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#### **INFORMATION NOTE**

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from: General Secretariat  
to: Permanent Representatives Committee/Council

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Subject: Proposal for a Directive of the European Parliament and of the Council on the harmonisation of the laws of the Member States relating to the making available on the market of radio equipment  
– Outcome of the European Parliament's first reading  
(Strasbourg, 10 to 13 March 2014)

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#### **I. INTRODUCTION**

In accordance with the provisions of Article 294 of the TFEU and the joint declaration on practical arrangements for the codecision procedure <sup>1</sup>, a number of informal contacts have taken place between the Council, the European Parliament and the Commission with a view to reaching an agreement on this dossier at first reading, thereby avoiding the need for second reading and conciliation.

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<sup>1</sup> OJ C 145, 30.6.2007, p.5

In this context, the rapporteur, Mrs Barbara WEILER (S&D-DE) presented one compromise amendment (amendment 99) to the proposal for a Directive, on behalf of the Committee on Internal Market and Consumer Affairs. One further amendment (amendment 100) which referred to a statement of the Parliament was tabled. These amendments had been agreed during the informal contacts referred to above.

## II. VOTE

When it voted on 13 March 2014, the plenary adopted the two amendments (amendments 99 and 100) to the proposal for a Directive. The Commission's proposal as thus amended and the legislative resolution constitute the Parliament's first reading position<sup>1</sup>, it reflects what had been previously agreed between the institutions. The Council should therefore be in a position to approve the Parliament's position.

The act would then be adopted in the wording which corresponds to the Parliament's position.

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<sup>1</sup> The text of the amendment adopted and the European Parliament's legislative resolution are set out in the Annex. The amendment is presented in the form of a consolidated text, where changes to the Commission's proposal are highlighted in *bold and italics*. The symbol "■" indicates deleted text.

## Radio equipment \*\*\*I

**European Parliament legislative resolution of 13 March 2014 on the proposal for a directive of the European Parliament and of the Council on the harmonisation of the laws of the Member States relating to the making available on the market of radio equipment (COM(2012)0584 – C7-0333/2012 – 2012/0283(COD))**

**(Ordinary legislative procedure: first reading)**

*The European Parliament,*

- having regard to the Commission proposal to Parliament and the Council (COM(2012)0584),
  - having regard to Article 294(2) and Article 114 of the Treaty on the Functioning of the European Union, pursuant to which the Commission submitted the proposal to Parliament (C7-0333/2012),
  - having regard to Article 294(3) of the Treaty on the Functioning of the European Union,
  - having regard to the opinion of the European Economic and Social Committee of 13 February 2013<sup>1</sup>,
  - having regard to the undertaking given by the Council representative by letter of 17 January 2014 to approve Parliament's position, in accordance with Article 294(4) of the Treaty on the Functioning of the European Union,
  - having regard to Rule 55 of its Rules of Procedure,
  - having regard to the report of the Committee on the Internal Market and Consumer Protection (A7-0316/2013),
1. Adopts its position at first reading hereinafter set out;
  2. Approves its statement annexed hereto, which will be published in the L series of the *Official Journal of the European Union* together with the final legislative act;
  3. Calls on the Commission to refer the matter to Parliament again if it intends to amend its proposal substantially or replace it with another text;
  4. Instructs its President to forward its position to the Council, the Commission and the national parliaments.

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<sup>1</sup> OJ C 133, 9.5.2013, p. 58.

**P7\_TC1-COD(2012)0283**

**Position of the European Parliament adopted at first reading on 13 March 2014 with a view to the adoption of Directive 2014/.../EU of the European Parliament and of the Council on the harmonisation of the laws of the Member States relating to the making available on the market of radio equipment *and repealing Directive 1999/5/EC***

(Text with EEA relevance)

THE EUROPEAN PARLIAMENT AND THE COUNCIL OF THE EUROPEAN UNION,

Having regard to the Treaty on the Functioning of the European Union, and in particular Article 114 thereof,

Having regard to the proposal from the European Commission,

After transmission of the draft legislative act to the national *parliaments*,

Having regard to the opinion of the Economic and Social Committee<sup>1</sup>,

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Acting in accordance with the ordinary legislative procedure<sup>2</sup>,

Whereas:

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<sup>1</sup> *OJ C 133, 9.5.2013, p. 58.*

<sup>2</sup> Position of the European Parliament of 13 March 2014.

- (1) Directive 1999/5/EC of the European Parliament and of the Council <sup>1</sup> has been substantially amended several times. Since further amendments are to be made, it should be replaced in the interests of clarity.
- (2) Regulation (EC) No 765/2008 of the European Parliament and of the Council <sup>2</sup> lays down rules on the accreditation of conformity assessment bodies, provides a framework for the market surveillance of products and for controls on products from third countries, and lays down the general principles of the CE marking.
- (3) Decision No 768/2008/EC of the European Parliament and of the Council <sup>3</sup> lays down common principles and reference provisions intended to apply across *sectoral* legislation in order to provide a coherent basis for revision or recasts of that legislation. Directive 1999/5/EC should therefore be adapted to that Decision.

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<sup>1</sup> *Directive 1999/5/EC of the European Parliament and of the Council of 9 March 1999 on radio equipment and telecommunications terminal equipment and the mutual recognition of their conformity* (OJ L 91, 7.4.1999, p. 10).

<sup>2</sup> *Regulation (EC) No 765/2008 of the European Parliament and of the Council of 9 July 2008 setting out the requirements for accreditation and market surveillance relating to the marketing of products and repealing Regulation (EEC) No 339/93* (OJ L 218, 13.8.2008, p. 30).

<sup>3</sup> *Decision No 768/2008/EC of the European Parliament and of the Council of 9 July 2008 on a common framework for the marketing of products, and repealing Council Decision 93/465/EEC* (OJ L 218, 13.8.2008, p. 82).

- (4) The essential requirements *laid down* in Directive 1999/5/EC which are relevant to fixed-line terminal equipment, i.e. to ensure the protection of health and safety *of persons and of domestic animals and the protection of property* and an adequate level of electromagnetic compatibility, are appropriately covered by Directive 2014/.../EU of the European Parliament and of the Council <sup>1\*</sup> and Directive 2014/.../EU of the European Parliament and of the Council <sup>2\*\*</sup>. This Directive should therefore not apply to fixed-line terminal equipment.
- (5) Competition issues in the market for terminal equipment are appropriately covered by Commission Directive 2008/63/EC <sup>3</sup>, in particular through the obligation for national regulatory authorities to ensure the publication of details of technical interface specifications for network access. It is therefore not necessary to include in this Directive requirements facilitating competition in the market for terminal equipment covered by Directive 2008/63/EC.

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<sup>1</sup> *Directive 2014/.../EU of the European Parliament and of the Council of ... on the harmonisation of the laws of the Member States relating to the making available on the market of electrical equipment designed for use within certain voltage limits (OJ L ...).*

\* *OJ: please insert number of the Directive (COD 2011/0357) in the recital and the number, date of adoption and publication reference of the Directive in footnote 1.*

<sup>2</sup> *Directive 2014/.../EU of the European Parliament and of the Council of ... on the harmonisation of the laws of the Member States relating to electromagnetic compatibility (OJ L ...).*

\*\* *OJ: please insert number of the Directive (COD 2011/0351) in the recital and the number, date of adoption and publication reference of the Directive in footnote 2.*

<sup>3</sup> *Commission Directive 2008/63/EC of 20 June 2008 on competition in the markets in telecommunications terminal equipment (OJ L 162, 21.6.2008, p. 20).*

(6) Equipment which intentionally *emits or receives* radio waves *for the purpose of radio communication or radiodetermination* makes systematic use of radio spectrum. In order to ensure an efficient use of *radio* spectrum so as to avoid harmful interference, all such equipment should fall within the scope of this Directive █ .

█

(7) The objectives with respect to safety requirements laid down in Directive *2014/.../EU*<sup>\*</sup> are sufficient to cover radio equipment, and should therefore be the reference and made applicable by virtue of this Directive. In order to avoid unnecessary duplications of provisions other than those concerning such requirements, Directive *2014/.../EU*<sup>\*</sup> should not apply to radio equipment.

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<sup>\*</sup> *OJ: please insert the number of the Directive (COD 2011/0357).*



- (8) The essential requirements in the area of electromagnetic compatibility laid down by Directive **2014/.../EU**\* are sufficient to cover radio equipment, and should therefore be the reference and made applicable by virtue of this Directive. In order to avoid unnecessary duplications of provisions other than those concerning essential requirements, Directive **2014/.../EU**\* should not apply to radio equipment.
- (9) *This Directive should apply to all forms of supply, including distance selling.*

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*OJ: please insert the number of the Directive (COD 2011/0351).*

- (10) *In order to ensure that radio equipment uses the radio spectrum effectively and supports the efficient use of radio spectrum, radio equipment should be constructed so that: in the case of a transmitter, when the transmitter is properly installed, maintained and used for its intended purpose it generates radio waves emissions that do not create harmful interference, while unwanted radio waves emissions generated by the transmitter (e.g. in adjacent channels) with a potential negative impact on the goals of radio spectrum policy should be limited to such a level that, according to the state of the art, harmful interference is avoided; and, in the case of a receiver, it has a level of performance that allows it to operate as intended and protects it against the risk of harmful interference, in particular from shared or adjacent channels, and, in so doing, supports improvements in the efficient use of shared or adjacent channels.*
- (11) Although receivers do not themselves cause harmful interference, reception capabilities are an increasingly important factor in ensuring the efficient use of radio spectrum by way of an increased resilience of receivers against harmful interference and unwanted signals on the basis of the *relevant* essential requirements *of Union harmonisation legislation*.

- (12) ■ Interworking via networks with other radio equipment and connection with interfaces of the appropriate type throughout the Union *is necessary in some cases*. Interoperability between radio equipment and accessories such as chargers *simplifies the* use of radio equipment *and reduces* unnecessary waste *and costs*. *A renewed effort to develop a common charger for particular categories or classes of radio equipment is necessary, in particular for the benefit of consumers and other end-users; this Directive should therefore include specific requirements in that area. In particular, mobile phones that are made available on the market should be compatible with a common charger.*
- (13) The protection of personal data and privacy of users and of subscribers of radio equipment and the protection from fraud may be enhanced by particular features of radio equipment. Radio equipment should therefore in appropriate cases be designed in such a way that it supports those features.

- (14) Radio equipment can be instrumental in providing access to emergency services. Radio equipment should therefore in appropriate cases be designed in such a way that it supports the features required for access to those services.
- (15) Radio equipment is important to the well-being and employment of people with disabilities, who represent a substantial and growing proportion of the population of Member States. Radio equipment should therefore in appropriate cases be designed in such a way that people with disabilities may use it without or with only minimal adaptation.
- (16) The compliance of some categories of radio equipment with the essential requirements set out in this Directive may be affected by the inclusion of software or modification of its existing software. The user, the radio equipment or a third party should only be able to load software into the radio equipment where this does not compromise the subsequent compliance of that radio equipment with the applicable essential requirements.

