MDR Database - MDRQ #1673

MDRQ # 1545 (Released): CHAPTER X FINAL PROVISIONS

MDRQ # 1610 (Released): Article 123 Entry into force and date of application

ANNEXES

11/04/2017 02:17 PM - Anonymous

Status:	Released	
Priority:	Low	

Description

ANNEXES

Subissues:

MDRQ # 1674: I General safety and performance requirements	**Released**
MDRQ # 1675: II Technical documentation	**Released**
MDRQ # 1676: III Technical documentation on post-market surveillance	**Released**
MDRQ # 1677: IV EU declaration of conformity	**Released**
MDRQ # 1678: V CE marking of conformity	**Released**
MDRQ # 1679: VI Information to be submitted upon the registration of devices and econom...	**Released**
MDRQ # 1680: VII Requirements to be met by notified bodies	**Released**
MDRQ # 1681: VIII Classification rules	**Released**
MDRQ # 1682: IX Conformity assessment based on a quality management system and assessme...	**Released**
MDRQ # 1683: X Conformity assessment based on type examination	**Released**
MDRQ # 1684: XI Conformity assessment based on product conformity verification	**Released**
MDRQ # 1685: XII Certificates issued by a notified body	**Released**
MDRQ # 1686: XIII Procedure for custom-made devices	**Released**
MDRQ # 1687: XIV Clinical evaluation and post-market clinical follow-up	**Released**
MDRQ # 1688: XV Clinical investigations	**Released**
MDRQ # 1689: XVI List of groups of products without an intended medical purpose referre...	**Released**