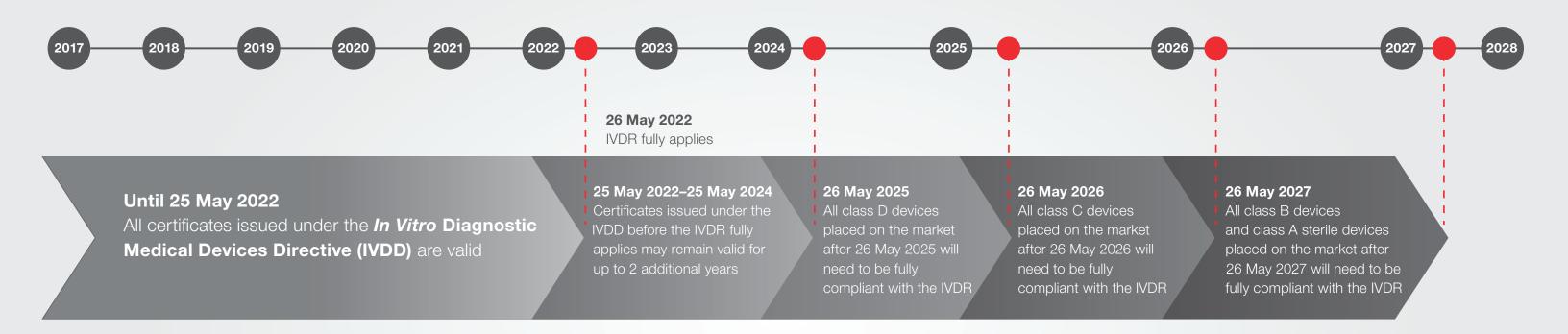
IVDR transition timelines



From 26 May 2022

No change: Any class A nonsterile products placed on the market after 25 May 2022 will need to be fully compliant with the IVDR

From 26 May 2017

Devices that conform with 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



Learn more about how Thermo Fisher Scientific can support you through the IVDR transition »