

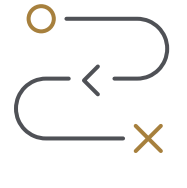


Issue of Permit to Import Medicines for Exhibitions

This service allows obtaining a permit to import medicinal products, narcotic drugs, controlled or semi-controlled drugs, or pharmaceutical preparations for the local agent holding a valid medical store license issued by the Ministry of Health and Prevention.

 Department name Drug	 Sector Health Regulation	 Main Service Clearance, Import and Export Permits	 Service Code 110-02-004-000
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 Service Classification Transactional	 Variation / Auxiliary Variation	 Service Type Government to Business Government to Government
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Service Process

- 01 To apply for e-services, create an account on MOHAP website or smart app with a username and password.
- 02 Submit the application online with the required documents.
- 03 The employee concerned at the Ministry of Health and Prevention will evaluate the application.
- 04 The permit will be automatically issued by the system and the customer can download the file.



Required Documents

- A letter from the exhibition management mentioning the name of the company participating in the exhibition.
- An undertaking from the importer to re-export the products after the exhibition and not to sell them.
- Invoice for products from the factory (manufacturer) or authorized distributor (a relationship letter from the manufacturer is required).
- Information or catalogue of products to be imported.
- Quality certificates from the product manufacturer (EC & ISO or FDA Certificate).
- The commercial license of the intermediary company / importer.
- Airway bill or bill of lading (shipment must be refrigerated if shipped by sea).



Requirements & Conditions

- The purchase invoice should include country of origin, batch/serial number and dates of production and expiry (the validity date must be more than two-thirds of the product validity period).
- The product(s) are only to be displayed in the mentioned event and a clear statement regarding their status as unregistered non-priced should be displayed at the event.
- Any advertisement for the medical products for exhibition in local media is not allowed.
- It is prohibited to import products that contain alcohol or any ingredients that are internationally banned or controlled by the Ministry of Health and Prevention (such as chemical precursors, drugs, psychotropic substances, tissues, human or animal blood, and products that contain other biological components).
- It is prohibited to sell, use or distribute the exhibition products.

Service Channels



MOHAP Website: www.mohap.gov.ae



MOHAP Smart App

Resources

- [User Manual](#)



Average Service Time
1 working day



Payment channels
E-Payment



Target Audience
Global pharmaceutical factories, local pharmaceutical factories, marketing offices, drug stores, companies wishing to display their products at exhibitions



Service Locations

- MOHAP Website
www.mohap.gov.ae
- MOHAP Smart App



Related Services
Commercial license from Departments of Economy for the entity.

Import permit from the Federal Authority for Nuclear Regulation (FANR) for radiation devices.

Inspection before customs release of shipments and before distribution locally in the market (Department of Control, Audit and Inspection - MOHAP) for all medical products.

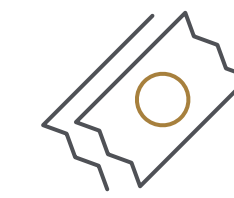


Service Bundle
This service is not linked to any bundles



Contact Details
Technical support
import.export@mohap.gov.ae

IT support
Applications.support@mohap.gov.ae



Service Fees

Application fee: AED 100

Entry Permit fee for medical products: AED 1,000

Sustainable Development Goals



Notes

The permit is valid for two months, and is only valid for displaying within exhibitions/events.

The medical products /medicines must be re-exported abroad as soon as the exhibition/event is completed.

FAQs

How much is the fee for issuing a permit to import medicines for exhibitions service?

Application fee: AED 100

Medical Products Entry Permit Fee: AED 1000

Should medical products imported for the purpose of exhibitions be re-exported?

Yes