Regulation (EU) No 1235/2010 of the European Parliament and of the Council of 15 December 2010 amending, as regards pharmacovigilance of medicinal products for human use, Regulation (EC) No 726/2004 laying down Community procedures for the authorisation and supervision of medicinal products for human and veterinary use and establishing a European Medicines Agency, and Regulation (EC) No 1394/2007 on advanced therapy medicinal products (Text with EEA relevance)

(**1**) OJ C 306, 16.12.2009, p. 22.

(**2**) OJ C 79, 27.3.2010, p. 50.

(**3**) OJ C 229, 23.9.2009, p. 19.

(**4**) Position of the European Parliament of 22 September 2010 (not yet published in the Official Journal) and Council Decision of 29 November 2010.

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

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

(**7**) OJ L 281, 23.11.1995, p. 31.

(**8**) OJ L 8, 12.1.2001, p. 1.

(**9**) See page 74 of this Official Journal.

(**10**) OJ L 184, 17.7.1999, p. 23.

(**11**) OJ L 218, 13.8.2008, p. 30.

(**12**) OJ L 324, 10.12.2007, p. 121.

Changes to legislation:
There are outstanding changes not yet made to Regulation (EU) No 1235/2010 of the European Parliament and of the Council. 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(r)