**Key documents**

**Recommendations**

- Commission Recommendation 2013/172/EU of 5 April 2013 on a common framework for a unique device identification system for medical devices in the Union
- Commission Recommendation 2013/473/EU of 24 September 2013 on the audits and assessments performed by notified bodies in the field of medical devices

**Classification**

- Reclassification of hip, knee and shoulder joint replacements:
- Reclassification of breast implants:

**Other amending and implementing legislation**

- Designation and the supervision of notified bodies:
- Common technical specification on IVDs:
  - Commission Implementing Decision 2019/1244/EU - OJ L193/1 of 19 July 2019
- Qualification of products depending on proanthocyanidins present in cranberry
- Eudamed2 - European databank on medical devices:
- Medical devices manufactured utilising tissues of animal origin:
- Electronic instructions for use of medical devices:
- Variant Creutzfeld-Jakob disease assays: