

Certra

Renewal software for Saudi medical-device
Authorized Representatives.

September 2026

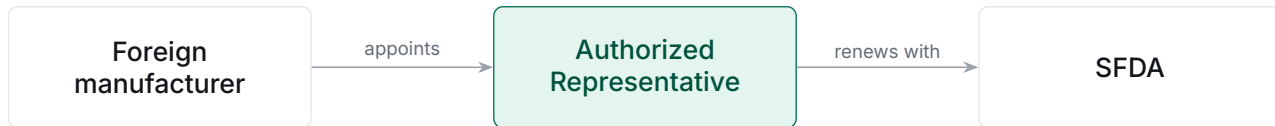
PROBLEM

Miss a renewal and the device leaves the market.

1,608

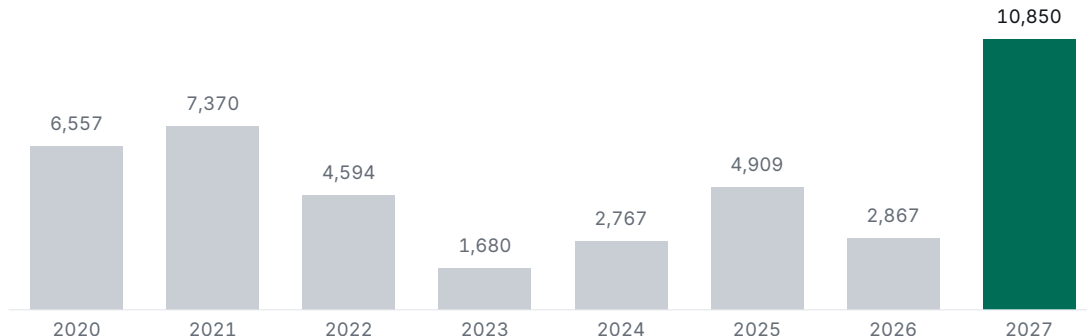
Authorized Representatives renew these certificates, mostly from spreadsheets.

The Authorized Representative carries the risk.



10,850 certificates expire in 2027.

The largest single expiry year on the register, and 13,717 across 2026 and 2027.



Device listings by MDMA expiry year. SFDA public device register, extract of 5 June 2026 (56,737 listings), analysed by MedDeviceGuide.

MARKET

A small, named market.

1,608

Authorized Representatives on
the register

428

hold 21–500 listings: our core
market

18

hold more than 500: enterprise

The core 428 are worth \$0.86–1.31M a year at list price.

SFDA register extract of 5 June 2026, analysed by MedDeviceGuide. Revenue: 211 ARs with 21–50 listings on Starter, 104 with 51–100 on Professional, and 113 with 101–500 on Professional or Portfolio. Listings include expired certificates, so this leans high.

Type a licence code. The portfolio is already there.

Portfolio

71 devices seeded from the SFDA register

Export

Search device, model or certificate number

12 months All manufacturers All classes All statuses Not started Clear

☐	Device	Manufacturer	Class	Certificate	Expiry ↑	Remaining	Renewal
☐	Anaesthesia Workstation XT-1004	PAUL HARTMANN	A	MDMA-2023-050857	13 Oct 2026	18d	Documents requested 6 blocking
☐	Digital Blood Pressure Monitor GX-2324	Intuitive Surgical, Inc.	A	MDMA-2023-051098	26 Oct 2026	31d	Documents requested 6 blocking
☐	Automated External Defibrillator PRO-6055	Boston Scientific Corporation	D	MDMA-2023-051028	15 Nov 2026	51d	Documents requested 10 blocking
☐	Digital Blood Pressure Monitor SR-7836	Vygon SA	B	MDMA-2023-051126	03 Dec 2026	69d	Documents requested 7 blocking

Straight from the register, with no data entry. Shown with demo data.

Bring the spreadsheet you already keep.

2-MESSY-DEVICE-REGISTER.XLSX · بيانات

× Start over

A list of devices 95%

COLUMNS WE RECOGNISED

Product → Device Model No → Model الشركة المصنعة → Manufacturer Class → Class Reg. No → Certificate تاريخ الانتهاء → Expiry

Dates read day-first, which the values in the column confirm.

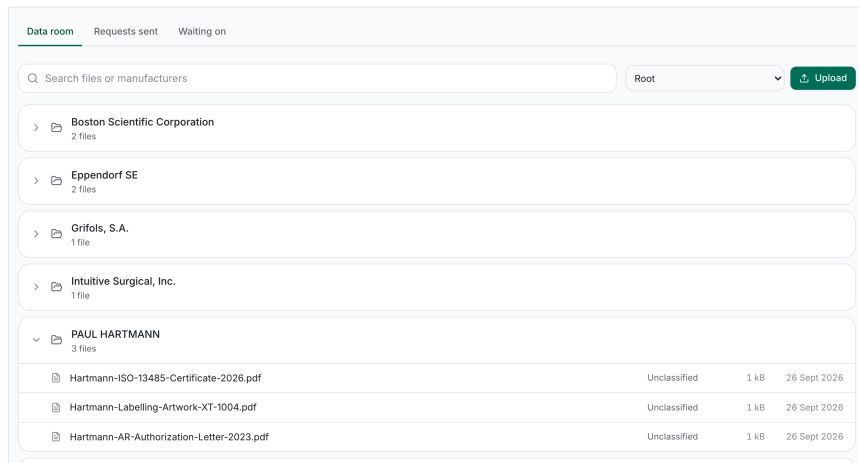
COLUMNS WE COULD NOT PLACE

Internal Owner · Chased?

