

Circular No.: (15) 2026

To: Authorized Representatives (AR's)

Subject: Updated Fee Schedules of services and requests for medical devices and products in the National Health Regulatory Authority

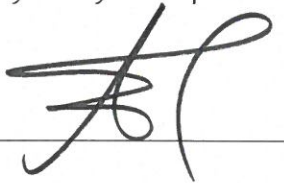
In reference to Resolution No. (24) of 2021, amending certain provisions of Resolution No. (20) of 2016 on the categories of fees for private healthcare facilities, and in accordance with Resolution No. (48) of 2020 on the control of the quality of medical devices and products, the National Health Regulatory Authority (NHRA) hereby informs all relevant stakeholders of the updated fee schedule applicable to medical devices and supplies services.

The fee schedule applies to licensing, registration, importation, traceability and other related services for medical devices and supplies, as set out below.

The applicable fees shall be settled in accordance with the procedures and payment mechanisms specified by NHRA, including through the relevant NHRA systems and platforms where applicable.

All stakeholders are requested to review the fee schedule and ensure compliance with the applicable payment requirements when submitting requests to NHRA.

Thank you for your cooperation.



Dr. Ahmed Mohamed Al Ansari

Chief Executive Officer



Annex 1: Medical Devices and Supplies Fee Schedule

Table 1: Medical Devices Authorized Representative Licensing Services

Service Type	Item	Type	Fees	Norm
Authorized Representative License Application fee			50	Each new order
Authorized Representative licensing	Class A	First license	1,000	First license
		License renewal	700	Annual
	Class B	First license	700	First license
		License renewal	500	Annual
	Class C	First license	500	First license
		License renewal	300	Annual
	Class D	First license	300	First license
		License renewal	200	Annual
Medical devices Warehouse licensing			1,000	First license
			800	Annual
Medical devices manufacturer licensing			5,000	First license
			3,000	Annual
Update medical device facilities data			50	Each request
Re-inspection of medical device facilities			50	Each request

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Table 2: Medical Device Registration Services

All medical device registrations have a 3-year validity period.

Service Type	Item	Type	Fees (Every 3 years)	Norm
Application to submit a medical device registration			5	Each application
Medical Device & Supplies Traceability fee			5	Each individual device
Medical device registration	Low risk	Registration/Renewal	150	Triannual per license
	Medium risk	Registration/Renewal	300	Triannual per license
	High risk	Registration/Renewal	1,000	Triannual per license
Update medical device registration data			20	Each device

Table 3: Medical Devices Importation Services

Service Type	Item	Type	Fees	Norm
Request approval to import a single device	Medical device	Registered	0.5	Each type
		Unregistered	2	Each type
	IVD/combined medical device	Registered	1	Each type
		Unregistered	3	Each type
Verify the validity of the quality certificate			5	Each type
Product classification request		regular	20	Each type
		Urgent	40	Each type
Demurrage for the device till approval			100	Per day per shipment
Issuing a product classification certificate			15	Each certificate
Sample examination			5	Each type
Inspecting the shipment at the port			10	Each type

Table 4: Other Service

Service Type	Item	Fees	Norm
Permit to use	Submit a request for approval to use the device	0.5	Each device
	Evaluate the viability of the product	50	Each device
Issuing certificates for medical manufacturer and devices, free sale and other paper documents		10	Each request
Product disposal supervising		25	Each request

The above fee schedule shall be applied from 1st May 2026, unless otherwise instructed by NHRA.

All fees are stated in Bahraini Dinar.

For inquiries or further clarification, stakeholders are requested to contact:

md-support@nhra.bh

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