

## External User Guide

### Issue of a Certificate of Free Sale of a Medical Product for Export

## Table of Contents

1. Brief Overview .....	3
2. Apply for Service .....	3
2.1 Create a New Application .....	4
2.2 Free Sale Certificate Application .....	4
2.2.1 Product Details.....	5
2.2.2 Pack Details.....	6
2.2.3 Free Sale Request Detail .....	7
2.2.4 Attachments.....	7
2.3 Review Application .....	8
2.4 Checkout .....	9
3. Take Required Actions .....	10
3.1 Application Correction.....	10
4. Rejected Applications.....	12
5. Printouts .....	12

## 1. Brief Overview

This manual is prepared to navigate applicants through the process of issuing the Certificate of Free Sale of a Medical Product for Export (FSC) for product packs in Emirates Drug Establishment (EDE) Portal.

This manual is designed to help applicants:

- Apply for a Certificate of Free Sale of a Medical Product for Export (FSC) for their medical products
- Manage Certificate of Free Sale of a Medical Product for Export (FSC) applications

After completing this manual, the applicant should be able to perform all activities related to FSC.

Please note: This service is only available to the Applicants listed below.

Applicant Type	Rules
<b>Agents</b>	Should have un-expired license Can only issue FSC for drugs registered under his name as agent
<b>Scientific Offices</b>	Should have un-expired license Can only issue FSC for drugs registered under his name as Applicant or MAH
<b>Local Manufacturers</b>	Should have un-expired license Can only issue FSC for drugs they (Manufacturer) are a part of in the manufacturing cycle

TABLE 1: TYPES OF APPLICANTS

## 2. Apply for Service

Portal users who have logged in successfully to EDE portal will be directed to Dashboard screen where they can apply to EDE's e-services. This user manual will focus on the FSC service.

For applicants to apply to their available services, they must go through the following process:

1. Create a new application
2. Complete application
3. Review application
4. Proceed to checkout

The sections below will help user through the step-by-step process of applying to an application.

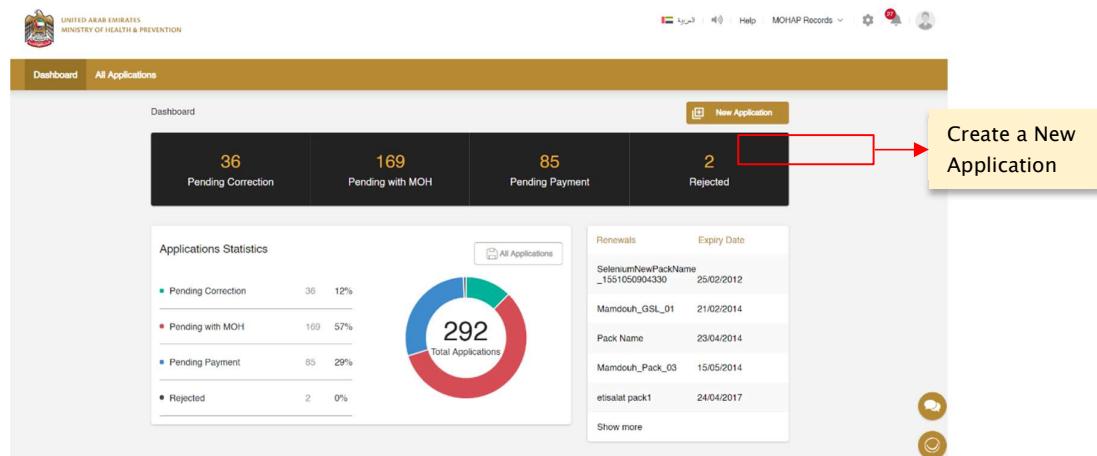


FIGURE A: DASHBOARD SCREEN

## 2.1 Create a New Application

The Applicant can request to classify their products based on EDE standards by:

Requesting to create a New Application form from the Applicant Portal and selecting the following service which falls under the **Drug, Certificates** services:

- Certificate of Free Sale of a Medical Product for Export (FSC)

To create a new application form from the Applicant Portal, the user must do the following:

- a. Click on one of the **New Application** buttons found on the Dashboard Screen (or All Applications Screen) – which opens New Application Screen
- b. Select **Free Sale Certificate**
- c. Click **Create Application** button

Once the user selects **Free Sale Certificate** as a service and creates an application, the user will be redirected to Product Details screen.

