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.