These are kept. Anything we cannot map is saved on the device as a note, so nothing in your sheet is lost.

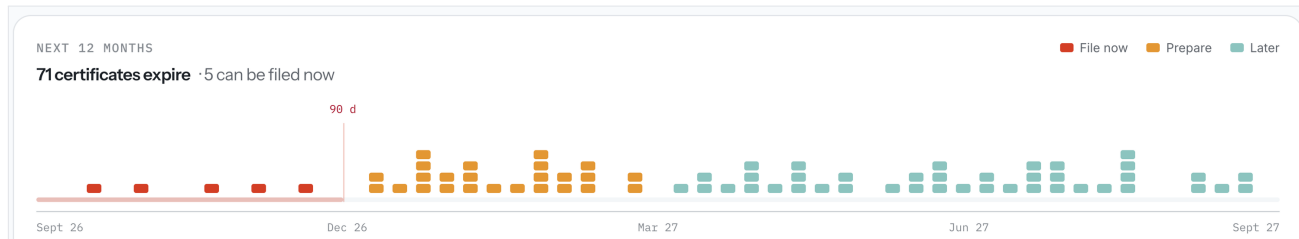
Reads your columns, even in Arabic. Flags what it cannot read. Writes nothing until you approve.

Every file, filed by manufacturer.



Uploads from manufacturers file themselves into the right folder.

See the renewal wall before you hit it.



Every certificate expiring in the next twelve months, and which can be filed today.

Request by link. Know when it lands.

THE MANUFACTURER UPLOADS. NO ACCOUNT.

4 still needed

1 received

Needed for the October renewal filing. Thank you.

For these devices

Anaesthesia Workstation MDMA-2023-050857

🕒 Declaration of Conformity

🔄 Replace

YOU ARE NOTIFIED. THE CHECKLIST TICKS ITSELF.

NOTIFICATIONS

✓ Mark all read

• PAUL HARTMANN uploaded a document

8 seconds ago

• PAUL HARTMANN uploaded a document

2 minutes ago

• Reminder: Dialysis Machine expires in 12 months

10 hours ago

See all notifications

Work each renewal as a team.

OWNERS AND INTERNAL COMMENTS

 **Internal comments**

Never shown to manufacturers.

Noura, Hartmann still owes the ISO 13485 certificate and the Declaration of Conformity. Can you get them in before we file?

26 Sept 2026

On it. Sending them the upload link now, and I will check the certificate scope against our device list when it lands.

26 Sept 2026

Post

PERMISSIONS SET PER PERSON

Start from what a Specialist gets, then change it for this person. Their role stays the same.

☒ Confirm extracted values	☒ Import devices and manufacturers Added for this person
☐ Manage plan and billing	☐ Export or delete the workspace
☐ Manage the team	☐ Change workspace settings
☐ Request documents from manufacturers	☐ Resolve findings Removed for this person
☒ Upload documents	

[↶ Back to role default](#) [Cancel](#) **Save permissions**

Catch the mismatch before the regulator does.

Findings

123 blockers, 0 warnings

Every finding cites its sources. Click one to open the document at that page.

[All severities](#) [Run a full scan](#)

▼ **Digital Blood Pressure Monitor**
6 findings

6 blocking

[Open this renewal →](#)

Blocker

ISO 13485 quality management system certificate has not been received. It is required for a Class A device and the file cannot be submitted without it.

Seen 3 times since 24 Sept 2026

Your records · Checklist: ISO 13485 quality management system certificate

ISO 13485 quality management system certificate

SFDA register · SFDA register record

A

Resolve

The AI flags and cites. A person decides.

EC Certificate — Quality Management System SPECIMEN — generated sample for product demonstration. Not a real certificate. ISO 13485:2016 — Medical devices — Quality management systems Certificate number: MD 704221 / QMS 13485 Issued by: TÜV SÜD Product Service GmbH Ridlerstrasse 65, 80339 Munich, Germany Certified organisation: Fresenius Kabi Deutschland GmbH Site 1: Else-Kroener-Strasse 1, 61352 Bad Homburg, Germany Site 2: Borkenberg 14, 61440 Oberursel, Germany Scope: Design, development, manufacture and distribution of infusion pumps, infusion systems and associated disposable administration sets. Date of issue: 2023-11-14 Valid until: 2026-11-13 This certificate remains the property of the certification body.	CERTIFICATE NUMBER MD 704221 / QMS 13485 Confidence 96% · Page 1 Correct EXPIRY DATE 2026-11-13 Confidence 98% · Page 1 Correct ISSUE DATE 2023-11-14 Confidence 95% · Page 1
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Every extracted value points to the words it came from.

WHERE WE ARE

The product is built. The business has not started.

DONE

Full product, end to end
Arabic and English
Billing integrated

NOT YET

Customers
Revenue
Live register access

Priced by portfolio size.

\$99

Starter, up to 50 devices

\$299

Professional, up to 200

\$699

Portfolio, up to 1,000

Per month. Annual billing is ten months' price. Enterprise is priced separately.

Next 90 days: customers, not code.

1

Deploy in the Kingdom

2

Connect the live
register

3

Talk to ten ARs

4

First paid pilot

Certra

We verify renewals. We never write the technical file.