






Evaluation of Mandatory Reports of Adverse Localized Negative Reactions to Drugs

This service enables companies to request the evaluation of mandatory reports on the side effects and adverse reactions to local and internationally manufactured medicines imported into the UAE.

 Department name Drug	 Sector Health Regulation	 Main Service Pharmaceutical vigilance	 Service Code 110-45-003-000
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 Service Classification Transactional	 Variation / Auxiliary Variation	 Service Type Government to Business
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Service Process

- 01 Login to the MOHAP website or smart application to apply for e-services and send the report.
- 02 Review documents and requirements.
- 03 The report will then be sent to the World Health Organization (Uppsala Monitoring Center).
- 04 If necessary, circulars will be issued regarding the report.



Required Documents

- No documents required



Requirements & Conditions

- The report should list all required patient data.
- The report should state all required data of the writer.
- The report should define the required drug data.
- The report should clearly state the side effects experienced from the drug.

Service Channels



MoHAP Website: www.mohap.gov.ae



MoHAP Smart App and UAE RADR

Resources

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Average Service Time

10 working days



Payment channels

None - The service is free



Target Audience

Scientific offices, medical stores, local and international pharmaceutical factories, patients, health care practitioners, health institutions



Service Locations

- MoHAP website
www.mohap.gov.ae
- MoHAP Smart Application



Related Services

This service is not linked to other services



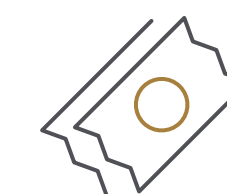
Service Bundle

This service is not linked to any other bundle



Contact Details

Email:
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Service Fees

Free

Sustainable Development Goals



Notes

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FAQs

None