- (17) *In order to supplement or amend certain non-essential elements of this Directive, the power to adopt acts in accordance with Article 290 of the Treaty on the Functioning of the European Union (TFEU) should be delegated to the Commission. It is of particular importance that the Commission carry out appropriate consultations during its preparatory work, including at expert level. The Commission, when preparing and drawing up delegated acts, should ensure a simultaneous, timely and appropriate transmission of relevant documents to the European Parliament and to the Council.*
- (18) *In order to effectively address the needs related to interoperability, protection of **personal data and privacy of the user and of the subscriber**, protection from fraud **■**, access to emergency services, use by users with a disability **■** or the prevention of non-compliant combinations of radio equipment and software **■**, the power to adopt acts in accordance with Article 290 TFEU should be delegated to the Commission in respect of **the** specification of categories or classes of radio equipment that have to comply with **one or more of the** additional essential requirements **set out in this Directive which address those needs**.*

- (19) Verification by radio equipment of the compliance of its combination with software should not be abused in order to prevent its use with software provided by independent parties. The availability to public authorities, manufacturers and users of information on the compliance of intended combinations of radio equipment and software should contribute to facilitate competition. In order to *achieve those objectives*, the power to adopt acts in accordance with Article 290 *TFEU* should be delegated to the Commission in respect of *the specification of* categories or classes of radio equipment for which manufacturers have to provide information on the compliance of intended combinations of radio equipment and software *with the essential requirements set out in this Directive*.

(20) A requirement to register in a central *system* radio equipment to be placed on the market may enhance the efficiency and effectiveness of market surveillance and thereby contribute to ensuring a high level of compliance with *this* Directive. Such a requirement entails additional burden to economic operators and should therefore be introduced only for those categories of radio equipment where a high level of compliance has not been attained. In order to *ensure the application of such a requirement*, the power to adopt acts in accordance with Article 290 *TFEU* should be delegated to the Commission in respect of *the specification of the* ■ *categories of radio equipment which manufacturers have to register within a central system and the elements of the technical documentation to be provided* on the basis of the information on the compliance *of radio equipment* to be provided by Member States *and following an evaluation of the risk of non-implementation of the essential requirements*.

- (21) Radio equipment which complies with the relevant essential requirements should be allowed to circulate freely. Such equipment should be allowed to be put into service and used for its intended purpose, where applicable in accordance with rules on authorisations for the use of radio spectrum and the provision of the service concerned.
- (22) In order to avoid unnecessary barriers to trade in radio equipment within the *internal market*, Member States should notify, under Directive 98/34/EC of the European Parliament and of the Council <sup>1</sup>, other Member States and the Commission of their projects in the area of technical regulations, such as radio interfaces, unless those technical regulations allow Member States to comply with binding Union acts such as Commission decisions on the harmonised use of radio spectrum adopted under Decision No 676/2002/EC of the European Parliament and of the Council <sup>2</sup>, *or where they correspond to radio equipment which can be put into service and used without restrictions within the Union.*

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<sup>1</sup> *Directive 98/34/EC of the European Parliament and of the Council of 22 June 1998 laying down a procedure for the provision of information in the field of technical standards and regulations and of rules on Information Society services* (OJ L 204, 21.7.1998, p. 37).

<sup>2</sup> *Decision No 676/2002/EC of the European Parliament and of the Council of 7 March 2002 on a regulatory framework for radio spectrum policy in the European Community (Radio Spectrum Decision)* (OJ L 108, 24.4.2002, p. 1).



- (23) The provision of information on the equivalence of regulated radio interfaces and their conditions of use reduces barriers for the access of radio equipment to the internal market. The Commission should therefore assess and establish the equivalence of regulated radio interfaces and make such information available in the form of radio equipment classes.
- (24) In accordance with Commission Decision 2007/344/EC<sup>1</sup>, Member States are to use the Frequency Information System (EFIS) of the *European Communications Office (ECO)* in order to make comparable information regarding the use of radio spectrum in each Member State available to the public via the internet. Manufacturers can search in EFIS frequency information for all Member States prior to the placing on the market of radio equipment and thereby evaluate whether and under which conditions such radio equipment may be used within each Member State. There is therefore no need to include in this Directive additional provisions, such as prior notification, allowing manufacturers to be informed of the conditions of use of radio equipment using non-harmonised frequency bands.

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<sup>1</sup> *Commission Decision 2007/344/EC of 16 May 2007 on harmonised availability of information regarding spectrum use within the Community* (OJ L 129, 17.5.2007, p. 67).

- (25) For the purpose of promotion of research and demonstration activities it should be possible, in the context of trade fairs, exhibitions and similar events, to *display* radio equipment which does not comply with this Directive and cannot be placed on the market, on the condition that exhibitors ensure that sufficient information is provided to the visiting public.
- (26) Economic operators should be responsible for the compliance of *radio equipment with this Directive*, in relation to their respective roles in the supply chain, so as to ensure a high level of protection of health and safety *of persons and of domestic animals, and the protection of property, an adequate level* of electromagnetic compatibility, an *effective and efficient* use of *radio* spectrum *and, where necessary, a high level of protection of other public interests*, and to guarantee fair competition on the Union market.

- (27) All economic operators intervening in the supply and distribution chain should take appropriate measures to ensure that they only make available on the market **radio equipment** which *is* in conformity with this Directive. It is necessary to provide for a clear and proportionate distribution of obligations which correspond to the role of each **economic** operator in the supply and distribution **chain**.
- (28) *In order to facilitate communication between economic operators, market surveillance authorities and consumers, Member States should encourage economic operators to include a website address in addition to the postal address.*
- (29) The manufacturer, having detailed knowledge of the design and production process, is best placed to carry out the **CE** conformity assessment procedure. Conformity assessment should therefore remain **solely** the obligation of the manufacturer **CE**.

- (30) The manufacturer should provide sufficient information on the intended use of the **radio** equipment so as to allow its use in compliance with the essential requirements. Such information may need to include a description of accessories such as antennas and of components such as software, and specifications of the installation process of the **radio** equipment.
- (31) The requirement ***laid down*** in Directive 1999/5/EC to include an EU declaration of conformity with equipment has been found to simplify and to enhance the information and the efficiency of market surveillance. The possibility to provide a simplified EU declaration of conformity has allowed ■ the burden associated with this requirement ***to be reduced*** without reduction of its effectiveness, and should therefore be provided for within this Directive. ***Furthermore, in order to ensure easy and efficient access to an EU declaration of conformity, including a simplified EU declaration of conformity, it should be possible to affix it to the packaging of the radio equipment concerned.***

- (32) It is necessary to ensure that *radio equipment* from third countries entering the Union market complies with **■** this Directive, and in particular that appropriate *conformity* assessment procedures have been carried out by manufacturers with regard to *that radio equipment*. Provision should therefore be made for importers to make sure that the *radio equipment* they place on the market complies with the requirements of this Directive and that they do not place on the market *radio equipment* which does not comply with such requirements or presents a risk. Provision should also be made for importers to make sure that conformity assessment procedures have been carried out and that marking *of radio equipment* and documentation drawn up by manufacturers are available for inspection by the *competent national* authorities.

- (33) *When placing radio equipment on the market, every importer should indicate on the radio equipment his name, registered trade name or registered trade mark and the postal address at which he can be contacted. Exceptions should be provided for in cases where the size or nature of the radio equipment does not allow it. This includes cases where the importer would have to open the packaging in order to put his name and address on the radio equipment.*
- (34) *The distributor makes radio equipment available on the market after it has been placed on the market by the manufacturer or the importer and should act with due care to ensure that its handling of the radio equipment does not adversely affect the compliance of the radio equipment.*

- (35) Any economic operator that either places radio equipment on the market under his own name or trade mark or modifies radio equipment in such a way that compliance with **■** this Directive may be affected should be considered to be the manufacturer and should assume the obligations of the manufacturer.
- (36) Distributors and importers, being close to the market place, should be involved in market surveillance tasks carried out by the competent national authorities, and should be prepared to participate actively, providing those authorities with all necessary information relating to the radio equipment concerned.

- (37) Ensuring traceability of radio equipment throughout the whole supply chain helps to make market surveillance simpler and more efficient. An efficient traceability system facilitates market surveillance authorities' task of tracing economic operators who made non-compliant radio equipment available on the market. ***When keeping the information required under this Directive for the identification of other economic operators, economic operators should not be required to update such information in respect of other economic operators who have either supplied them with radio equipment or to whom they have supplied radio equipment.***
- (38) This Directive should be limited to the expression of essential requirements. In order to facilitate conformity assessment with those requirements it is necessary to provide for a presumption of conformity for radio equipment which is in conformity with harmonised standards that are adopted in accordance with Regulation (EU) No 1025/2012 of the ***European Parliament and of the Council***<sup>1</sup> for the purpose of expressing detailed technical specifications of those requirements.

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<sup>1</sup> ***Regulation (EU) No 1025/2012 of the European Parliament and of the Council of 25 October 2012 on European standardisation, amending Council Directives 89/686/EEC and 93/15/EEC and Directives 94/9/EC, 94/25/EC, 95/16/EC, 97/23/EC, 98/34/EC, 2004/22/EC, 2007/23/EC, 2009/23/EC and 2009/105/EC of the European Parliament and of the Council and repealing Council Decision 87/95/EEC and Decision No 1673/2006/EC of the European Parliament and of the Council (OJ L 316, 14.11.2012, p. 12).***



- (39) Regulation (EU) No **1025/2012** provides for a procedure for objections to harmonised standards where those standards do not entirely satisfy *the* requirements of this Directive.
- (40) In order to enable economic operators to demonstrate and the competent authorities to ensure that radio equipment made available on the market conforms to the essential requirements, it is necessary to provide for conformity assessment procedures. Decision No 768/2008/EC establishes modules for conformity assessment procedures, which include procedures from the least to the most stringent, in proportion to the level of risk involved and the level of safety required. In order to ensure inter-sectoral coherence and to avoid ad-hoc variants, conformity assessment procedures should be chosen from among those modules.
- (41) Manufacturers should draw up an EU declaration of conformity to provide **■** information *required under this Directive* on the conformity of radio equipment with the requirements of this Directive and of the other relevant Union harmonisation legislation.

- (42) *To ensure effective access to information for market surveillance purposes, the information required to identify all applicable Union acts should be available in a single EU declaration of conformity. In order to reduce the administrative burden on economic operators, that single EU declaration of conformity may be a dossier made up of relevant individual declarations of conformity.*
- (43) The CE marking, indicating the conformity of *radio equipment*, is the visible consequence of a whole process comprising conformity assessment in a broad sense. General principles governing the CE marking are set out in Regulation (EC) No 765/2008. Rules governing the affixing of the CE marking should be laid down in this Directive.
- (44) The requirement to affix the CE marking on products is important for the information of consumers and public authorities. The possibility laid down in Directive 1999/5/EC to affix a reduced CE mark on small-*sized* equipment, provided that it remains visible and legible, has allowed the application of that requirement to be simplified without reducing its effectiveness, and should therefore be included in this Directive.

- (45) The requirement ***laid down*** in Directive 1999/5/EC to affix the CE marking on the packaging of equipment has been found to simplify the task of market surveillance, and should therefore be included in this Directive.
- (46) *Member States should take appropriate measures to ensure that radio equipment may be made available on the market only if, when properly installed and maintained and used for its intended purpose, it complies with the essential requirements set out in this Directive, and, in the case of the essential requirement to ensure the protection of the health and safety of persons and of domestic animals and the protection of property, also under conditions of use which can be reasonably foreseen. Radio equipment should be considered as non-compliant with that essential requirement only under conditions of use which can be reasonably foreseen, that is when such use could result from lawful and readily predictable human behaviour.*

- (47) *In view of the rapid pace of technological change towards a paperless environment, where radio equipment is fitted with an integral screen, the Commission should examine, as part of a review of the operation of this Directive, the feasibility of replacing the requirements for affixing: the manufacturer's name, registered trade name or registered trade mark and a single point or postal address at which they can be contacted, CE marking and EU declaration of conformity with either a function whereby such information is automatically displayed upon starting up the radio equipment, or a function allowing the end-user to select the display of the relevant information. Furthermore, as part of that examination of feasibility, where radio equipment fitted with an integral screen operates from an integral battery which does not hold an initial charge, the Commission should also consider the use of removable transparent integral screen covering labels which would display the same information.*
- (48) *Certain* conformity assessment procedures set out in this Directive require the intervention of conformity assessment bodies, which are notified by the Member States to the Commission.

