Commission Regulation (EU) No 712/2012 of 3 August 2012 amending Regulation (EC) No 1234/2008 concerning the examination of variations to the terms of marketing authorisations for medicinal products for human use and veterinary medicinal products (Text with EEA relevance)

ANNEX

ANNEX VI List of Member States referred in Article 24a...

(**1**) OJ L 311, 28.11.2001, p. 1.

(**2**) OJ L 311, 28.11.2001, p. 67.

(**3**) OJ L 136, 30.4.2004, p. 1.

(**4**) OJ L 168, 30.6.2009, p. 33.

(**5**) OJ L 334, 12.12.2008, p. 7.

(**6**) OJ L 348, 31.12.2010, p. 1.

(**7**) OJ L 348, 31.12.2010, p. 74.

Changes to legislation:
There are outstanding changes not yet made to Commission Regulation (EU) No 712/2012. Any changes that have already been made to the legislation appear in the content and are referenced with annotations.
View outstanding changes

Changes and effects yet to be applied to :

– Regulation revoked in part by S.I. 2019/775 Sch. 9 para. 1(u)