Directive 2014/30/EU of the European Parliament and of the Council of 26 February 2014 on the harmonisation of the laws of the Member States relating to electromagnetic compatibility (recast) (Text with EEA relevance)

CHAPTER 1

GENERAL PROVISIONS

Article 1 Subject matter
Article 2 Scope
Article 3 Definitions
Article 4 Making available on the market and/or putting into service
Article 5 Free movement of equipment
Article 6 Essential requirements

CHAPTER 2

OBLIGATIONS OF ECONOMIC OPERATORS

Article 7 Obligations of manufacturers
Article 8 Authorised representatives
Article 9 Obligations of importers
Article 10 Obligations of distributors
Article 11 Cases in which obligations of manufacturers apply to importers and distributors
Article 12 Identification of economic operators

CHAPTER 3

CONFORMITY OF EQUIPMENT

Article 13 Presumption of conformity of equipment
Article 14 Conformity assessment procedures for apparatus
Article 15 EU declaration of conformity
Article 16 General principles of the CE marking
Article 17 Rules and conditions for affixing the CE marking
Article 18 Information concerning the use of apparatus
Article 19 Fixed installations

CHAPTER 4

NOTIFICATION OF CONFORMITY ASSESSMENT BODIES

Article 20 Notification
Article 21 Notifying authorities
Article 22 Requirements relating to notifying authorities
Article 23 Information obligation on notifying authorities
Article 24 Requirements relating to notified bodies
Article 25 Presumption of conformity of notified bodies

Article 26 Subsidiaries of and subcontracting by notified bodies
Article 27 Application for notification
Article 28 Notification procedure
Article 29 Identification numbers and lists of notified bodies
Article 30 Changes to notifications
Article 31 Challenge of the competence of notified bodies
Article 32 Operational obligations of notified bodies
Article 33 Appeal against decisions of notified bodies
Article 34 Information obligation on notified bodies
Article 35 Exchange of experience
Article 36 Coordination of notified bodies

CHAPTER 5

UNION MARKET SURVEILLANCE AND CONTROL OF APPARATUS ENTERING THE UNION MARKET AND UNION SAFEGUARD PROCEDURE

Article 37 Union market surveillance and control of apparatus entering the Union market
Article 38 Procedure for dealing with apparatus presenting a risk at national level
Article 39 Union safeguard procedure
Article 40 Formal non-compliance

CHAPTER 6

COMMITTEE, TRANSITIONAL AND FINAL PROVISIONS

Article 41 Committee procedure
Article 42 Penalties
Article 43 Transitional provisions
Article 44 Transposition
Article 45 Repeal
Article 46 Entry into force and application
Article 47 Addressees

ANNEX I

ESSENTIAL REQUIREMENTS

1. General requirements

2. Specific requirements for fixed installations

ANNEX II

MODULE A: INTERNAL PRODUCTION CONTROL

1. Internal production control is the conformity assessment procedure whereby the...

2. Electromagnetic compatibility assessment

3. Technical documentation

4. Manufacturing

5. CE marking and EU declaration of conformity
   5.1. The manufacturer shall affix the CE marking to each individual...
   5.2. The manufacturer shall draw up a written EU declaration of...

6. Authorised representative

ANNEX III

PART A

Module B: EU-type examination
1. EU-type examination is the part of a conformity assessment procedure...
2. EU-type examination shall be carried out by assessment of the...
3. The manufacturer shall lodge an application for EU-type examination with...
4. The notified body shall examine the technical documentation to assess...
5. The notified body shall draw up an evaluation report that...
6. Where the type meets the requirements of this Directive that...
7. The notified body shall keep itself apprised of any changes...
8. Each notified body shall inform its notifying authority concerning the...
9. The manufacturer shall keep a copy of the EU-type examination...
10. The manufacturer’s authorised representative may lodge the application referred to...

PART B

Module C: conformity to type based on internal production control...
1. Conformity to type based on internal production control is the...
2. Manufacturing
3. CE marking and EU declaration of conformity
   3.1. The manufacturer shall affix the CE marking to each individual...
   3.2. The manufacturer shall draw up a written EU declaration of...
4. Authorised representative

ANNEX IV

EU declaration of conformity (No Xxxx)

1. Apparatus model/Product (product, type, batch or serial number):

2. Name and address of the manufacturer or his authorised representative:...

3. This declaration of conformity is issued under the sole responsibility...

4. Object of the declaration (identification of apparatus allowing traceability; it...

5. The object of the declaration described above is in conformity...

6. References to the relevant harmonised standards used, including the date...

7. Where applicable, the notified body … (name, number) performed …...

8. Additional information:

ANNEX V

ANNEX VI

(**1**) OJ C 181, 21.6.2012, p. 105.

(**2**) Position of the European Parliament of 5 February 2014 (not yet published in the Official Journal) and decision of the Council of 20 February 2014.

(**3**) OJ L 390, 31.12.2004, p. 24.

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

(**5**) OJ L 218, 13.8.2008, p. 82.

(**6**) OJ L 91, 7.4.1999, p. 10.

(**7**) OJ L 316, 14.11.2012, p. 12.

(**8**) OJ L 55, 28.2.2011, p. 13.