

Directive 2011/62/EU of the European Parliament and of the Council of 8 June 2011 amending Directive 2001/83/EC on the Community code relating to medicinal products for human use, as regards the prevention of the entry into the legal supply chain of falsified medicinal products (Text with EEA relevance)

- Article 1 Directive 2001/83/EC is hereby amended as follows: Article 1 is...
- Article 2 (1) Member States shall bring into force the laws, regulations...
- Article 3 At the latest 5 years after the date of application...
- Article 4 In order to adopt the delegated acts referred to in...
- Article 5 This Directive shall enter into force on the 20th day...
- Article 6 This Directive is addressed to the Member States.
- Signature

- (1) OJ C 317, 23.12.2009, p. 62.
- (2) OJ C 79, 27.3.2010, p. 50.
- (3) Position of the European Parliament of 16 February 2011 (not yet published in the Official Journal) and decision of the Council of 27 May 2011.
- (4) OJ L 311, 28.11.2001, p. 67.
- (5) OJ L 281, 23.11.1995, p. 31.
- (6) OJ L 210, 7.8.1985, p. 29.
- (7) Judgment of the Court of 19 May 2009 in Joined Cases C-171/07 and C-172/07 *Apothekerkammer des Saarlandes and Others v Saarland* ECR [2009] I-4171, paragraphs 19 and 31.
- (8) Judgment of the Court of 19 May 2009 in Joined Cases C-171/07 and C-172/07 *Apothekerkammer des Saarlandes and Others v Saarland* ECR [2009] I-4171, paragraphs 34 and 35.
- (9) OJ L 55, 28.2.2011, p. 13.
- (10) OJ C 321, 31.12.2003, p. 1.
- (11) OJ L 348, 31.12.2010, p. 74.