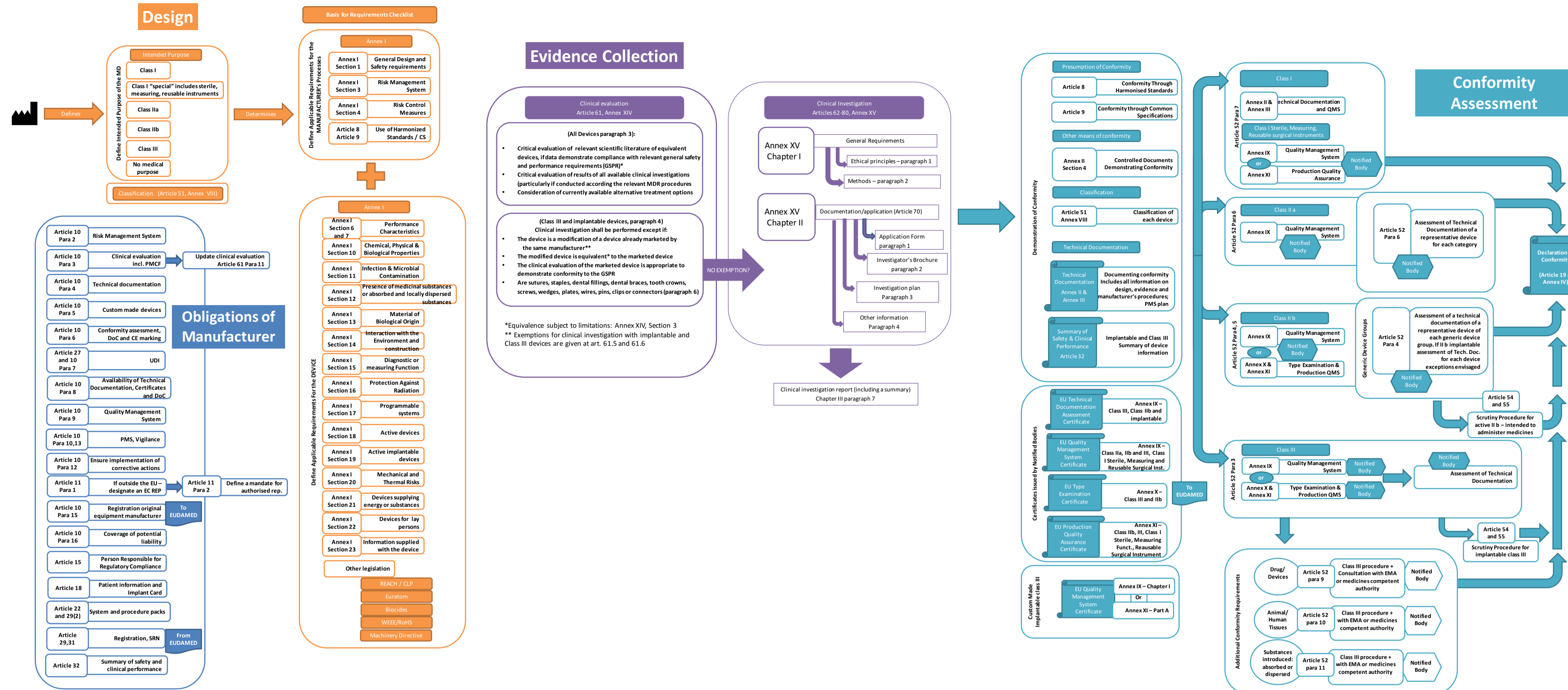


Overview of requirements under
the Medical Devices Regulation
Regulation 2017/745/EU on Medical Devices
December 2017