- (49) Experience has shown that the criteria set out in Directive 1999/5/EC that conformity assessment bodies have to fulfil to be notified to the Commission are not sufficient to ensure a uniformly high level of performance of notified bodies throughout the Union. It is, however, essential that all notified bodies perform their functions to the same level and under conditions of fair competition. That requires the setting of obligatory requirements for conformity assessment bodies wishing to be notified in order to provide conformity assessment services.
- (50) *If a conformity assessment body demonstrates conformity with the criteria laid down in harmonised standards, it should be presumed to comply with the corresponding requirements set out in this Directive.*
- (51) *In order to ensure a consistent level of conformity assessment quality it is also necessary to set requirements for notifying authorities and other bodies involved in the assessment, notification and monitoring of notified bodies.*

- (52) The system set out in this Directive should be complemented by the accreditation system provided for in Regulation (EC) No 765/2008. Since accreditation is an essential means of verifying the competence of conformity assessment bodies, it should also be used for the purposes of notification.
- (53) Transparent accreditation as provided for in Regulation (EC) No 765/2008, ensuring the necessary level of confidence in conformity certificates, should be considered by the national public authorities throughout the Union as the preferred means of demonstrating the technical competence of conformity assessment bodies. However, national authorities may consider that they possess the appropriate means of carrying out *that* evaluation themselves. In such cases, in order to ensure the appropriate level of credibility of evaluations carried out by other national authorities, they should provide the Commission and the other Member States with the necessary documentary evidence demonstrating the compliance of the conformity assessment bodies evaluated with the relevant regulatory requirements.

- (54) Conformity assessment bodies frequently subcontract parts of their activities linked to the assessment of conformity or have recourse to a subsidiary. In order to safeguard the level of protection required for radio equipment to be placed on the Union market, it is essential that conformity assessment subcontractors and subsidiaries fulfil the same requirements as notified bodies in relation to the performance of conformity assessment tasks. Therefore, it is important that the assessment of the competence and the performance of bodies to be notified and the monitoring of bodies already notified cover also activities carried out by subcontractors and subsidiaries.
- (55) It is necessary to increase the efficiency and transparency of the notification procedure and, in particular, to adapt it to new technologies so as to enable online notification.

- (56) Since notified bodies may offer their services throughout the Union, it is appropriate to give the other Member States and the Commission the opportunity to raise objections concerning a notified body. It is therefore important to provide for a period during which any doubts or concerns as to the competence of conformity assessment bodies can be clarified before they start operating as notified bodies.
- (57) In the interests of competitiveness, it is crucial that notified bodies apply the conformity assessment procedures without creating unnecessary burdens for economic operators. For the same reason, and to ensure equal treatment of economic operators, consistency in the technical application of the conformity assessment procedures needs to be ensured. That can best be achieved through appropriate coordination and cooperation between notified bodies.



- (58) In order to ensure legal certainty, it is necessary to clarify that rules on Union market surveillance and control of products entering the Union market provided for in Regulation (EC) No 765/2008 apply to radio equipment *covered by this Directive. This Directive should not prevent Member States from choosing the competent authorities to carry out those tasks.*
- (59) Directive 1999/5/EC already provides for a safeguard procedure which applies only in the event of disagreement between Member States over measures taken by a Member State. In order to increase transparency and to reduce processing time, it is necessary to improve the existing safeguard procedure, with a view to making it more efficient and drawing on the expertise available in Member States.

(60) *The decisions of the Commission adopted under Decision No 676/2002/EC may include conditions for the availability and efficient use of radio spectrum which may have as a consequence the limitation of the total number of items of radio equipment put into service, such as a "sunset" date, a maximum penetration rate or a maximum number of items of radio equipment in each Member State or throughout the Union. Those conditions enable the market to be opened up to new radio equipment while limiting the risk of harmful interference by accumulation of an excessive number of items of radio equipment put into service, even though that equipment individually complies with the essential requirements set out in this Directive. Infringing such conditions may create a risk to the essential requirements, particularly a risk of harmful interference.*

- (61) The existing system should be supplemented by a procedure under which interested parties are informed of measures intended to be taken with regard to radio equipment presenting a risk to the health or safety of persons or to other aspects of public interest **protection** covered by █ this Directive. It should also allow market surveillance authorities, in cooperation with the relevant economic operators, to act at an earlier stage in respect of such equipment.
- (62) Where the Member States and the Commission agree as to the justification of a measure taken by a Member State, no further involvement of the Commission should be required, except where non-compliance can be attributed to shortcomings of a harmonised standard.
- (63) In order to ensure uniform conditions for the implementation of this Directive, implementing powers should be conferred on the Commission. Those powers should be exercised in accordance with Regulation (EU) No 182/2011 of the European Parliament and of the Council █<sup>1</sup>.

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<sup>1</sup> ***Regulation (EU) No 182/2011 of the European Parliament and of the Council of 16 February 2011 laying down the rules and general principles concerning mechanisms for control by Member States of the Commission's exercise of implementing powers*** (OJ L 55 █, 28.2.2011, p. 13).

- (64) *The advisory procedure should be used for the adoption of implementing acts specifying how to present information in cases of restrictions on putting into service or of existing requirements for authorisation of use; and requesting the notifying Member State to take the necessary corrective measures in respect of a notified body that does not meet or no longer meets the requirements for its notification.*
- (65) *The examination procedure should be used for the adoption of implementing acts: determining whether certain categories of electrical or electronic products meet the definition of “radio equipment”; laying down the operational rules for making the information on compliance available; laying down the operational rules for registration and the operational rules for affixing the registration number on radio equipment; and establishing the equivalence between notified radio interfaces and assigning a radio equipment class. It should also be used with respect to compliant radio equipment which presents a risk to the health or safety of persons or to other aspects of public interest protection.*

- (66) *The Commission should adopt immediately applicable implementing acts where, in duly justified cases relating to compliant radio equipment which presents a risk to the health or safety of persons, imperative grounds of urgency so require.*
- (67) *In line with established practice, the committee set up by this Directive can play a useful role in examining matters concerning the application of this Directive raised either by its chair or by a representative of a Member State in accordance with its rules of procedure.*
- (68) *When matters relating to this Directive, other than its implementation or infringements, are being examined, i.e. in a Commission expert group, the European Parliament should in line with existing practice receive full information and documentation and, where appropriate, an invitation to attend such meetings.*

(69) *The Commission should, by means of implementing acts and, given their special nature, acting without the application of Regulation (EU) No 182/2011, determine whether measures taken by Member States in respect of non-compliant radio equipment are justified or not.*

■

(70) The Member States should lay down rules on penalties applicable to infringements of the ■ provisions *of national law* adopted pursuant to this Directive and ensure that *those rules* are *enforced*. *The* penalties *provided for* should be effective, proportionate and dissuasive.

- (71) It is necessary to provide for transitional arrangements that allow the making available on the market and putting into service of radio equipment that has already been placed on the market in accordance with Directive 1999/5/EC.
- (72) *The European Data Protection Supervisor has been consulted.*
- (73) Since the objective of this Directive, namely to ensure that radio equipment made available on the market fulfils requirements providing a high level of protection of health and safety, adequate level of electromagnetic compatibility and an *effective and* efficient use of *radio* spectrum so as to avoid harmful interference while guaranteeing the proper functioning of the internal market, cannot be sufficiently achieved by the Member States *but* can *rather*, by reason of its scale and effects, be better achieved at Union level, the Union may adopt measures, in accordance with the principle of subsidiarity as set out in Article 5 of the Treaty on European Union. In accordance with the principle of proportionality, as set out in that Article, this Directive does not go beyond what is necessary in order to achieve that objective.

(74) *Directive 1999/5/EC should be repealed.*

(75) In accordance with the Joint Political Declaration *of 28 September 2011* of Member States and the Commission on explanatory documents<sup>1</sup>, Member States have undertaken to accompany, in justified cases, the notification of their transposition measures with one or more documents explaining the relationship between the components of a directive and the corresponding parts of national transposition instruments. With regard to this Directive, the legislator considers the transmission of such documents to be justified,

HAVE ADOPTED THIS DIRECTIVE:

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<sup>1</sup> *OJ C 369, 17.12.2011. p. 14.*



CHAPTER I  
GENERAL PROVISIONS

Article 1

Subject matter and scope

1. This Directive establishes a regulatory framework for the making available on the market and putting into service in the Union of radio equipment.
2. This Directive shall not apply to equipment listed in Annex I.
3. This Directive shall not apply to radio equipment exclusively used for activities concerning public security, defence, State security, including the economic well-being of the State in the case of activities pertaining to State security matters, and the activities of the State in the area of criminal law.
4. Radio equipment falling within the scope of this Directive shall not be subject to Directive **2014/.../EU\***, except as set out in ***point (a) of*** Article 3(1) **■** of this Directive.

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\*

***OJ: please insert the number of the Directive (COD 2011/0357).***

Article 2  
Definitions

1. For the purposes of this Directive, the following definitions **■** apply:
  - (1) ‘radio equipment’ means *an electrical or electronic* product, which intentionally emits *and/or receives* radio waves *for the purpose of radio communication and/or radiodetermination*, or *an electrical or electronic* product which must be completed with an accessory, such as antenna, so as to *intentionally* emit *and/or receive* radio waves *for the purpose of radio communication and/or radiodetermination*;
  - (2) ‘radio communication’ means *communication by means of radio waves*;
  - (3) ‘radiodetermination’ means *the determination of the position, velocity and/or other characteristics of an object, or the obtaining of information relating to those parameters, by means of the propagation properties of radio waves*;

- (4) ‘radio waves’ means electromagnetic waves of frequencies *lower than* 3 000 GHz, propagated in space without artificial guide;
- (5) ‘*radio* interface’ means *the specification of the regulated use of radio spectrum*;
- (6) ‘radio equipment class’ means a class identifying particular categories of radio equipment which, under this Directive, are considered similar and those *radio* interfaces for which the *radio* equipment is designed;
- (7) ‘harmful interference’ means harmful interference as defined in *point (r) of Article 2 of Directive 2002/21/EC* of the European Parliament and of the Council<sup>1</sup>;

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<sup>1</sup> *Directive 2002/21/EC of the European Parliament and of the Council of 7 March 2002 on a common regulatory framework for electronic communications networks and services (Framework Directive)* (OJ L 108, 24.4.2002, p. 33).

