DEADLINE · NOVEMBER 28, 2026

EUDAMED readiness: can you tick all of these?

If you manufacture, import, or sell medical devices in the EU, use of the EUDAMED UDI/Device registration module becomes mandatory on November 28, 2026.

☐ Actor registration & SRN

Your organisation is registered in EUDAMED and holds a Single Registration Number.

☐ Basic UDI-DI assigned

Every device family has a Basic UDI-DI linked to its technical documentation and certificates.

☐ UDI from an issuing entity

Your UDI-DIs come from an EU-designated issuing entity (GS1, HIBCC, ICCBBA or IFA).

☐ Complete device data

All required EUDAMED UDI/Device attributes are collected, validated and ready to submit.

☐ Labels match the data

Printed labels and EUDAMED records agree field by field, and your printing process is validated.

☐ Legacy & on-market devices planned

You know which devices already on the EU market need registering and by when.

☐ Upload route chosen

Manual entry, XML bulk upload, or machine-to-machine, matched to your portfolio size.

☐ Owner & maintenance plan

Someone owns ongoing updates when products, labels or certificates change.

MISSING ONE?

Book a free 15–30 minute video consultation call. Our team will identify the gaps.
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Take the interactive version at globaludi.com/readiness.html. General guidance only; timing for devices already on the market depends on your device and certificate status.