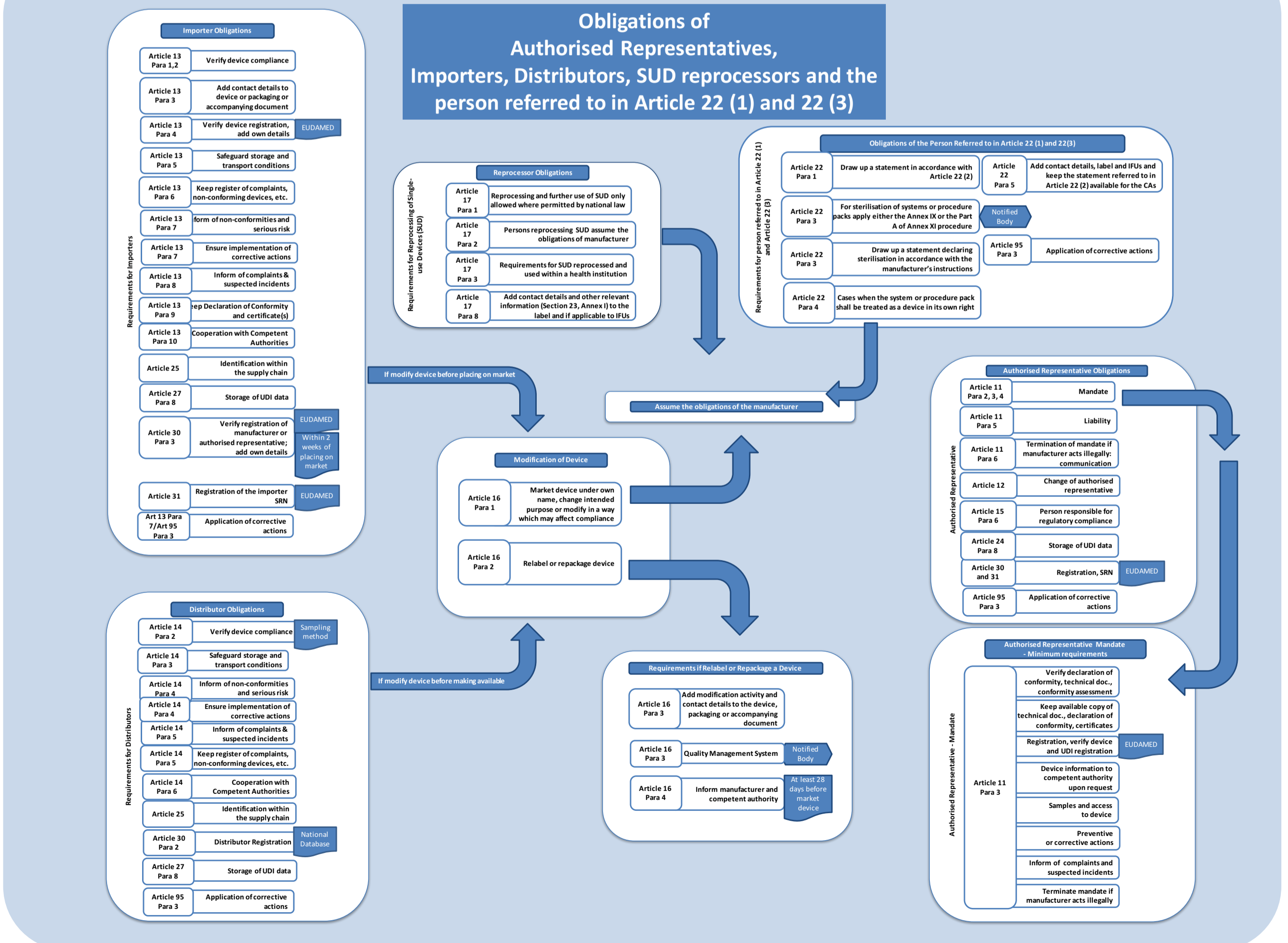
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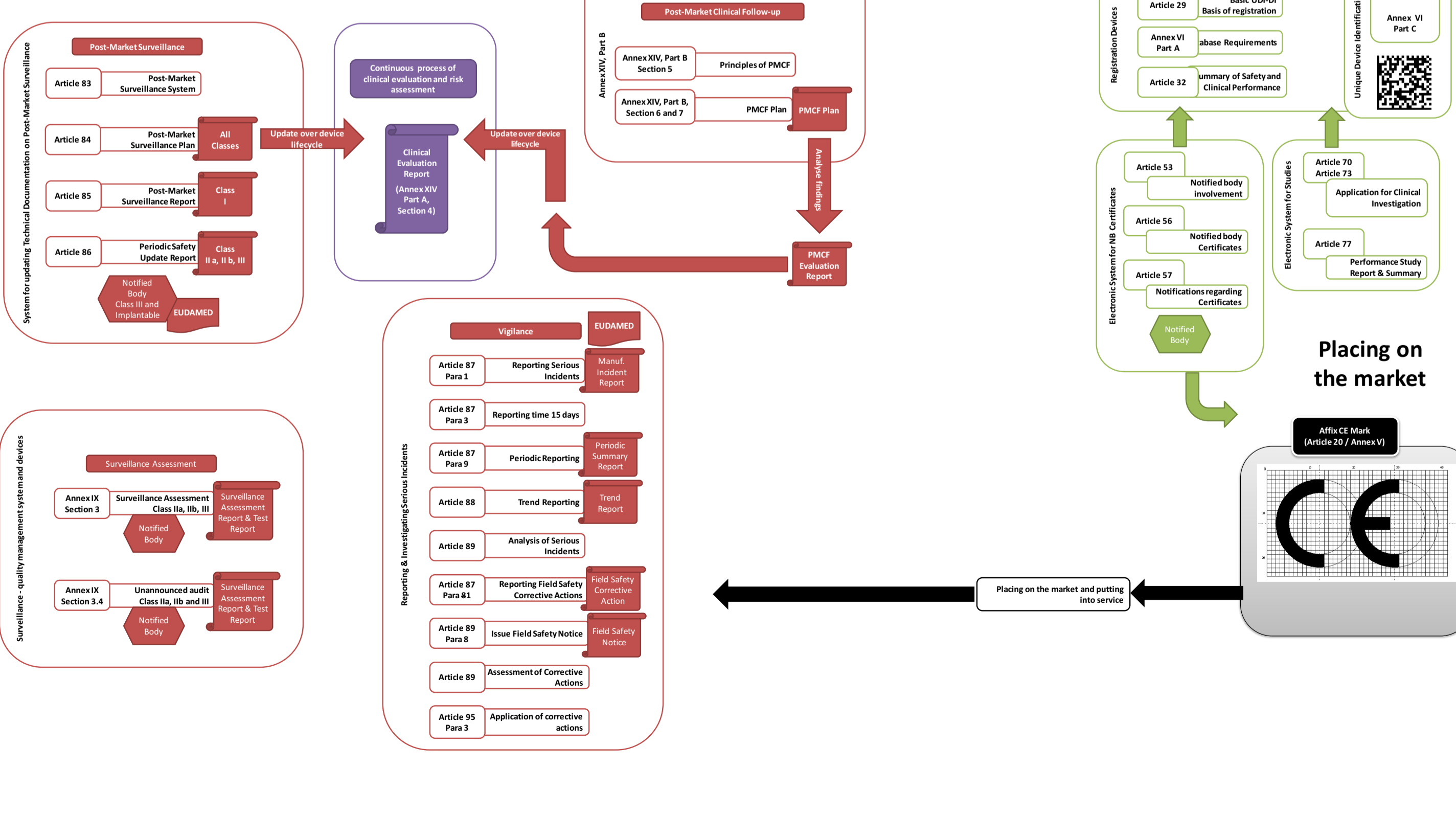
For more information please contact the regulations & industrial policy department: regulatory@medtecheurope.org