- (8) *‘electromagnetic disturbance’ means electromagnetic disturbance as defined in point 5 of Article 3(1) of Directive 2014/.../EU\**;
- (9) ‘making available on the market’ means any supply of radio equipment for distribution, consumption or use *on* the **█** Union market in the course of a commercial activity, whether in return for payment or free of charge;
- (10) ‘placing on the market’ means the first making available of radio equipment on the **█** Union market;
- (11) *‘putting into service’ means the first use of radio equipment in the Union by its end-user;*

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\*

*OJ: please insert the number of the Directive (COD 2011/0351).*

- (12) ‘manufacturer’ means any natural or legal person who manufactures radio equipment or has radio equipment designed or manufactured, and markets that equipment under his name or trade mark;
- (13) ‘authorised representative’ means any natural or legal person established within the **EU** Union who has received a written mandate from a manufacturer to act on his behalf in relation to specified tasks;
- (14) ‘importer’ means any natural or legal person established within the **EU** Union who places radio equipment from a third country on the **EU** Union market;
- (15) ‘distributor’ means any natural or legal person in the supply chain, other than the manufacturer or the importer, who makes radio equipment available on the market;

- (16) ‘economic operators’ means the manufacturer, the authorised representative, the importer and the distributor;
- (17) ‘technical specification’ means a document that prescribes technical requirements to be fulfilled by radio equipment;
- (18) ‘harmonised standard’ means harmonised standard as defined in *point (c) of point 1 of Article 2* of Regulation (EU) No **1025/2012**;
- (19) ‘accreditation’ means accreditation as defined in *point 10 of Article 2 of Regulation (EC) No 765/2008*;
- (20) ‘national accreditation body’ means national accreditation body as defined in *point 11 of Article 2 of Regulation (EC) No 765/2008*;

- (21) ‘conformity assessment’ means the process demonstrating whether the essential requirements *of this Directive* relating to radio equipment have been fulfilled;
- (22) ‘conformity assessment body’ means a body that performs conformity assessment activities;
- (23) ‘recall’ means any measure aimed at achieving the return of radio equipment that has already been made available to the *end-user*;
- (24) ‘withdrawal’ means any measure aimed at preventing radio equipment in the supply chain from being made available on the market;

(25) *'Union harmonisation legislation' means any Union legislation harmonising the conditions for the marketing of products;*

(26) *'CE marking' means a marking by which the manufacturer indicates that the radio equipment is in conformity with the applicable requirements set out in Union harmonisation legislation providing for its affixing.*

2. *The Commission may adopt implementing acts to determine whether certain categories of electrical or electronic products meet the definition set out in point 1 of paragraph 1 of this Article. Those implementing acts shall be adopted in accordance with the examination procedure referred to in Article 45(3).*

■



Article 3  
Essential requirements

1. Radio equipment shall be constructed so as to ensure:
  - (a) the protection of health and safety of *persons and of domestic animals and the protection of property*, including the objectives with respect to safety requirements set out in *Directive 2014/.../EU\**, but with no voltage limit applying;
  - (b) *an adequate level* of electromagnetic compatibility as set out in *Directive 2014/.../EU\*\**.
2. Radio equipment shall be so constructed that *it both effectively uses and supports the efficient use of radio spectrum in order* to avoid harmful interference.

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\* *OJ: please insert the number of the Directive (COD 2011/0357).*

\*\* *OJ: please insert the number of the Directive (COD 2011/0351).*

3. Radio equipment *within certain categories or classes* shall be so constructed that it complies with the following essential requirements:
- (a) radio equipment interworks with accessories, *in particular with common chargers*;
  - (b) *radio equipment interworks via networks with other radio equipment*;
  - (c) *radio equipment can be connected to interfaces of the appropriate type throughout the Union*;
  - (d) radio equipment does not harm the network or its functioning nor misuse network resources, thereby causing an unacceptable degradation of service;
  - (e) radio equipment incorporates safeguards to ensure that the personal data and privacy of the user and of the subscriber are protected;

- (f) radio equipment supports certain features ensuring *protection from* fraud;
- (g) radio equipment supports certain features ensuring access to emergency services;
- (h) radio equipment supports certain features in order to facilitate its use by users with a disability;
- (i) radio equipment supports certain features in order to ensure that software can only be loaded into the radio equipment where the compliance of the combination of the radio equipment and software has been demonstrated.

The Commission shall be empowered to adopt delegated acts in accordance with Article 44 specifying which categories or classes of radio equipment are concerned by each of the requirements set out in points (a) to (i) of the first subparagraph of this paragraph.

## Article 4

### Provision of information on the compliance of combinations of radio equipment and software

1. Manufacturers of radio equipment and of software allowing radio equipment to be used as intended shall provide the Member States and the Commission with information on the compliance of intended combinations of radio equipment and software with the essential requirements set out in Article 3. ***Such information shall result from a conformity assessment carried out in accordance with Article 17, and shall be given in the form of a statement of compliance which includes the elements set out in Annex VI. Depending on the specific combinations of radio equipment and software, the information shall precisely identify the radio equipment and the software which have been assessed, and it shall be continuously updated.***

2. The Commission shall be empowered to adopt delegated acts in accordance with Article 44 specifying which categories or classes of radio equipment are concerned by the requirement *set out* in **■** paragraph 1 *of this Article*.
3. ***The Commission shall adopt implementing acts laying down the operational rules for making the information on compliance available for the categories and classes specified by the delegated acts adopted pursuant to paragraph 2 of this Article. Those implementing acts shall be adopted in accordance with the examination procedure referred to in Article 45(3).***

## Article 5

### Registration of radio equipment *types* within some categories

1. As from ...<sup>\*</sup>, manufacturers shall register radio equipment types within categories of radio equipment affected by a low level of compliance with the essential requirements set out in Article 3 within a central system referred to in paragraph 4 of this Article prior to radio equipment within those categories being placed on the market. ***When registering such radio equipment types, manufacturers shall provide some, or where justified all, elements of the technical documentation listed in points (a), (d), (e), (f), (g), (h) and (i) of Annex V.*** The Commission shall allocate to each registered radio equipment type a registration number, which manufacturers shall affix on radio equipment placed on the market.
2. The Commission shall be empowered to adopt delegated acts in accordance with Article 44 specifying which categories of radio equipment are concerned by the requirement set out in **■** paragraph 1 ***of this Article, and the elements of the technical documentation to be provided,*** taking into account the information on the compliance of radio equipment provided by Member States in accordance with Article 47(1) ***and following an evaluation of the risk of non-implementation of the essential requirements.***

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***OJ: please insert the date: four years after the date of entry into force of this Directive.***

3. *The Commission shall adopt implementing acts laying down the operational rules for registration and the operational rules for affixing the registration number on radio equipment for the categories specified by the delegated acts adopted pursuant to paragraph 2 of this Article. Those implementing acts shall be adopted in accordance with the examination procedure referred to in Article 45(3).*
4. The Commission shall make available a central system allowing manufacturers to register the required information. *That system shall ensure appropriate control of access to information of confidential nature.*
5. *Following the date of application of a delegated act adopted pursuant to paragraph 2 of this Article, the reports prepared in accordance with Article 47(1) and (2) shall evaluate its impacts.*

## Article 6

### ***Making available*** on the market

Member States shall ***take appropriate measures to*** ensure that radio equipment is made available on the market only if it complies with this Directive **■** .

## Article 7

### Putting into service and use

Member States shall allow the putting into service and use of radio equipment ***if*** it complies with this Directive ***when it is properly installed, maintained and used for its intended purpose***. Without prejudice to their obligations under Decision No 676/2002/EC and to the conditions attached to authorisations for the use of frequencies in conformity with Union law, in particular under ***Article 9(3) and ■ (4)*** of Directive 2002/21/EC, Member States may only introduce additional requirements for the putting into service and/or use of radio equipment for reasons related to the ***effective and efficient*** use of the radio spectrum, to the avoidance of harmful interference, ***to the avoidance of electromagnetic disturbances*** or to public health.



## Article 8

### Notification of *radio* interface specifications and assignment of radio equipment classes

1. Member States shall notify, in accordance with the procedure set out in Directive 98/34/EC, the *radio* interfaces which they intend to regulate *except*:
  - (a) *the radio interfaces which fully and without any deviation comply with the Commission decisions on the harmonised use of radio spectrum adopted pursuant to Decision No 676/2002/EC; and*
  - (b) *the radio interfaces which, in accordance with implementing acts adopted pursuant to paragraph 2 of this Article, correspond to radio equipment which can be put into service and used without restrictions within the Union.*
  
2. The Commission shall *adopt implementing acts* establishing the equivalence between notified *radio* interfaces and assigning a radio equipment class, details of which shall be published in the *Official Journal of the European Union*. Those implementing acts shall be adopted in accordance with the *examination* procedure referred to in Article 45(3).

## Article 9

### Free movement of radio equipment

1. Member States shall not **■** impede, for reasons relating to **■** aspects covered by this Directive, the *making available* on the market in their territory of radio equipment *which complies with* this Directive.
2. At trade fairs, exhibitions **■** and similar events, Member States shall not create any obstacles to the display of radio equipment which does not comply with this Directive, provided that a visible sign clearly indicates that such radio equipment may not be *made available on the market or put into service* until it has been *brought into conformity with this Directive. Demonstration of radio equipment may only take place provided that adequate measures, as prescribed by Member States, have been taken to avoid harmful interference, electromagnetic disturbances and risk to the health or safety of persons or of domestic animals or to property.*

CHAPTER II  
OBLIGATIONS OF ECONOMIC OPERATORS

Article 10

Obligations of manufacturers

1. When placing *their* radio equipment on the market, manufacturers shall ensure that it has been designed and manufactured in accordance with the essential requirements set out in Article 3.
2. *Manufacturers shall ensure that radio equipment shall be so constructed that it can be operated in at least one Member State without infringing applicable requirements on the use of radio spectrum.*
3. Manufacturers shall draw up the technical documentation referred to in Article 21 and carry out the *relevant* conformity assessment procedure referred to in Article 17 or have it carried out.

Where compliance of radio equipment with the applicable requirements has been demonstrated by that *conformity assessment* procedure, manufacturers shall draw up an EU declaration of conformity and affix the CE marking.

4. Manufacturers shall keep the technical documentation and the EU declaration of conformity for 10 years after *the* radio equipment has been placed on the market.
5. Manufacturers shall ensure that procedures are in place for series production to remain in conformity *with this Directive*. Changes in radio equipment design or characteristics and changes in the harmonised standards or in *other* technical specifications by reference to which conformity of *radio equipment* is declared shall be adequately taken into account.

When deemed appropriate with regard to the risks presented by radio equipment, manufacturers shall, *to protect the health and safety of end-users*, carry out sample testing of radio equipment made available on the market, investigate, and, if necessary, keep a register of complaints, of non-conforming radio equipment and radio equipment recalls, and shall keep distributors informed of any such monitoring.

6. Manufacturers shall ensure that radio equipment *which they have placed on the market* bears a type, batch or serial number or other element allowing its identification, or, where the size or nature of the radio equipment does not allow it, that the required information is provided on the packaging, or in a document accompanying the radio equipment.

7. Manufacturers shall indicate *on the radio equipment* their name, registered trade name or registered trade mark and the *postal* address at which they can be contacted ■ or, where the size or nature of radio equipment does not allow it, on its packaging, or in a document accompanying the radio equipment. The address shall indicate a single point at which the manufacturer can be contacted. ***The contact details shall be in a language easily understood by end-users and market surveillance authorities.***
  
8. Manufacturers shall ensure that *the* radio equipment is accompanied by instructions and safety information in a language which can be easily understood by consumers and other *end-users*, as determined by the Member State concerned. Instructions shall include the information required to use radio equipment in accordance with its intended use. Such information shall include, where applicable, a description of accessories and components, including software, which allow the radio equipment to operate as intended. ***Such instructions and safety information, as well as any labelling, shall be clear, understandable and intelligible.***

The following information shall also be included *in the case of radio equipment intentionally emitting radio waves*:

- (a) frequency band(s) in which the radio equipment operates;
- (b) *maximum* radio-frequency power transmitted in the frequency band(s) in which the radio equipment operates.

