


EUDAMED - European Database on Medical Devices

EUDAMED Time line

The European Commission planning – June 2022



Q4 2023	Q1-Q2 2024	Q2 2024	Q2 2024	Q4 2024	Q2 2026
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 <i>Official Journal of the European Union (OJEU)</i> The full EUDAMED system (all 6 modules) is released.	End of 6 months transitional period after publication of the notice in the OJEU 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.