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)

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

Signature

(**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.