

UDI World Overview

Authority		Database	Issuing Agencies	Current state	Deadline
	FDA (Food and Drug Administration)	GUIDID (Global Unique Device Identification Database)	GS1, HIBBC, ICCBBA	Last category submission by September 24, 2022: Class I and devices that have not been classified into class I, class II or class III.	2014 - 2022
	European Commission	EUDAMED (European database on medical devices)	GS1, HIBBC, ICCBBA, IFA	Database registration opened (SRN). Playground available. MDR fully applies from May 26, 2021.	2021 - 2027
	Swissmedic	eGov Portal (for the regulating authority) BASEC (Business Administration System for Ethics Committees, for the ethics committee)	GS1, HIBCC, ICCBBA, IFA	The Mutual Recognition Agreement (MRA) between the EU and Switzerland may be updated. If not, an amendment to the Medical Devices Ordinance (Eventual-MepV) will be put into force by the on May 26, 2021. For the time being, registration in EUDAMED is voluntary.	
	NMPA (National Medical Products Administration)	MDUID (Medical Device Unique Identification Database)	GS1 China	Database registration opened. Class III products data collection/transfer.	2021 - 2026
	SFDA (Saudi Food & Drug Administration)	Saudi-DI	GS1, HIBCC, ICCBBA	Database registration opened. Class D products data submission & marking.	2021 - 2024
	MFDS (Ministry of Food & drug Safety)	IMDIS (Integrated Medical Device Information System)	GS1	Database Registration opened. Class II products data collection/transfer.	2019 - 2022
	TGA (Therapeutic Goods Administration)	AusUDID	GS1, HIBCC, ICCBBA	Implementation in progress.	2021-2027
	Health Canada	MDALL (Medical Devices Active Licence Listing)	GS1, HIBCC, ICCBBA	Implementation in progress. Manufacturer are required to obtain a Medical Device Establishment License (MDEL).	2023-2030
	MHRA (Medicines & Healthcare products Regulatory Agency)	MHRA Database		CE marking and certificates are accepted for the UK market until June 30, 2023. More info on our website: UDI compliance during Brexit	2021-2022