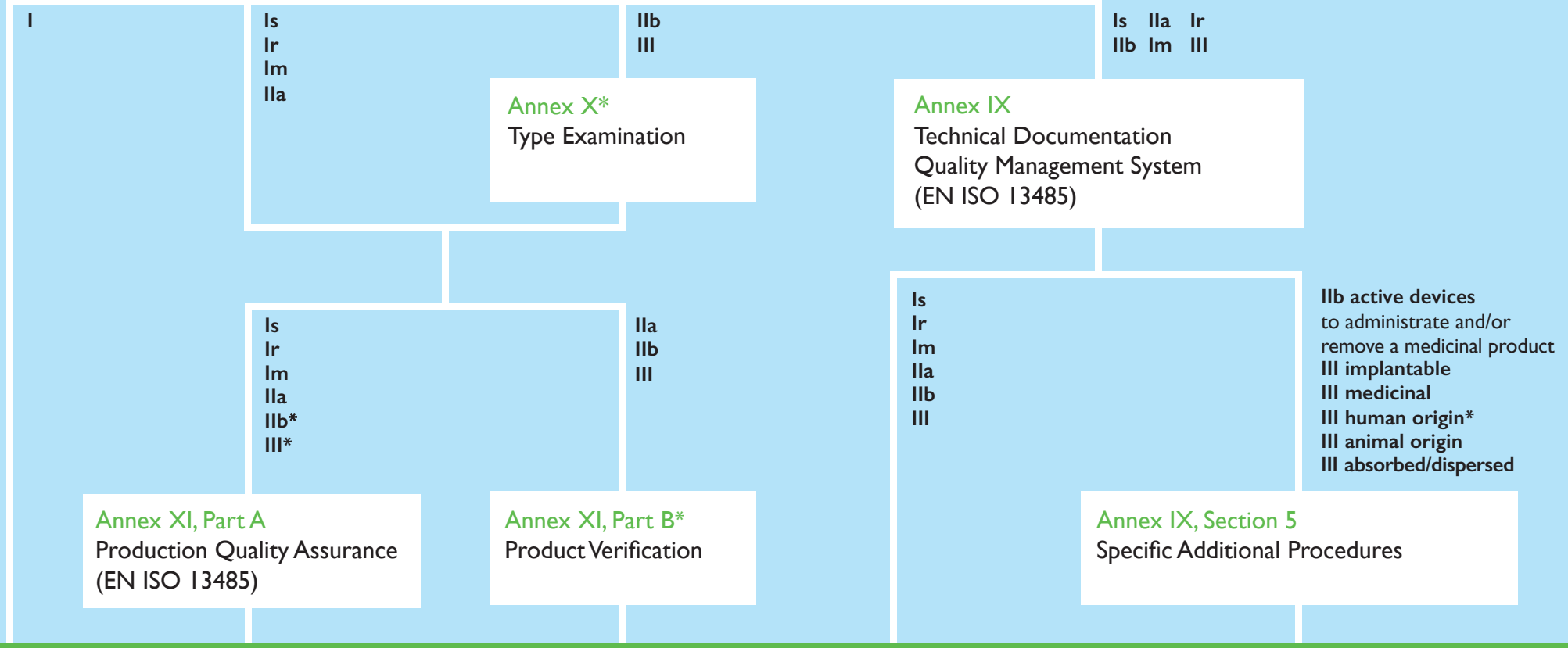
MDR Conformity Assessment Procedure OVERVIEW

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



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CE 0482

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