

Welcome to Shipment Clearance System



Figure 64 - Shipment Clearance Modules

- Select the Approved Permit to be added and Click Add button to include it in the shipment clearance request.
- You can add multiple permits in one clearance request.

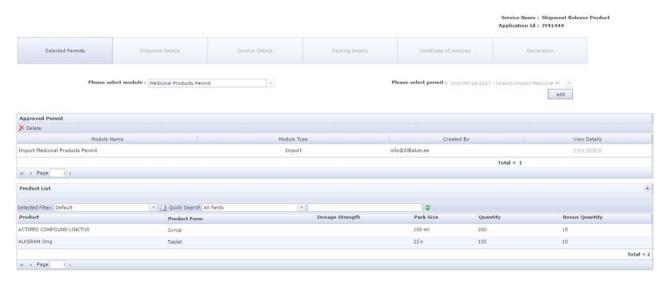


Figure 65 - Shipment Clearance Add Permits

• Double click each permit in the Approve permit list to list its items.

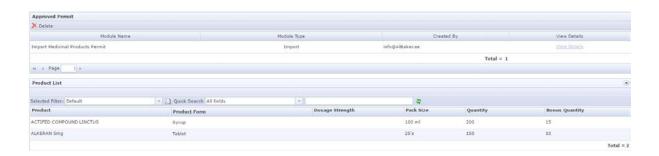


Figure 66 - Permit Items

 Double click each item to view and verify its details. The details layout will change depend on the type of the item, such as, Medicinal Product, Raw Material, etc. and then click next.

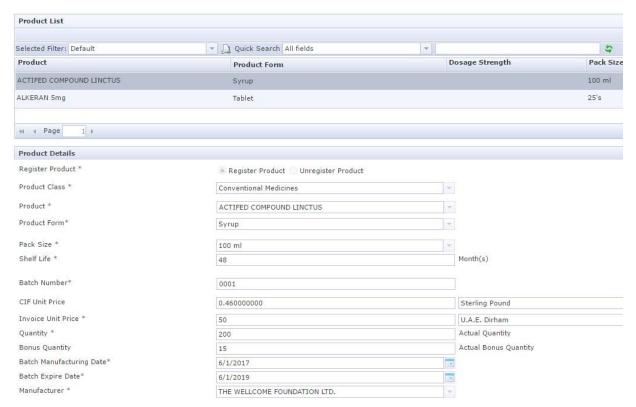


Figure 67 - Permit Item Details

 Fill in the shipment details and add the relevant shipment documents and click Next button.

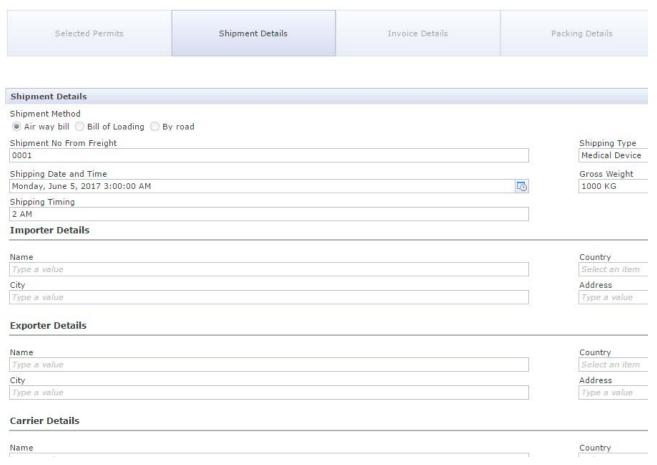


Figure 68 - Shipment Details

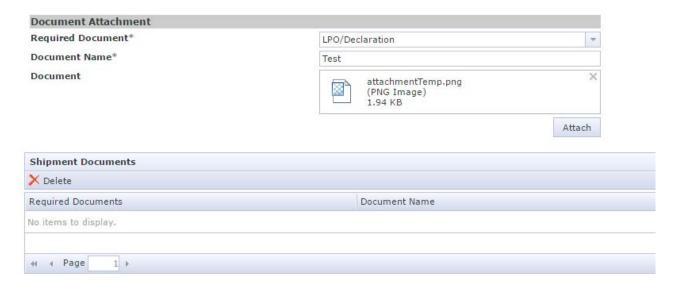


Figure 69 - Shipment Documents

 Fill in the invoice details with relevant attachments for each associated invoice and Click Save Invoice button to add an invoice.