9. *Manufacturers shall ensure that each item of radio equipment is accompanied by* a copy of the  EU declaration of conformity *or by* a simplified EU declaration of conformity. Where  a simplified EU declaration of conformity is provided, it shall *contain* the exact internet  address where the full *text of the* EU declaration of conformity can be obtained.

10. *In cases of restrictions on putting into service or of requirements for authorisation of use*, information available on the packaging shall allow the identification of the Member States or the geographical area within a Member State where *restrictions on putting into service or requirements for authorisation of use exist*. Such information shall be completed in the instructions accompanying the radio equipment. The Commission may adopt implementing acts specifying how to present that information. Those implementing acts shall be adopted in accordance with the advisory procedure referred to in Article 45(2).



11. Manufacturers who consider or have reason to believe that radio equipment which they have placed on the market is not in conformity with this Directive shall immediately take the corrective measures *necessary* to bring that radio equipment into conformity, to withdraw it or recall it, if appropriate. Furthermore, where the radio equipment presents a risk, manufacturers shall immediately inform the competent national authorities of the Member States in which they made *the* radio equipment available *on the market* to that effect, giving details, in particular, of the non-compliance, of any corrective measures taken *and of the results thereof*.
12. Manufacturers shall, further to a reasoned request from a competent national authority, provide it with all the information and documentation *in paper or electronic form* necessary to demonstrate the conformity of *the* radio equipment *with this Directive*, in a language which can be easily understood by that authority. They shall cooperate with that authority, at its request, on any action taken to eliminate the risks posed by radio equipment which they have placed on the market.

## Article 11

### Authorised representatives

1. A manufacturer may, by a written mandate, appoint an authorised representative.

The obligations laid down in Article 10(1) and the *obligation to draw up* ■ technical documentation *laid down in Article 10(3)* shall not form part of the authorised representative's mandate.

2. An authorised representative shall perform the tasks specified in the mandate received from the manufacturer. The mandate shall allow the authorised representative to do at least the following:
  - (a) keep the EU declaration of conformity and the technical documentation at the disposal of national *market* surveillance authorities for 10 years after the radio equipment has been placed on the market;

- (b) further to a reasoned request from a competent national authority, provide that authority with all the information and documentation necessary to demonstrate the conformity of radio equipment;
- (c) cooperate with the competent national authorities, at their request, on any action taken to eliminate the risks posed by radio equipment covered by the authorised representative's mandate.

## Article 12

### Obligations of importers

1. Importers shall place only compliant radio equipment on the market.

2. Before placing radio equipment on the market importers shall ensure that the appropriate conformity assessment procedure *referred to in Article 17* has been carried out by the manufacturer *and that the radio equipment is so constructed that it can be operated in at least one Member State without infringing applicable requirements on the use of radio spectrum*. They shall ensure that the manufacturer has drawn up the technical documentation, that the radio equipment bears the CE marking and is accompanied by the information *and documents* referred to in Article 10(8), (9) and (10), and that the manufacturer has complied with the requirements set out in Article 10(6) and (7).

Where an importer considers or has reason to believe that radio equipment is not in conformity with the essential requirements set out in Article 3, he shall not place the radio equipment on the market until it has been brought into conformity. Furthermore, where *the* radio equipment presents a risk, the importer shall inform the manufacturer and the market surveillance authorities to that effect.

3. Importers shall indicate *on the radio equipment* their name, registered trade name or registered trade mark and the *postal* address at which they can be contacted ■ or, where that is not possible, on its packaging or in a document accompanying the radio equipment. This includes cases where the size of radio equipment does not allow it, or where importers would have to open the packaging in order to indicate their name and address on the radio equipment. *The contact details shall be in a language easily understood by end-users and market surveillance authorities.*
4. Importers shall ensure that the radio equipment is accompanied by instructions and safety information in a language which can be easily understood by consumers and other *end-users*, as determined by the Member State concerned.
5. Importers shall ensure that, while radio equipment is under their responsibility, *its* storage or transport conditions do not jeopardise its compliance with the essential requirements set out in Article 3.

6. When deemed appropriate with regard to the risks presented by radio equipment, importers shall, to protect the health and safety of *end-users*, carry out sample testing of radio equipment made available on the market, investigate, and, if necessary, keep a register of complaints, of non-conforming radio equipment and radio equipment recalls, and shall keep distributors informed of *any* such monitoring.
  
7. Importers who consider or have reason to believe that radio equipment which they have placed on the market is not in conformity with this Directive shall immediately take the corrective measures necessary to bring that radio equipment into conformity, to withdraw it or recall it, if appropriate. Furthermore, where the radio equipment presents a risk, importers shall immediately inform the competent national authorities of the Member States in which they made the radio equipment available *on the market* to that effect, giving details, in particular, of the non-compliance and of any corrective measures taken.

8. Importers shall, for **10** years after the radio equipment has been placed on the market, keep a copy of the EU declaration of conformity at the disposal of the market surveillance authorities and ensure that the technical documentation can be made available to those authorities, upon request.
9. Importers shall, further to a reasoned request from a competent national authority, provide it ■ with all the information and documentation *in paper or electronic form* necessary to demonstrate the conformity of radio equipment in a language which can be easily understood by that authority. They shall cooperate with that authority, at its request, on any action taken to eliminate the risks posed by ■ radio equipment which they have placed on the market.

## Article 13

### Obligations of distributors

1. When making radio equipment available on the market distributors shall act with due care in relation to the requirements of this Directive.
2. Before making radio equipment available on the market distributors shall verify that the radio equipment bears the **■** CE marking, that it is accompanied by the **■** documents *required by this Directive* and by *the* instructions and safety information in a language which can be easily understood by consumers and other *end-users* in the Member State in which the radio equipment is to be made available on the market, and that the manufacturer and the importer have complied with the requirements set out in *Article 10(2) and* (6) to (10) and Article 12(3) *respectively*.



Where a distributor considers or has reason to believe that radio equipment is not in conformity with the essential requirements set out in Article 3, he shall not make the radio equipment available on the market until it has been brought into conformity. Furthermore, where the radio equipment presents a risk, the distributor shall inform the manufacturer or the importer to that effect as well as the market surveillance authorities.

3. Distributors shall ensure that, while radio equipment is under their responsibility, *its* storage or transport conditions do not jeopardise its compliance with the essential requirements set out in Article 3.

4. Distributors who consider or have reason to believe that radio equipment which they have made available on the market is not in conformity with this Directive shall make sure that the corrective measures necessary to bring that radio equipment into conformity, to withdraw it or recall it, if appropriate, are taken. Furthermore, where the radio equipment presents a risk, distributors shall immediately inform the competent national authorities of the Member States in which they made the radio equipment available *on the market* to that effect, giving details, in particular, of the non-compliance and of any corrective measures taken.
  
5. Distributors shall, further to a reasoned request from a competent national authority, provide it **■** with all the information and documentation *in paper or electronic form* necessary to demonstrate the conformity of radio equipment. They shall cooperate with that authority, at its request, on any action taken to eliminate the risks posed by radio equipment which they have made available on the market.

## Article 14

### Cases in which obligations of manufacturers apply to importers and distributors

An importer or distributor shall be considered a manufacturer for the purposes of this Directive and he shall be subject to the obligations of the manufacturer under Article 10, where he places radio equipment on the market under his name or trade mark or modifies radio equipment already placed on the market in such a way that compliance with ■ this Directive may be affected.

## Article 15

### Identification of economic operators

Economic operators shall, on request, identify the following to the market surveillance authorities:

- (a) any economic operator who has supplied them with radio equipment;
- (b) any economic operator to whom they have supplied radio equipment.

Economic operators shall be able to present the information referred to in the first paragraph for ■ 10 years after they have been supplied with the radio equipment and for ■ 10 years after they have supplied the radio equipment.

CHAPTER III  
CONFORMITY OF RADIO EQUIPMENT

Article 16

Presumption of conformity *of radio equipment*

Radio equipment which is in conformity with harmonised standards or parts thereof the references of which have been published in the *Official Journal of the European Union* shall be presumed to be in conformity with the essential requirements *set out in Article 3* covered by those standards or parts thereof .

■

## Article 17

### Conformity assessment procedures

1. *The manufacturer shall perform a conformity assessment of the radio equipment with a view to meeting the essential requirements set out in Article 3. The conformity assessment shall take into account all intended operating conditions and, for the essential requirement set out in point (a) of Article 3(1), the assessment shall also take into account the reasonably foreseeable conditions. Where the radio equipment is capable of taking different configurations, the conformity assessment shall confirm whether the radio equipment meets the essential requirements set out in Article 3 in all possible configurations.*
2. Manufacturers *shall* demonstrate compliance of radio equipment with the essential requirements *set out* in *Article 3(1)* ■ using any of the following conformity assessment procedures:

- (a) internal production control ■ set out in Annex II;
- (b) EU-type examination *that is* followed by the conformity to type *based on internal production control* set out in Annex III;
- (c) conformity based on full quality assurance ■ set out in Annex IV.

3. Where, in assessing the compliance of radio equipment with the essential requirements *set out* in *Article* 3(2) and (3), the manufacturer has applied harmonised standards the references of which have been published in the *Official Journal of the European Union*, he *shall* use any of the following procedures:

- (a) internal production control ■ set out in Annex II;
- (b) EU-type examination *that is* followed by the conformity to type *based on internal production control* set out in Annex III;
- (c) conformity based on full quality assurance ■ set out in Annex IV.

4. Where, in assessing the compliance of radio equipment with the essential requirements *set out* in *Article* 3(2) and (3), the manufacturer has not applied or has applied only in part harmonised standards the references of which have been published in the *Official Journal of the European Union*, or where such harmonised standards do not exist, radio equipment shall be submitted with regard to those essential requirements to either of the following procedures:
- (a) EU-type examination *that is* followed by the conformity to type *based on internal production control* set out in Annex III;
  - (b) conformity based on full quality assurance ■ set out in Annex IV.

## Article 18

### EU declaration of conformity

1. The EU declaration of conformity shall state that the fulfilment of the essential requirements set out in Article 3 has been demonstrated.
2. The EU declaration of conformity shall have the model structure *set out in Annex VI*, shall contain the elements set out in *that* Annex ■ and shall be continuously updated. It shall be translated into the language or languages required by the Member State in which ■ the radio equipment is placed or made available *on the market*.

The simplified EU declaration of conformity referred to in Article 10(9) shall contain the elements set out in Annex VII and shall be continuously updated. It shall be translated into the language or languages required by the Member State in which the radio equipment is placed or made available on the market. The full *text of the* EU declaration of conformity shall be available at the internet ■ address referred to in the simplified EU declaration of conformity, in a language or languages required by the Member State in which the radio equipment is placed or made available on the market.



3. Where radio equipment is subject to more than one Union act requiring an EU declaration of conformity, a single EU declaration of conformity shall be drawn up in respect of all such Union acts. That declaration shall contain the identification of the *Union* acts concerned including *their* publication references.
4. By drawing up the EU declaration of conformity, the manufacturer shall assume responsibility for the compliance of the radio equipment *with the requirements laid down in this Directive*.

#### Article 19

##### General principles of the CE marking

1. The CE marking shall be subject to the general principles set out in Article 30 of Regulation (EC) No 765/2008.
2. On account of the nature of radio equipment, the height of the CE marking affixed to radio equipment may be lower than 5 mm, provided that it remains visible and legible.

## Article 20

### Rules and conditions for affixing the CE marking *and the identification number of the notified body*

1. The CE marking shall be affixed visibly, legibly and indelibly to the radio equipment or to its data plate, unless that is not possible or not warranted on account of the nature of radio equipment. The CE marking shall also be affixed visibly and legibly to the packaging.
2. The CE marking shall be affixed before the radio equipment is placed on the market.
3. The CE marking shall be followed by the identification number of the notified body where the conformity assessment procedure set out in Annex IV is applied.