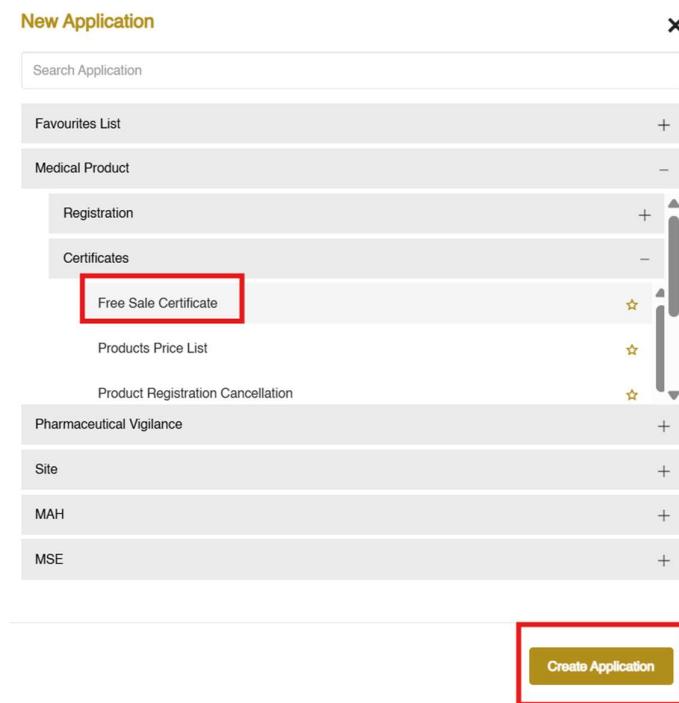


FIGURE B: NEW APPLICATION SCREEN

## 2.2 Free Sale Certificate Application

To complete a Free Sale Certificate application, the applicant must fill the following three sections:

1. Product Details
2. Free Sale Request
3. Attachments

At any application stage, the applicant can perform the following actions:

- ❖ Save application

- ❖ Discard application
- ❖ Proceed to next section
- ❖ Go back to previous section

For an applicant to move from one section to another, they are required to fill the section's mandatory fields and tables.



FIGURE C: PROGRESS BAR

Warnings:

- In case the applicant did **not** complete a mandatory field, a warning message marked in red will be displayed under the field text box highlighting the note: **This Field is Required**.
- In case the applicant did **not** fill a table, a warning message marked in red will be displayed under the table box highlighting the note: **Table Cannot be empty**.
- In case the applicant inputs a non-numeric character in a field that only accepts numbers, a warning message marked in red will be displayed under the field text box highlighting the note: **This field accepts numbers only**.

Display icons:

- allows user to Delete data (usually found under Actions)
- allows user to Edit data (usually found under Actions)

Tips: Each application section will contain a progress bar that shows the input progress of the application data

### 2.2.1 Product Details

In the Product Details screen, the applicant is required to search and select the Medical Product's Name / Trade Name. Consequently, the screen will display the product details along with the pack list for the product selected.

To proceed with Free Sale Certificate request, the applicant must click on icon under Registered Packs to proceed to the Pack Details screen.

The screenshot shows a software interface for managing medical product registrations. At the top, a progress bar indicates '0% Completed'. Below it, three steps are listed: 'Product details' (step 1), 'Free Sale Request' (step 2), and 'Attachments' (step 3). A note in a yellow box states: 'All fields are mandatory, except for those labeled as optional.' A search bar is present with the text 'Select a product ..' and 'Products Name / Trade Name (Optional)' containing 'LORALIFE'. Below the search bar, detailed product information is shown: 'Products Name / Trade Name (Optional)' (LORALIFE), 'Product Class (Optional)' (Conventional Medicines), and 'Product Subclass (Optional)' (N/A). The 'Active Ingredients' section contains a table with one row: 'Active Ingredient Name' (loratadine), 'Description' (loratadine), and 'Strength' (N/A). The 'Body System' section contains a table with one row: 'Body System' (RESPIRATORY SYSTEM) and 'Body Subsystem' (ANTIHISTAMINES, HYPOSENSITISATION, ALLERGIC E). The 'Registered Packs' section contains a table with one row: 'Pack Name' (Loralife 10mn), 'Strength' (10mg/tablet), 'Pack Sizes' (2), 'Registration Date' (25/05/2003), 'Expiry Date' (05/02/2028), and an 'Actions' column with a red box around the edit icon.