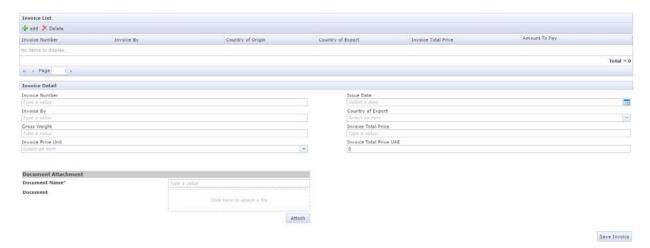


Figure 70 - Invoice Details

You can also update or delete an existing invoice from the invoice list.



Figure 71 - Invoice List

- Click next when all invoices are done.
- Fill in the packaging details and attach the relevant documents and click
 Save Packaging Info button to save.

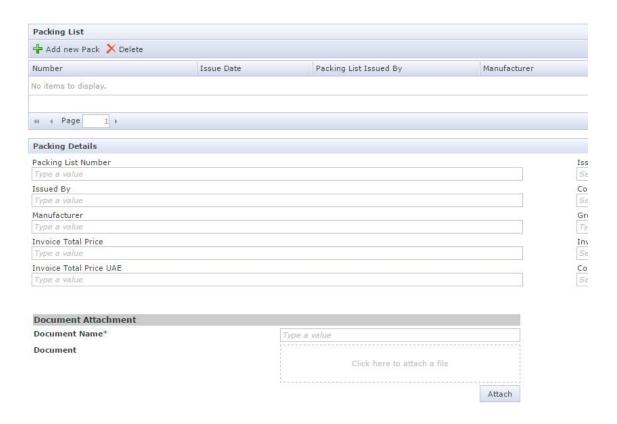


Figure 72 - Packaging Details

 You can also update or delete an existing packaging detail from the Packaging list.



Figure 73 - Packaging List

• Click next when all packaging information is done.

 Fill in the Certificate of Analysis (COA) details for each applicable item in the selected permit and attach the relevant documents and click Save COA button to save.

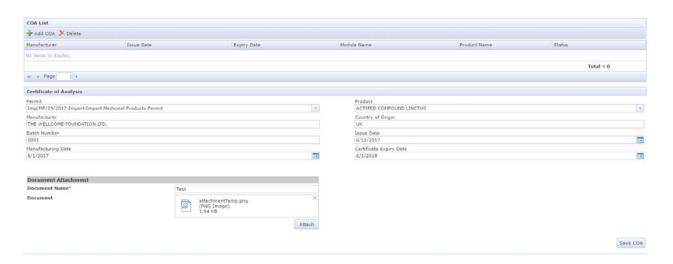


Figure 74 - Certificate of Analysis

You can also update or delete an existing COA from the COA list.



Figure 75 - COA List

- Click next when all COA are done.
- Confirm the declaration and click Submit button to submit the request to Drug Department.

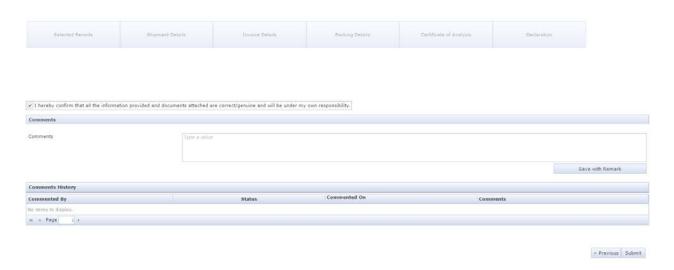


Figure 76 - Shipment Declaration

6.5. Payments

- For Import/Export Pre-Permits and for Shipment Clearance requests, the System will redirect the user to online Payment (e-dirham) website, whenever a payment is applicable.
- The applicant shall be shown the following message before redirecting to the payment website:
 - "Please note that your application is not submitted yet. Once you have completed the payment process, you shall be redirected back to the Drug Import Export eservice website for submitting your request (when the payment is processed successfully).

