Directive 2011/24/EU of the European Parliament and of the Council of 9 March 2011 on the application of patients' rights in cross-border healthcare

CHAPTER I

GENERAL PROVISIONS

Article 2

Relationship with other Union provisions

This Directive shall apply without prejudice to:

(a) Council Directive 89/105/EEC of 21 December 1988 relating to the transparency of measures regulating the prices of medicinal products for human use and their inclusion in the scope of national health insurance systems[(1)];

(b) Council Directive 90/385/EEC of 20 June 1990 on the approximation of the laws of the Member States relating to active implantable medical devices[(2)], Council Directive 93/42/EEC of 14 June 1993 concerning medical devices[(3)] and Directive 98/79/EC of the European Parliament and of the Council of 27 October 1998 on in vitro diagnostic medical devices[(4)];

(c) Directive 95/46/EC and Directive 2002/58/EC of the European Parliament and of the Council of 12 July 2002 concerning the processing of personal data and the protection of privacy in the electronic communications sector[(5)];

(d) Directive 96/71/EC of the European Parliament and of the Council of 16 December 1996 concerning the posting of workers in the framework of the provision of services[(6)];

(e) Directive 2000/31/EC;

(f) Council Directive 2000/43/EC of 29 June 2000 implementing the principle of equal treatment between persons irrespective of racial or ethnic origin[(7)];

(g) Directive 2001/20/EC of the European Parliament and of the Council of 4 April 2001 on the approximation of the laws, regulations and administrative provisions of the Member States relating to the implementation of good clinical practice in the conduct of clinical trials on medicinal products for human use[(8)];

(h) Directive 2001/83/EC of the European Parliament and of the Council of 6 November 2001 on the Community code relating to medicinal products for human use[(9)];

(i) Directive 2002/98/EC of the European Parliament and of the Council of 27 January 2003 setting standards of quality and safety for the collection, testing, processing, storage and distribution of human blood and blood components[(10)];

(j) Regulation (EC) No 859/2003;

(k) Directive 2004/23/EC of the European Parliament and of the Council of 31 March 2004 on setting standards of quality and safety for the donation, procurement, testing, processing, preservation, storage and distribution of human tissues and cells[(11)];

(l) Regulation (EC) No 726/2004 of the European Parliament and of the Council of 31 March 2004 laying down Community procedures for the authorisation and supervision of medicinal products for human and veterinary use and establishing a European Medicines Agency[(12)];

(m) Regulation (EC) No 883/2004 and Regulation (EC) No 987/2009 of the European Parliament and of the Council of 16 September 2009 laying down the procedure for implementing Regulation (EC) No 883/2004 on the coordination of social security systems[(13)];

(n) Directive 2005/36/EC;

(o) Regulation (EC) No 1082/2006 of the European Parliament and of the Council of 5 July 2006 on a European grouping of territorial cooperation (EGTC)[(14)];

(p) Regulation (EC) No 1338/2008 of the European Parliament and of the Council of 16 December 2008 on Community statistics on public health and health and safety at work[(15)];

(q) Regulation (EC) No 593/2008 of the European Parliament and of the Council of 17 June 2008 on the law applicable to contractual obligations (Rome I)[(16)], Regulation (EC) No 864/2007 of the European Parliament and of the Council of 11 July 2007 on the law applicable to non-contractual obligations (Rome II)[(17)] and other Union rules on private international law, in particular rules related to court jurisdiction and the applicable law;

(r) Directive 2010/53/EU of the European Parliament and of the Council of 7 July 2010 on standards of quality and safety of human organs intended for transplantation[(18)];

(s) Regulation (EU) No 1231/2010.

(**1**) OJ L 40, 11.2.1989, p. 8.

(**2**) OJ L 189, 20.7.1990, p. 17.

(**3**) OJ L 169, 12.7.1993, p. 1.

(**4**) OJ L 331, 7.12.1998, p. 1.

(**5**) OJ L 201, 31.7.2002, p. 37.

(**6**) OJ L 18, 21.1.1997, p. 1.

(**7**) OJ L 180, 19.7.2000, p. 22.

(**8**) OJ L 121, 1.5.2001, p. 34.

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

(**10**) OJ L 33, 8.2.2003, p. 30.

(**11**) OJ L 102, 7.4.2004, p. 48.

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

(**13**) OJ L 284, 30.10.2009, p. 1.

(**14**) OJ L 210, 31.7.2006, p. 19.

(**15**) OJ L 354, 31.12.2008, p. 70.

(**16**) OJ L 177, 4.7.2008, p. 6.

(**17**) OJ L 199, 31.7.2007, p. 40.

(**18**) OJ L 207, 6.8.2010, p. 14.