FIGURE D: PRODUCT DETAILS SCREEN

Rule ID	Rules
R1	The selected pack's MAH Country should be UAE.
R2	The selected pack should be partially/fully manufactured in UAE, i.e. if it's manufactured by one manufacturer, its country should be UAE, and if there are many manufacturers, at least one of them, its country should be UAE.
R3	The selected pack's Product Class should <b>not</b> be Medical Device.
R4	Pack registration should be non-expired.

FIGURE E: PRODUCT DETAILS RULES

### 2.2.2 Pack Details

The Pack Details section is broken down into many subsections, each containing essential details of the registered pack. Below are the following subsections:

- General Information
- Package Insert Details
- Product Form
- Active Ingredients\*
- Inactive Ingredients\*
- Routes List
- Market Authorization Holder
- PV Officer\*

- i. Target Animals\*\*
- j. GCC Registration
- k. Registration Status Country of Origin
- l. Registration Status Other Countries
- m. Patent Status
- n. Pack Sizes
- o. Additional Information (Optional)

The landing page of the Pack Details section is the General Information sub-section. To proceed to the next screen, please proceed to click on the **Next** button.

Please note: This fields/attachments in this section are Read-Only.

### 2.2.3 Free Sale Request Detail

For the applicant to proceed with the CPP request, they are required to provide the following information:

- a. Importing Country
- b. Remarks

Once the applicant provides the section's required information, they can move to the next section by clicking the **Next** button.

FIGURE F: FREE SALE REQUEST DETAILS SCREEN

### 2.2.4 Attachments

This section is where an applicant can upload General Attachments before proceeding to submit the application.

The user can add more attachments to their CPP application by performing the following:

1. Enter the Document Name
2. Click the Add Attachment icon 
3. Upload Attachment

Once the applicant is done with this section, they can move to the Review section by clicking **Review** button.

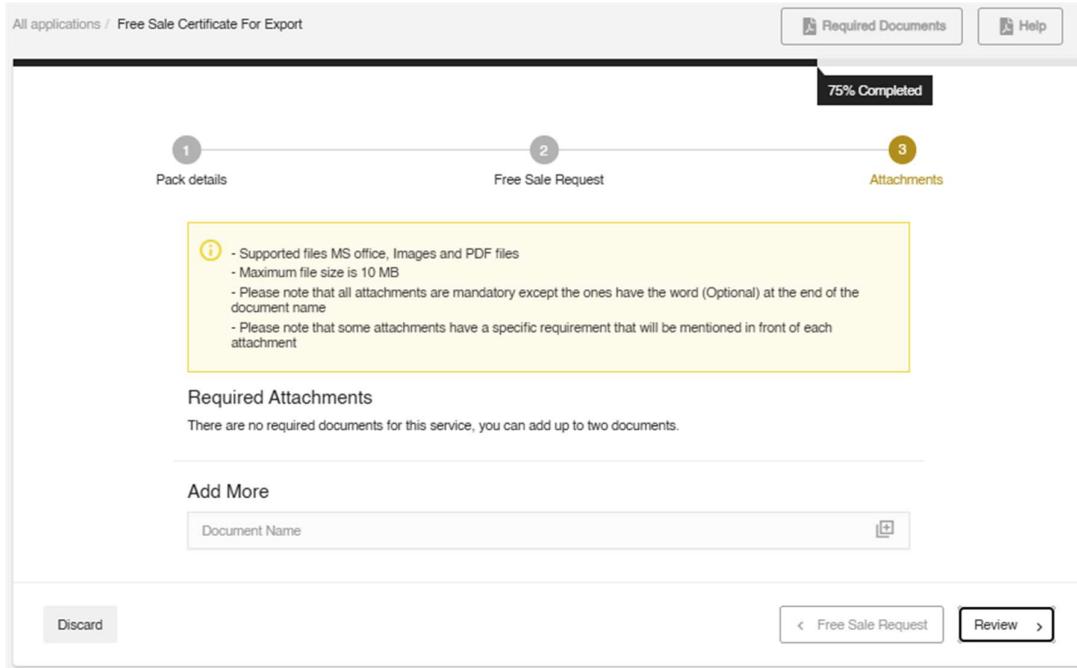


Figure G: Attachments Screen

## 2.3 Review Application

After the applicant completes filling in all application details, they must click on the **Review** button to proceed with application submission. The Review section provides a full summary of the application and allows the user to perform the following actions:

- ❖ Edit Application
- ❖ Save Application
- ❖ Discard Application
- ❖ Go Back to Previous Section
- ❖ Read & Accept General Terms & Conditions
- ❖ Submit Application

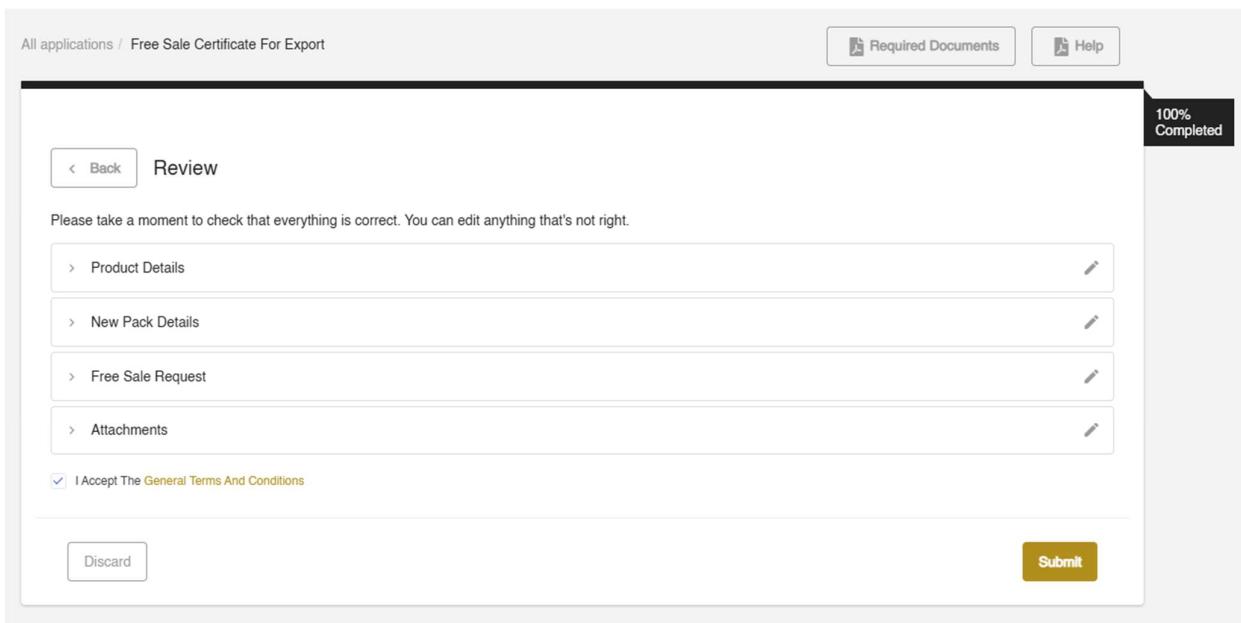
Once the applicant reviews application and accepts the General Terms and Conditions, they must click the **Submit** button to proceed to checkout.

Please note: to proceed with application submission, the applicant must read & accept the General Terms and Conditions.

Warning: In case the user did **not** fill all mandatory fields, a warning message will be displayed noting that applicant must fill all required fields before submitting the application.

Display icon:

-  allows user to Edit application details
-  allows user to view table



All applications / Free Sale Certificate For Export

Required Documents Help

100% Completed

< Back Review

Please take a moment to check that everything is correct. You can edit anything that's not right.

> Product Details 

> New Pack Details 

> Free Sale Request 

> Attachments 

I Accept The [General Terms And Conditions](#)

Discard Submit

Figure H: Review Screen

## 2.4 Checkout

After an application has been submitted, the applicant will be redirected to Checkout screen where they will be asked to pay the following:

- Application Fees: AED 1000 per Application

To proceed with payment, the applicant must click on the **Checkout** button shown at the bottom-right corner of the Checkout screen which will redirect the applicant to EDE's Payment Gateway (e-dirham).

Tip: Overrun Pop-up Blocker to open EDE's Payment Gaterway (e-dirham)

[back](#)

By submitting your application, you are obliged to pay the following fees:

price	
Issuing Of Certificate CPP	1000
Total	1000



Figure I: Checkout Screen

### 3. Take Required Actions

Once an application has been reviewed by a EDE officer, the officer might request the applicant to take actions before proceeding with application. The applicant will get notified via SMS or email once the officer requires an action.

