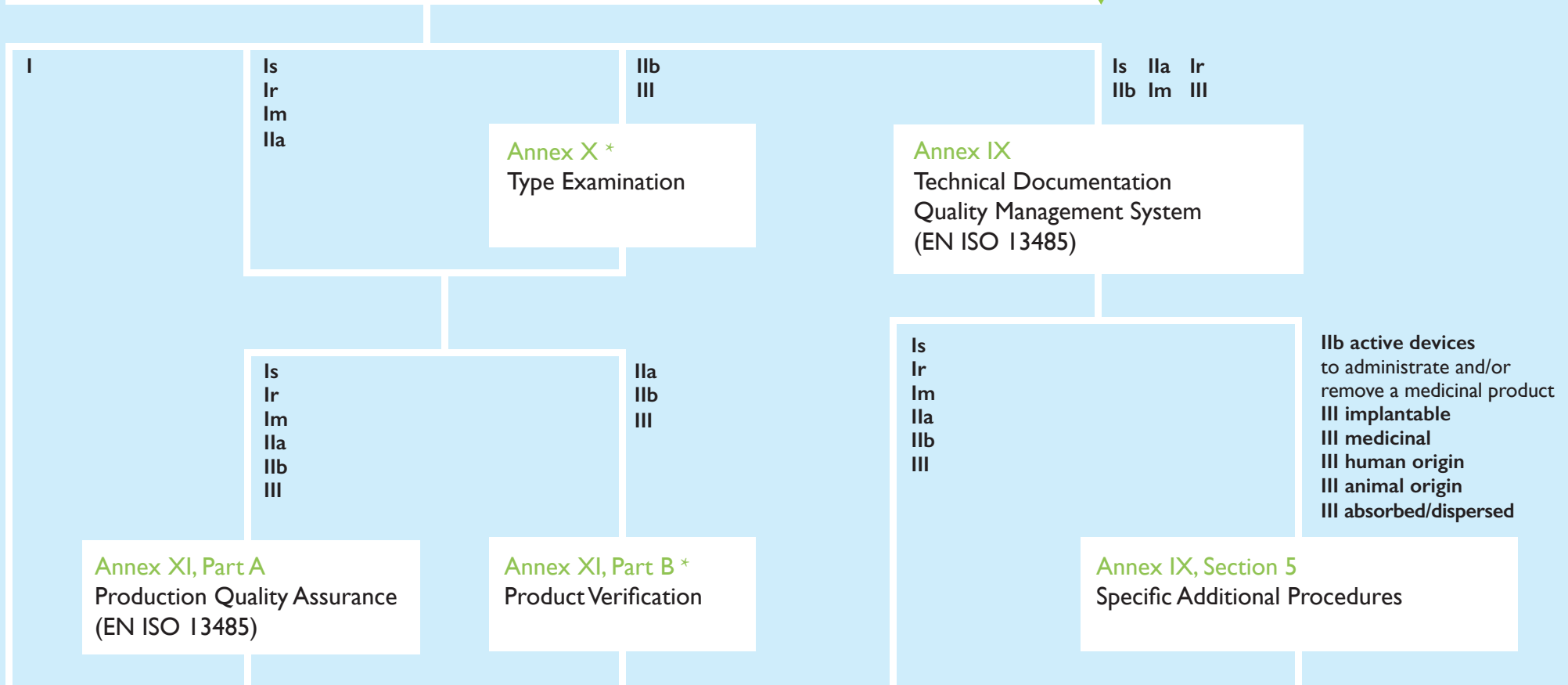


# MDR Conformity Assessment Procedure **OVERVIEW**

- Annex I General Safety and Performance Requirements
- Annex II Technical Documentation
- Annex III Technical Documentation on Post Market Surveillance
- Annex IV EU Declaration of Conformity
- Annex VI UDI – Unique Device Identification
- Annex VIII Classification Rules



\*This service is not offered by MEDCERT, and the marking "CE 0482" may not be affixed under this annex.