The identification number of the notified body shall have the same height as the CE marking.

*The identification number of the notified body* shall be affixed by the *notified* body itself or, under its instructions, by the manufacturer or his authorised representative.

4. *Member States shall build upon existing mechanisms to ensure correct application of the regime governing the CE marking and shall take appropriate action in the event of improper use of that marking.*

#### Article 21

##### Technical documentation

1. The technical documentation shall contain all relevant data or details of the means used by the manufacturer to ensure that radio equipment complies with the *essential* requirements set out in Article 3. It shall, at least, contain the *elements* set out in Annex V.
2. The technical documentation shall be drawn up before radio equipment is placed on the market and shall be continuously updated.

3. The technical documentation and correspondence relating to any EU-type examination procedure shall be drawn up in an official language of the Member State in which the notified body is established or in a language acceptable to that body.



4. Where the technical documentation does not comply with paragraphs 1, 2 or 3 of this Article, and in so doing fails to present sufficient relevant data or means used to ensure compliance of radio equipment with the essential requirements set out in Article 3, the market surveillance authority may *ask* the manufacturer *or the importer* to have a test performed by a body acceptable to the market surveillance authority at the expense of the manufacturer *or the importer* within a specified period in order to verify compliance with the essential requirements set out in Article 3.

CHAPTER IV  
NOTIFICATION OF CONFORMITY ASSESSMENT BODIES

Article 22  
Notification

Member States shall notify the Commission and the other Member States of bodies authorised to carry out third-party conformity assessment tasks under this Directive.

Article 23  
Notifying authorities

1. Member States shall designate a notifying authority that shall be responsible for setting up and carrying out the necessary procedures for the assessment and notification of conformity assessment bodies and the monitoring of notified bodies, including compliance with Article 28.

2. Member States may decide that the assessment and monitoring referred to in paragraph 1 shall be carried out by a national accreditation body within the meaning of and in accordance with Regulation (EC) No 765/2008.
3. Where the notifying authority delegates or otherwise entrusts the assessment, notification or monitoring referred to in paragraph 1 to a body which is not a governmental entity, that body shall be a legal entity and shall comply *mutatis mutandis* with the requirements laid down in Article 24 ■ . In addition it shall have arrangements to cover liabilities arising out of its activities.
4. The notifying authority shall take full responsibility for the tasks performed by the body referred to in paragraph 3.

## Article 24

### Requirements relating to notifying authorities

1. A notifying authority shall be established in such a way that no conflict of interest with conformity assessment bodies occurs.
2. A notifying authority shall be organised and operated so as to safeguard the objectivity and impartiality of its activities.
3. A notifying authority shall be organised in such a way that each decision relating to notification of a conformity assessment body is taken by competent persons different from those who carried out the assessment.
4. A notifying authority shall not offer or provide any activities that conformity assessment bodies perform or consultancy services on a commercial or competitive basis.

5. A notifying authority shall safeguard the confidentiality of the information it obtains.
6. A notifying authority shall have a sufficient number of competent personnel at its disposal for the proper performance of its tasks.

#### Article 25

##### Information obligation on notifying authorities

Member States shall inform the Commission of their procedures for the assessment and notification of conformity assessment bodies and the monitoring of notified bodies, and of any changes thereto.

The Commission shall make that information publicly available.



## Article 26

### Requirements relating to notified bodies

1. For the purposes of notification, a conformity assessment body shall meet the requirements laid down in paragraphs 2 to 11.
2. A conformity assessment body shall be established under national law *of a Member State* and have legal personality.
3. A conformity assessment body shall be a third-party body independent of the organisation or the radio equipment it assesses.

A body belonging to a business association or professional federation representing undertakings involved in the design, manufacturing, provision, assembly, use or maintenance of radio equipment which it assesses, may, on condition that its independence and the absence of any conflict of interest are demonstrated, be considered such a body.

4. A conformity assessment body, its top level management and the personnel responsible for carrying out the conformity assessment tasks shall not be the designer, manufacturer, supplier, installer, purchaser, owner, user or maintainer of the radio equipment which they assess, nor the ■ representative of any of those parties. This shall not preclude the use of assessed radio equipment that is necessary for the operations of the conformity assessment body or the use of such radio equipment for personal purposes.

A conformity assessment body, its top level management and the personnel responsible for carrying out the conformity assessment tasks shall not be directly involved in the design, manufacture or construction, the marketing, installation, use or maintenance of that radio equipment, or represent the parties engaged in those activities. They shall not engage in any activity that may conflict with their independence of judgement or integrity in relation to conformity assessment activities for which they are notified. This shall in particular apply to consultancy services.

Conformity assessment bodies shall ensure that the activities of their subsidiaries or subcontractors do not affect the confidentiality, objectivity or impartiality of their conformity assessment activities.

5. Conformity assessment bodies and their personnel shall carry out the conformity assessment activities with the highest degree of professional integrity and the requisite technical competence in the specific field and shall be free from all pressures and inducements, particularly financial, which might influence their judgement or the results of their conformity assessment activities, especially as regards persons or groups of persons with an interest in the results of those activities.
6. A conformity assessment body shall be capable of carrying out all the conformity assessment tasks assigned to it by *Annexes III and IV* in relation to which it has been notified, whether those tasks are carried out by the conformity assessment body itself or on its behalf and under its responsibility.

At all times and for each conformity assessment procedure and each kind or category of radio equipment in relation to which it has been notified, a conformity assessment body shall have at its disposal the necessary:

- (a) personnel with technical knowledge and sufficient and appropriate experience to perform the conformity assessment tasks;
- (b) descriptions of procedures in accordance with which conformity assessment is carried out, ensuring the transparency and the ability of reproduction of those procedures. It shall have appropriate policies and procedures in place that distinguish between tasks it carries out as a notified body and other activities;
- (c) procedures for the performance of activities which take due account of the size of an undertaking, the sector in which it operates, its structure, the degree of complexity of radio equipment technology in question and the mass or serial nature of the production process.

*A conformity assessment body* shall have the means necessary to perform the technical and administrative tasks connected with the conformity assessment activities in an appropriate manner.

7. The personnel responsible for carrying out conformity assessment *tasks* shall have the following:
  - (a) sound technical and vocational training covering all the conformity assessment activities in relation to which the conformity assessment body has been notified;
  - (b) satisfactory knowledge of the requirements of the assessments they carry out and adequate authority to carry out those assessments;
  - (c) appropriate knowledge and understanding of the essential requirements set out in Article 3, of the applicable harmonised standards and of the relevant provisions of Union harmonisation legislation and of national legislation;

(d) the ability to draw up *EU-type examination* certificates *or quality system approvals*, records and reports demonstrating that assessments have been carried out.

8. The impartiality of the conformity assessment bodies, their top level management and of the ■ personnel *responsible for carrying out the conformity assessment tasks* shall be guaranteed.

The remuneration of the top level management and ■ personnel *responsible for carrying out the conformity assessment tasks* of a conformity assessment body shall not depend on the number of assessments carried out or on the results of those assessments.

9. Conformity assessment bodies shall take out liability insurance unless liability is assumed by the State in accordance with national law, or the Member State itself is directly responsible for the conformity assessment.

10. The personnel of a conformity assessment body shall observe professional secrecy with regard to all information obtained in carrying out their tasks under *Annexes III and IV* or any provision of national law giving effect to them, except in relation to the competent authorities of the Member State in which its activities are carried out. Proprietary rights shall be protected.
  
11. Conformity assessment bodies shall participate in, or ensure that their **■** personnel *responsible for carrying out the conformity assessment tasks* are informed of, the relevant standardisation activities, the regulatory activities in the area of radio equipment and frequency planning, and the activities of the notified body coordination group established under the relevant Union harmonisation legislation and *shall* apply as general guidance the administrative decisions and documents produced as a result of the work of that group.

## Article 27

### Presumption of conformity of *notified* bodies

Where a conformity assessment body demonstrates its conformity with the criteria laid down in the relevant harmonised standards or parts thereof the references of which have been published in the *Official Journal of the European Union* it shall be presumed to comply with the requirements set out in Article 26 in so far as the applicable harmonised standards cover those requirements.

## Article 28

### Subsidiaries of and subcontracting by notified bodies

1. Where a notified body subcontracts specific tasks connected with conformity assessment or has recourse to a subsidiary, it shall ensure that the subcontractor or the subsidiary meets the requirements set out in Article 26 and shall inform the notifying authority accordingly.
2. Notified bodies shall take full responsibility for the tasks performed by subcontractors or subsidiaries wherever these are established.



3. Activities may be subcontracted or carried out by a subsidiary only with the agreement of the client.
4. Notified bodies shall keep at the disposal of the notifying authority the relevant documents concerning the assessment of the qualifications of the subcontractor or the subsidiary and the work carried out by them under *Annexes III and IV*.

#### Article 29

##### Application for notification

1. A conformity assessment body shall submit an application for notification to the notifying authority of the Member State in which it is established.

2. *The application for notification* shall be accompanied by a description of the conformity assessment activities, the conformity assessment module or modules and the radio equipment for which that body claims to be competent, as well as by an accreditation certificate, where one exists, issued by a national accreditation body attesting that the conformity assessment body fulfils the requirements laid down in Article 26.
3. Where the conformity assessment body concerned cannot provide an accreditation certificate, it shall provide the notifying authority with all the documentary evidence necessary for the verification, recognition and regular monitoring of its compliance with the requirements laid down in Article 26.

## Article 30

### Notification procedure

1. Notifying authorities may notify only conformity assessment bodies which have satisfied the requirements laid down in Article 26.
2. They shall notify the Commission and the other Member States using the electronic notification tool developed and managed by the Commission.
3. The notification shall include full details of the conformity assessment activities, the conformity assessment module or modules and **■** the radio equipment concerned and the relevant attestation of competence.

4. Where a notification is not based on an accreditation certificate as referred to in Article 29(2), the notifying authority shall provide the Commission and the other Member States with documentary evidence which attests to the conformity assessment body's competence and the arrangements in place to ensure that that body will be monitored regularly and will continue to satisfy the requirements laid down in Article 26.
5. The body concerned may perform the activities of a notified body only where no objections are raised by the Commission or the other Member States within two weeks of a notification where an accreditation certificate is used or within two months of a notification where accreditation is not used.

Only such a body shall be considered a notified body for the purposes of this Directive.

6. ***The notifying authority shall notify the Commission and the other Member States of any subsequent relevant changes to the notification.***

## Article 31

### Identification numbers and lists of notified bodies

1. The Commission shall assign an identification number to a notified body.

It shall assign a single such number even where the body is notified under several Union acts.

2. The Commission shall make publicly available the list of the bodies notified under this Directive, including the identification numbers that have been *assigned* to them and the activities for which they have been notified.

The Commission shall ensure that *the* list is kept up to date.

## Article 32

### Changes to notifications

1. Where a notifying authority has ascertained or has been informed that a notified body no longer meets the requirements laid down in Article 26, or that it is failing to fulfil its obligations, the notifying authority shall restrict, suspend or withdraw notification as appropriate, depending on the seriousness of the failure to meet those requirements or fulfil those obligations. It shall immediately inform the Commission and the other Member States accordingly.
2. In the event of restriction, suspension or withdrawal of notification, or where the notified body has ceased its activity, the notifying Member State shall take appropriate steps to ensure that the files of that body are either processed by another notified body or kept available for the responsible notifying and market surveillance authorities at their request.