You shall now be redirected to the E-Dirham website for payment."

Select the payment method





Figure 77 - E-Dirham Payment Method

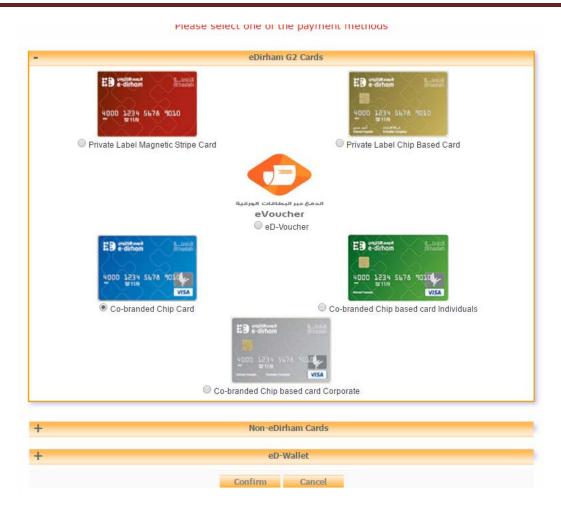


Figure 78 - E-Dirham Payment Method Selection

Review amount and click confirm to continue





Figure 79 - Payment Method Confirmation

 Provide the payment credentials (Card Number and Code, etc.) and click submit.



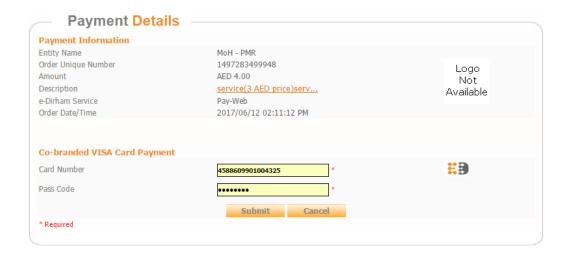


Figure 80 - Payment Credentials

Save the payment receipt as proof (optional), and click OK.

Transaction Id:
111111123002609
Service Provider:
Ministry of Health
Payment Result :
Receipt Id:
Date:
Date .
Payment Id:
1497283499948
Service 1
e Service Description :
The value of import permit for any of the medical products or raw materials or substances, chemical precursors
Ok
Copyright © 2014, Ministry of Health. All rights reserved.

Figure 81 - Payment Receipt

- You shall be redirected to the Drug Import Export System
- Select the Application for which you made the payment; the application will be with "Check Payment Status" status.



Figure 82 - Pending Payment Confirmation

You will be taken to the Payment Status Form

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Home | Payment Status You shall be able to Submit your request once your payment is successful: . Click "Check Payment Status" button to get the latest status of your payment. . Click "Submit Request" button once the Payment Status becomes "Successful". . Click "Retry" button if the Payment Status becomes "Failed". Order Reference Number Shp/MP/14/2017 Amount to Pay (AED) 200.00 Payment Status Failed Transaction Status Failed Check Payment Status Retry

Figure 83 - Payment Status

- Click "Check Payment Status" button to get the latest status of your payment.
- Click "Submit Request" button once the Payment Status becomes "Successful".
- Click "Retry" button if the Payment Status becomes "Failed".
- Once you click the submit button, your application shall be submitted to the Drug Department for processing.

6.6. View Applications

Click on View Application system will open the task list.

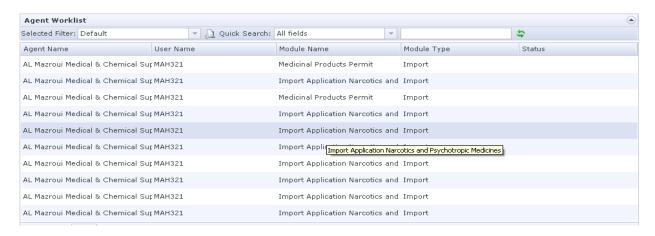


Figure 84 - View Applications in Workspace