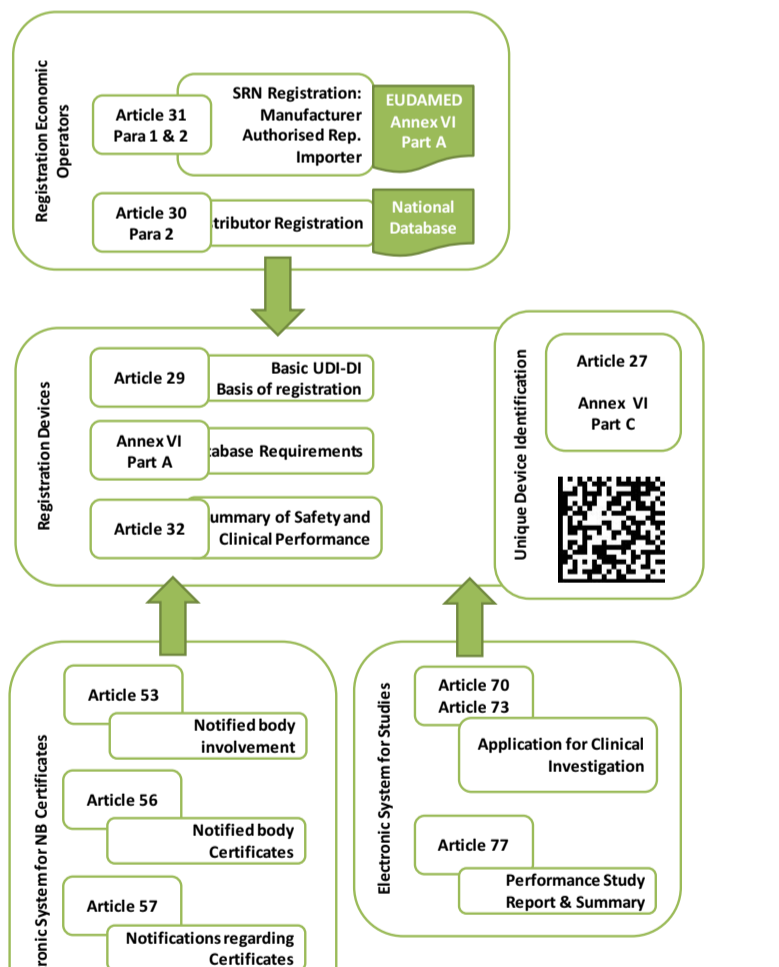
Obligations of
Authorised Representatives,
Importers, Distributors, SUD reprocessors and the
person referred to in Article 22 (1) and 22 (3)



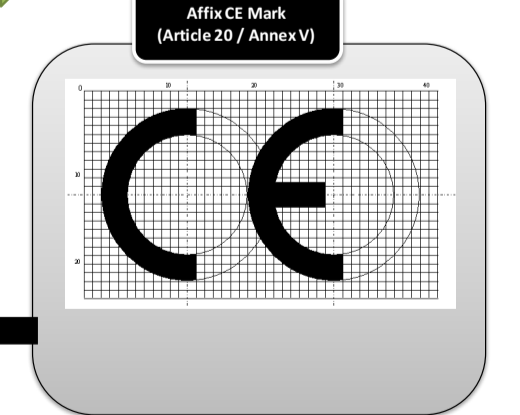
Post-Market Surveillance
and Vigilance



Registration



Placing on
the market



Overview of Regulation 2017/745/EU on Medical Devices