## Article 33

### Challenge of the competence of notified bodies

1. The Commission shall investigate all cases where it doubts, or doubt is brought to its attention regarding, the competence of a notified body or the continued fulfilment by a notified body of the requirements and responsibilities to which it is subject.
2. The notifying Member State shall provide the Commission, on request, with all information relating to the basis for the notification or the maintenance of the competence of the *notified* body concerned.
3. The Commission shall ensure that all sensitive information obtained in the course of its investigations is treated confidentially.

4. Where the Commission ascertains that a notified body does not meet or no longer meets the requirements for its notification, it shall ***adopt an implementing act requesting*** the notifying Member State **█** to take the necessary corrective measures, including ***withdrawal of notification*** if necessary.

***That implementing act shall be adopted in accordance with the advisory procedure referred to in Article 45(2).***

#### Article 34

##### Operational obligations of notified bodies

1. Notified bodies shall carry out conformity assessments in accordance with the conformity assessment procedures provided for in Annexes III and IV.
2. Conformity assessments shall be carried out in a proportionate manner, avoiding unnecessary burdens for economic operators. Conformity assessment bodies shall perform their activities taking due account of the size of an undertaking, the sector in which it operates, its structure, the degree of complexity of the radio equipment technology in question and the mass or serial nature of the production process.



In so doing they shall nevertheless respect the degree of rigour and the level of protection required for the compliance of the radio equipment with this Directive.

3. Where a notified body finds that the essential requirements set out in Article 3 or corresponding harmonised standards or *other* technical specifications have not been met by a manufacturer, it shall require that manufacturer to take appropriate corrective measures and shall not issue *an EU-type examination certificate or a quality system approval*.
4. Where, in the course of the monitoring of conformity following the issue of *an EU-type examination certificate or a quality system approval*, a notified body finds that radio equipment no longer complies, it shall require the manufacturer to take appropriate corrective measures and shall suspend or withdraw the *EU-type examination certificate or the quality system approval* if necessary.
5. Where corrective measures are not taken or do not have the required effect, the notified body shall restrict, suspend or withdraw any *EU-type examination certificates or quality system approvals*, as appropriate.

## Article 35

### Appeal against decisions of notified bodies

Member States shall ensure that an appeal procedure against decisions of the notified bodies is available.

## Article 36

### Information obligation on notified bodies

1. Notified bodies shall inform the notifying authority of the following:
  - (a) any refusal, restriction, suspension or withdrawal of *an EU-type examination certificate or a quality system approval in accordance with the requirements of Annexes III and IV*;
  - (b) any circumstances affecting the scope of *or* conditions for notification;
  - (c) any request for information which they have received from market surveillance authorities regarding conformity assessment activities;

- (d) on request, conformity assessment activities performed within the scope of their notification and any other activity performed, including cross-border activities and subcontracting.
2. Notified bodies shall, *in accordance with the requirements of Annexes III and IV*, provide the other bodies notified under this Directive carrying out similar conformity assessment activities covering the same categories of radio equipment with relevant information on issues relating to negative and, on request, positive conformity assessment results.
3. Notified bodies shall fulfil information obligations under Annexes III and IV.

#### Article 37

#### Exchange of experience

The Commission shall provide for the organisation of exchange of experience between the Member States' national authorities responsible for notification policy.

Article 38

Coordination of notified bodies

The Commission shall ensure that appropriate coordination and cooperation between bodies notified under this Directive are put in place and properly operated in the form of a sectoral group of notified bodies.

Member States shall ensure that the bodies notified by them participate in the work of that group, directly or by means of designated representatives.

CHAPTER V

UNION MARKET SURVEILLANCE, CONTROL OF *RADIO EQUIPMENT* ENTERING THE UNION MARKET AND *UNION SAFEGUARD PROCEDURE*

Article 39

Union market surveillance and control of *radio equipment* entering the Union market

Article 15(3) and Articles 16 to 29 of Regulation (EC) No 765/2008 shall apply to radio equipment.

## Article 40

### Procedure for dealing with radio equipment presenting a risk at national level

1. Where the market surveillance authorities of one Member State **■** have sufficient **■** reason to believe that radio equipment covered by this Directive presents a risk to the health or safety of persons or to other aspects of public interest protection covered by this Directive, they shall carry out an evaluation in relation to the radio equipment concerned covering all *relevant* requirements laid down in this Directive. The relevant economic operators shall cooperate as necessary with the market surveillance authorities *for that purpose*.

Where, in the course of *the* evaluation *referred to in the first subparagraph*, the market surveillance authorities find that the radio equipment does not comply with the requirements laid down in this Directive, they shall without delay require the relevant economic operator to take all appropriate corrective *actions* to bring the radio equipment into compliance with those requirements, to withdraw the radio equipment from the market, or to recall it within a reasonable period, commensurate with the nature of the risk, as they may prescribe.

The market surveillance authorities shall inform the relevant notified body accordingly.

Article 21 of Regulation (EC) No 765/2008 shall apply to the measures referred to in the second subparagraph *of this paragraph*.

2. Where the market surveillance authorities consider that non-compliance is not restricted to their national territory, they shall inform the Commission and the other Member States of the results of the evaluation and of the actions which they have required the economic operator to take.
3. The economic operator shall ensure that all appropriate corrective action is taken in respect of all radio equipment concerned that it has made available on the market throughout the Union.

4. Where the relevant economic operator does not take adequate corrective action within the period referred to in the second subparagraph of paragraph 1, the market surveillance authorities shall take all appropriate provisional measures to prohibit or restrict *the* radio equipment being made available on their national market, to withdraw *the* radio equipment from that market or to recall it.

The market surveillance authorities shall inform the Commission and the other Member States, without delay, of those measures.

5. The information referred to in *the second subparagraph of* paragraph 4 shall include all available details, in particular the data necessary for the identification of the non-compliant radio equipment, the origin of the radio equipment, the nature of the non-compliance alleged and the risk involved, the nature and duration of the national measures taken and the arguments put forward by the relevant economic operator. In particular, the market surveillance authorities shall indicate whether the non-compliance is due to either of the following:

- (a) failure of *the* radio equipment to meet *the relevant essential* requirements *set out in Article 3; or*
  - (b) shortcomings in the harmonised standards referred to in Article 16 conferring a presumption of conformity.
6. Member States other than the Member State initiating the procedure *under this Article* shall without delay inform the Commission and the other Member States of any measures adopted and of any additional information at their disposal relating to the non-compliance of *the* radio equipment concerned, and, in the event of disagreement with the *adopted* national measure, of their objections.
7. Where, within *three months* of receipt of the information referred to in *the second subparagraph of* paragraph 4, no objection has been raised by either a Member State or the Commission in respect of a provisional measure taken by a Member State, that measure shall be deemed justified.



8. Member States shall ensure that appropriate restrictive measures, *such as withdrawal of the radio equipment from the market*, are taken in respect of the *radio equipment* concerned without delay.

#### Article 41

##### Union safeguard procedure

1. Where, on completion of the procedure set out in *Article 40(3)* and **■** (4), objections are raised against a measure taken by a Member State, or where the Commission considers a national measure to be contrary to Union legislation, the Commission shall without delay enter into consultation with the Member States and the relevant economic operator or operators and shall evaluate the national measure. On the basis of the results of that evaluation, the Commission shall *adopt an implementing act determining* whether the national measure is justified or not.

The Commission shall address its decision to all Member States and shall immediately communicate it to them and the relevant economic operator or operators.

2. If the national measure is considered justified, all Member States shall take the *necessary* measures ■ to ensure that the non-compliant radio equipment is withdrawn or recalled from their market, and shall inform the Commission accordingly. If the national measure is considered unjustified, the Member State concerned shall withdraw *that* measure.
3. Where the national measure is considered justified and the non-compliance of the radio equipment is attributed to shortcomings in the harmonised standards referred to in *point (b) of Article 40(5)* of this Directive, the Commission shall apply the procedure provided for in *Article 11 of Regulation (EU) No 1025/2012*.

## Article 42

### Compliant radio equipment which presents a risk ■

1. Where, having *carried out* an evaluation under Article 40(1), a Member State finds that although radio equipment is in compliance with this Directive, it presents a risk to the health or safety of persons or to other aspects of public interest protection covered by this Directive, it shall require the relevant economic operator to take all appropriate measures to ensure that the radio equipment concerned, when placed on the market, no longer presents that risk, to withdraw the radio equipment from the market or to recall it within a reasonable period, commensurate with the nature of the risk, as it may prescribe.
2. The economic operator shall ensure that corrective action is taken in respect of all the radio equipment concerned *that he has made available on the market throughout the Union*.

3. The Member State shall immediately inform the Commission and the other Member States. That information shall include all available details, in particular the data necessary for the identification of the radio equipment concerned, the origin and the supply chain of radio equipment, the nature of the risk involved and the nature and duration of the national measures taken.
4. The Commission shall without delay enter into consultation with the Member States and the relevant economic operator or operators and shall evaluate the national measures taken. On the basis of the results of that evaluation, the Commission shall decide *by means of implementing acts* whether the *national* measure is justified or not and, where necessary, propose appropriate measures.

*The implementing acts referred to in the first subparagraph of this paragraph shall be adopted in accordance with the examination procedure referred to in Article 45(3).*

*On duly justified imperative grounds of urgency relating to the protection of health and safety of persons, the Commission shall adopt immediately applicable implementing acts in accordance with the procedure referred to in Article 45(4).*

5. The Commission shall address its decision to all Member States and shall immediately communicate it to them and the relevant economic operator or operators.

#### Article 43

#### Formal non-compliance

1. Without prejudice to Article 40, where a Member State makes one of the following findings, it shall require the relevant economic operator to put an end to the non-compliance concerned:
  - (a) the CE marking has been affixed in violation of Article 30 of Regulation (EC) No 765/2008 or of Article 20 of this Directive;
  - (b) the CE marking has not been affixed;

- (c) *the identification number of the notified body, where the conformity assessment procedure set out in Annex IV is applied, has been affixed in violation of Article 20 or has not been affixed;*
- (d) the EU declaration of conformity has not been drawn up;
- (e) the EU declaration of conformity has not been drawn up correctly;
- (f) technical documentation is either not available or not complete;
- (g) *the information referred to in Article 10(6) or (7) or Article 12(3) is absent, false or incomplete;*
- (h) information on the intended use of radio equipment, the EU declaration of conformity *or* usage restrictions as set out in Article 10(8), (9) and (10) does not accompany the radio equipment;

- (i) requirements on identification of economic operators set out in Article 15 are not fulfilled;
  - (j) Article 5 is not complied with.
2. Where the non-compliance referred to in paragraph 1 persists, the Member State concerned shall take all appropriate measures to restrict or prohibit corresponding radio equipment being made available on the market or ensure that it is withdrawn or recalled from the market.

