MDR Regulation 2017/475

MEDICAL DEVICE COORDINATION GROUP

Link to the European comission

1. Notified bodies oversight (NBO)
2. Standards
3. Clinical investigation and evaluation (CIE)
4. Post-market surveillance and vigilance (PMSV)
5. Market Surveillance (MS)
6. Borderline and classification (B&C)
7. New technologies
8. EUDAMED
9. Unique device identification (UDI)
10. International matters
11. In vitro diagnostic medical devices (IVD)
12. Nomenclature
13. “Annex XVI” products