EUDAMED - European Database on Medical Devices

EUDAMED Time line

The European Commission targets

Q4 2022	Q1-Q2 2023	Q2 2023	Q2 2023	Q4 2023	Q2 2025
End of the EUDAMED MVP[1] development for all six modules	Independent **Audit**	Audit results presented to the Medical Devices Coordination Group (MDCG)	EUDAMED has achieved full functionality following the outcome of the Audit **Publication of a Commission notice** in the *Official Journal of the European Union* **(OJEU)** The full EUDAMED system is ready Only the first 3 modules, with features available on voluntary basis, are in production	End of 6 months transitional period after publication of the notice in the OJEU Fully functional EUDAMED (all 6 modules) goes live The use of **EUDAMED** becomes **mandatory** as regards obligations and requirements related to **Actors, Vigilance, Clinical Investigation & Performance Studies and Market Surveillance modules**	End of 24 months transitional period after publication of the notice in the OJEU The use of **EUDAMED** becomes **mandatory** as regards obligations and requirements related to **UDI/Device and NB & Certificate modules**

[1] EUDAMED Minimum Viable Product (MVP) means that the system developed implements at least the minimum Medical Devices Regulations requirements and allows competent authorities and all stakeholders to comply with their legal obligations.