



# Transition Timelines from the Directives to the medical devices Regulation



**Until 25 May 2021**  
All certificates issued under the medical devices Directives (AIMDD/MDD) are valid until their date of expiry

**26 May 2021 – 25 May 2024**  
Certificates issued under the AIMDD/MDD before the MDR fully applies may remain valid until 25 May 2024 under certain conditions \*

**26 May 2024 – 27 May 2025**  
MDD devices already placed on the market may continue to be made available



**From May 26 2017**  
Devices that conform with the medical devices Regulation (MDR) may be placed on the market

**From May 2024**  
All devices placed on the market must be in conformity with the MDR

**26 MAY 2017** MDR enters into force (and is partially applicable)

**26 MAY 2021** MDR applies



ACRONYMS  
AIMDD: Directive 90/385/EEC MDD: Directive 93/42/EEC MDR: Regulation (EU) 2017/745

\* In addition, MDD Class I devices that would require the involvement of a Notified Body under the MDR may continue to be placed on the market until 25 May 2024 under certain conditions.

Health

Funded under the Third Health Programme 2014 -2020



# Transition Timelines from the Directive to the *in vitro* diagnostic medical devices Regulation



**Until 25 May 2022**  
All certificates issued under the *in vitro* diagnostic medical devices Directive (IVDD) are valid until their date of expiry

**26 May 2022 – 25 May 2024**  
certificates issued under the IVDD from 25 May 2017 expire latest by 27 May 2024

**26 May 2024 – 27 May 2025**  
IVDD devices already placed on the market may continue to be made available



**From 26 MAY 2017**  
Devices that conform to the *in vitro* diagnostic medical devices Regulation (IVDR) may be placed on the market

**From 26 MAY 2024**  
All devices placed on the market must be in conformity with the IVDR

**26 MAY 2017** The IVDR enters into force

**26 MAY 2022** The IVDR applies



ACRONYMS  
IVDD: Directive 98/79/EC IVDR: Regulation (EU) 2017/746

Health

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