

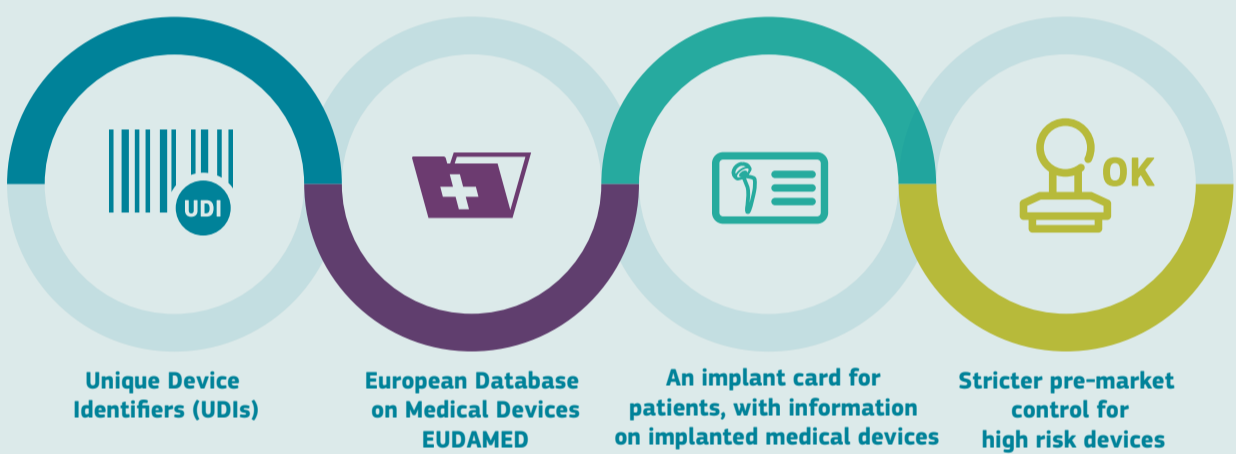
Medical Devices Regulation (MDR) and *In Vitro* Diagnostic Medical Devices Regulation (IVDR)

The European Commission has adopted 2 new Regulations – the Medical Devices Regulation (MDR) and the *In Vitro* Diagnostic Medical Devices Regulation (IVDR) - to bring EU legislation up to date with medical advances and to ensure better protection of public health and patient safety.

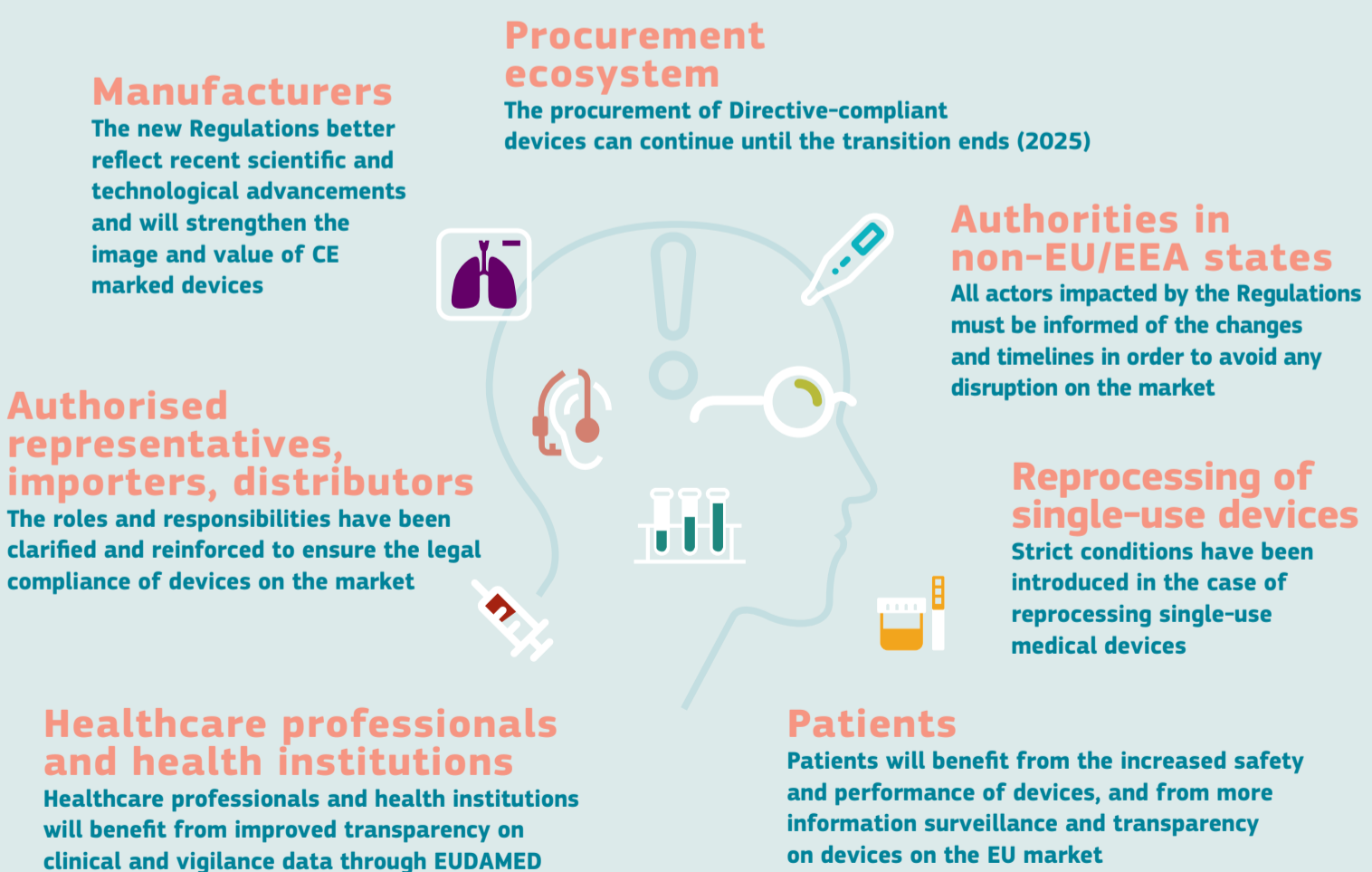
THE NEW REGULATIONS



SOME OF THE NEW FEATURES:



SOME THINGS TO KEEP IN MIND...



For a complete overview of the impact of the new Regulations and the roles and responsibilities of all stakeholders, check the Medical Devices section on the DG GROW website: https://ec.europa.eu/growth/sectors/medical-devices_en

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