For a user to view and take these actions, they must:

- Click on the **Application Number** in All Applications screen – which redirects user to Submitted Application screen
- Click on **Required Actions** tab
- Click on the Action – which opens an Action window

Figure J: Submitted Application Screen

The request the officer can ask an applicant is the following:

- Application Correction

This action will be explained in detail in the following sections.

#### 3.1 Application Correction

In case a EDE officer identifies fields/attachments in application that require modification, the officer will send back the application to applicant for correction as part of the Drug Classification reviewing process.

For the applicant to view and change the fields/attachments that require correction, the applicant must:

- Click on **Correction** button found in Required Actions tab – which redirects applicant to Correction screen
- Make changes to fields/attachments\*
- Add comment in Comment box

- d. Click on **Submit Correction** button
- e. Confirm correction

Once the applicant confirms correction, the application will be sent back to officer for further review.

Rule: \*Only fields/attachments needed for correction are displayed for change

Pending Correction

Application Type: Certificate of Pharmaceutical Products (CPP)  
Submitted On: 27/03/2019

Application details    Required actions (1)    Certificates    Application History

DRCPP-2019-000072  
Correction  
Mar 27, 2019

Figure K: Submitted Application Screen – Required Actions



Figure L: Field Correction Cursor

Original Composition Certificate Signed By Authorized Person (Optional)

Attachment 2.txt

Insert Which is Last Approved By MOHAP To Be Signed By Authorized Person (Optional)

Insert which is last approved by MOHAP to be Signed by Authc. X

Figure M: Modified Field

Figure N: Unmodified Field

Attachments

2/2 Resolved Request(s)

All fields are mandatory, except for those labeled as optional.

Required Attachments

Insert Which Is Last Approved By MOHAP To Be Signed By Authorized Person (Optional)

Attachment 1.txt

Was Attachment

Original Composition Certificate Signed By Authorized Person (Optional)

Attachment 2.txt

Was Attachment

Comments:

Comment

Corrected the Attachments as required.

Click to Submit Correction

Figure O: Correction Screen

Display icons: allows the user to undo change

**Tips:**

- i. Field Correction Tool helps user navigate from one unsolved request to another
- ii. Fields required for change are highlighted in **Red**
- iii. Modified fields are highlighted in **Yellow**.

## 4. Rejected Applications

Once an application has been rejected by a EDE Officer, an email or an SMS will be sent to the applicant containing the application's latest updates and results.

To view the rejection comments given by EDE Officer and the step at which the application got rejected, please click on the **Application History** button.

Date	Action	Comment
09-04-2019	Payment Received	Transaction was processed successfully.
09-04-2019	submit	
09-04-2019	Approve	Approved

**Figure P: Application History**

To resubmit a rejected application, please perform the following actions:

- a. Click on  icon shown in the All Applications screen (or Submitted Application screen) – which redirected to Patient Details screen
- b. Make changes based on EDE Officer's rejection comments
- c. Proceed to submit application

**Figure Q: Submitted Application Screen – Application Details**

## 5. Printouts

Once an application has been reviewed and approved by a EDE officer, the officer will issue the applicant a Certificate associated with the service type. The applicant will get notified via SMS or email once the officer issues the certificate.

For a user to view and printout a certificate, they must:

- a. Click on the **Application Number** in All Applications screen – which redirects user to Submitted Application screen

- b. Click on the **Certificate** tab
- c. Click on the Attachment – which opens the certificate as a PDF
- d. Print PDF

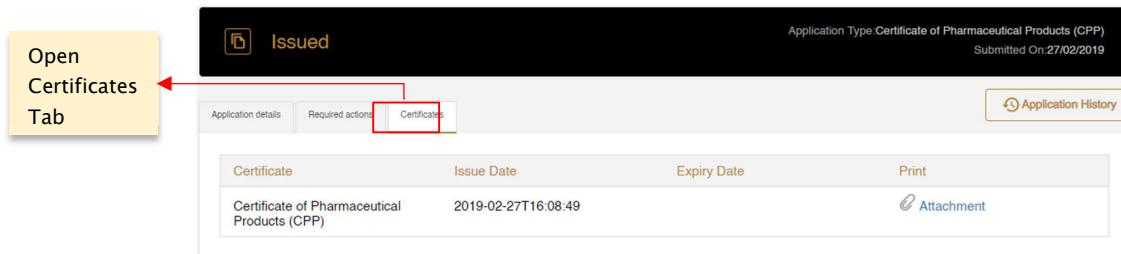


Figure R: Submitted Application Screen – Printout