## CHAPTER VI

### ***DELEGATED ACTS AND IMPLEMENTING ACTS AND THE COMMITTEE***

#### Article 44

#### ***Exercise of the delegation***

1. ***The power to adopt delegated acts is conferred on the Commission subject to the conditions laid down in this Article.***

2. *The power to adopt delegated acts referred to in the second subparagraph of Article 3(3), Article 4(2) and Article 5(2) shall be conferred on the Commission for a period of five years from ...<sup>\*</sup>. The Commission shall draw up a report in respect of the delegation of power not later than nine months before the end of the five-year period. The delegation of power shall be tacitly extended for periods of an identical duration, unless the European Parliament or the Council opposes such extension not later than three months before the end of each period.*
  
3. *The delegation of power referred to in the second subparagraph of Article 3(3), Article 4(2) and Article 5(2) may be revoked at any time by the European Parliament or by the Council. A decision to revoke shall put an end to the delegation of the power specified in that decision. It shall take effect the day following the publication of the decision in the Official Journal of the European Union or at a later date specified therein. It shall not affect the validity of any delegated acts already in force.*

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\*

*OJ: please insert the date of entry into force of this Directive.*



4. *As soon as it adopts a delegated act, the Commission shall notify it simultaneously to the European Parliament and to the Council.*
5. *A delegated act adopted pursuant to the second subparagraph of Article 3(3), Article 4(2) and Article 5(2) shall enter into force only if no objection has been expressed either by the European Parliament or the Council within a period of two months of notification of that act to the European Parliament and the Council or if, before the expiry of that period, the European Parliament and the Council have both informed the Commission that they will not object. That period shall be extended by two months at the initiative of the European Parliament or of the Council.*

Article 45

*Committee procedure*

1. *The Commission shall be assisted by the Telecommunication Conformity Assessment and Market Surveillance Committee. That committee shall be a committee within the meaning of Regulation (EU) No 182/2011.*

2. *Where reference is made to this paragraph, Article 4 of Regulation (EU) No 182/2011 shall apply.*
3. *Where reference is made to this paragraph, Article 5 of Regulation (EU) No 182/2011 shall apply.*
4. *Where reference is made to this paragraph, Article 8 of Regulation (EU) No 182/2011, in conjunction with Article 5 thereof, shall apply.*
5. *The committee shall be consulted by the Commission on any matter for which consultation of sectoral experts is required by Regulation (EU) No 1025/2012 or by any other Union legislation.*

*The committee may furthermore examine any other matter concerning the application of this Directive raised either by its chair or by a representative of a Member State in accordance with its rules of procedure.*

CHAPTER VII  
FINAL AND TRANSITIONAL PROVISIONS

Article 46

Penalties

Member States shall lay down rules on penalties applicable to infringements *by economic operators* of the *provisions of national law* adopted pursuant to this Directive and shall take all measures necessary to ensure that they are enforced. *Such rules may include criminal penalties for serious infringements.*

The penalties provided for shall be effective, proportionate and dissuasive.

■

## Article 47

### Review and reporting

1. Member States shall submit to the Commission regular reports on the application of this Directive by ... \* and at least every two years thereafter. The reports shall contain a presentation of the market surveillance activities performed by the Member States and provide information on whether and to what extent compliance with the requirements of this Directive has been attained, including in particular requirements on identification of economic operators.

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\* *OJ: please insert the date: three years after the entry into force of this Directive.*

2. The Commission shall review the operation of this Directive and report thereon to the European Parliament and to the Council, by ... \* and every five years thereafter. The report shall cover progress on drawing up the relevant standards, as well as any problems that have arisen in the course of implementation. The report shall also outline the activities of the *Telecommunication Conformity Assessment and Market Surveillance* Committee, assess progress in achieving an open competitive market for radio equipment at Union level and examine how the regulatory framework for the *making available* on the market and putting into service of radio equipment should be developed in order to achieve the following:
- (a) ensure that a coherent system is achieved at Union level for all radio equipment;
  - (b) allow for convergence of the telecommunications, audiovisual and information technology sectors;

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\* *OJ: please insert the date: four years after the entry into force of this Directive.*

- (c) enable regulatory measures to be harmonised at international level;
- (d) *reach a high level of consumer protection;*
- (e) *ensure that portable radio equipment interworks with accessories, in particular with common chargers;*
- (f) *where radio equipment is fitted with an integral screen, allow the display of the required information on the integral screen.*

█

## Article 48

### Transitional provisions

Member States shall not impede, *for the aspects covered by this Directive*, the making available on the market or putting into service of radio equipment covered by *this Directive* ■ which is in conformity with *the relevant Union harmonisation legislation applicable before ...* \* and which was placed on the market before ... \*\* .

## Article 49

### Transposition

1. Member States shall adopt and publish, by ... \*\*\*, the laws, regulations and administrative provisions necessary to comply with this Directive. They shall forthwith communicate *the text of those measures* to the Commission ■ .

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\* *OJ: Please insert the date: day after two years after the entry into force of this Directive.*

\*\* *OJ: Please insert the date: day after three years after the entry into force of this Directive.*

\*\*\* *OJ: Please insert the date: two years after the entry into force of this Directive.*

They shall apply those *measures* from ...\* .

When Member States adopt those *measures*, they shall contain a reference to this Directive or be accompanied by such a reference on the occasion of their official publication. They shall also include a statement that references in existing laws, regulations and administrative provisions to the Directive repealed by this Directive shall be construed as references to this Directive. Member States shall determine how such reference is to be made and how that statement is to be formulated.

2. Member States shall communicate to the Commission the texts of the main provisions of national law which they adopt in the field covered by this Directive.

#### Article 50

#### Repeal

Directive 1999/5/EC is repealed with effect from ...\*.

References to the repealed Directive shall be construed as references to this Directive and shall be read in accordance with the correlation table in Annex VIII.

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\*

*OJ: Please insert the date: day after two years after the entry into force of this Directive.*



Article 51  
Entry into force

This Directive shall enter into force on the twentieth day following that of its publication in the *Official Journal of the European Union*.

Article 52  
Addressees

This Directive is addressed to the Member States.

*Done at ...,*

*For the European Parliament*  
*The President*

*For the Council*  
*The President*

## ANNEX I

### EQUIPMENT NOT COVERED BY THIS DIRECTIVE

1. Radio equipment used by radio amateurs within the meaning of Article 1, definition 56, of the International Telecommunications Union (ITU) **Radio Regulations**, unless the equipment is *made available on the market*.

*The following shall be regarded as not being made available on the market:*

- (a) *radio kits for assembly and use by radio amateurs;*
- (b) *radio equipment modified by and for the use of radio amateurs;*
- (c) *equipment constructed by individual radio amateurs for experimental and scientific purposes related to amateur radio.*

2. Marine equipment falling within the scope of Council Directive 96/98/EC<sup>1</sup>.

3. *Airborne* products, parts and appliances falling within the *scope* of Article 3 of Regulation (EC) No 216/2008 of the European Parliament and of the Council<sup>2</sup>.

4. *Custom-built evaluation kits destined for professionals to be used solely at research and development facilities for such purposes.*

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<sup>1</sup> *Council Directive 96/98/EC of 20 December 1996 on marine equipment* (OJ L 46, 17.2.1997, p. 25).

<sup>2</sup> *Regulation (EC) No 216/2008 of the European Parliament and of the Council of 20 February 2008 on common rules in the field of civil aviation and establishing a European Aviation Safety Agency, and repealing Council Directive 91/670/EEC, Regulation (EC) No 1592/2002 and Directive 2004/36/EC* (OJ L 79, 19.3.2008, p. 1).

## ANNEX II

### CONFORMITY ASSESSMENT MODULE A

#### INTERNAL PRODUCTION CONTROL

1. Internal production control is the conformity assessment procedure whereby the manufacturer fulfils the obligations laid down in points 2, 3 and 4 of this Annex, and ensures and declares on his sole responsibility that the radio equipment concerned satisfies the *essential* requirements set out in Article 3.


2. Technical documentation

The manufacturer shall establish the technical documentation in accordance with Article 21.

3. Manufacturing

The manufacturer shall take all measures necessary so that the manufacturing process and its monitoring ensure compliance of the manufactured radio equipment with the technical documentation referred to in point 2 and with the relevant essential requirements set out in Article 3.

4. CE marking and *EU* declaration of conformity

4.1. The manufacturer shall affix the  CE marking in accordance with Articles 19 and 20 to each item of radio equipment that satisfies the applicable requirements of this Directive.

4.2. The manufacturer shall draw up a written *EU* declaration of conformity for each radio equipment type and keep it together with the technical documentation at the disposal of the national authorities for 10 years after the radio equipment has been placed on the market. The *EU* declaration of conformity shall identify the radio equipment for which it has been drawn up.

A copy of the *EU* declaration of conformity shall be made available to the relevant authorities upon request.

5. Authorised representative

The manufacturer's obligations set out in point 4 may be fulfilled by his authorised representative, on his behalf and under his responsibility, provided that they are specified in the mandate.

## ANNEX III

### CONFORMITY ASSESSMENT MODULES B AND C

#### EU-TYPE EXAMINATION AND CONFORMITY TO TYPE BASED ON INTERNAL PRODUCTION CONTROL

When reference is made to this Annex, the conformity assessment procedure shall follow Modules B (EU-type examination) and C (Conformity to type based on internal production control) of this Annex.

#### Module B

##### EU-type examination

1. EU-type examination is the part of a conformity assessment procedure in which a notified body examines the technical design of the radio equipment and verifies and attests that the technical design of the radio equipment meets the *essential* requirements *set out in Article 3*.
2. EU-type examination shall be carried out by assessment of the adequacy of the technical design of the radio equipment through examination of the technical documentation and supporting evidence referred to in point 3, without examination of a specimen (design type).

3. The manufacturer shall lodge an application for EU-type examination with a single notified body of his choice.

The application shall include:

- (a) the name and address of the manufacturer and, if the application is lodged by the authorised representative, his name and address as well;
- (b) a written declaration that the same application has not been lodged with any other notified body;
- (c) the technical documentation. The technical documentation shall make it possible to assess the radio equipment's conformity with the applicable requirements of *this Directive* and shall include an adequate analysis and assessment of the risk(s). The technical documentation shall specify the applicable requirements and cover, as far as relevant for the assessment, the design, manufacture and operation of the radio equipment. The technical documentation shall contain, wherever applicable, the elements set out in Annex V;

- (d) the supporting evidence for the adequacy of the technical design solution. That supporting evidence shall mention any documents that have been used, in particular where the relevant harmonised standards ■ have not been applied *or have not been fully applied*. The supporting evidence shall include, where necessary, the results of tests carried out *in accordance with other relevant technical specifications* by the appropriate laboratory of the manufacturer, or by another testing laboratory on his behalf and under his responsibility.
4. The notified body shall examine the technical documentation and supporting evidence to assess the adequacy of the technical design of the radio equipment.
5. The notified body shall draw up an evaluation report that records the activities undertaken in accordance with point 4 and their outcomes. Without prejudice to its obligations as provided in point 8, the notified body shall release the content of that report, in full or in part, only with the agreement of the manufacturer.



6. Where the type meets the requirements of *this Directive* that apply to *the* radio equipment concerned, the notified body shall issue an EU-type examination certificate to the manufacturer. *That* certificate shall contain the name and address of the manufacturer, the conclusions of the examination, *the aspects of the essential requirements covered by the examination*, the conditions (if any) for its validity and the necessary data for identification of the assessed type. The *EU-type examination* certificate may have one or more annexes attached.

The EU-type examination certificate and its annexes shall contain all relevant information to allow the conformity of manufactured radio equipment with the examined type to be evaluated and to allow for in-service control.

Where the type does not satisfy the applicable requirements of *this Directive*, the notified body shall refuse to issue an EU-type examination certificate and shall inform the applicant accordingly, giving detailed reasons for its refusal.

7. The notified body shall keep itself apprised of any changes in the generally acknowledged state of the art which indicate that the approved type may no longer comply with the applicable requirements of *this Directive*, and shall determine whether such changes require further investigation. If so, the notified body shall inform the manufacturer accordingly.

The manufacturer shall inform the notified body that holds the technical documentation relating to the EU-type examination certificate of all modifications to the *approved* type that may affect the conformity of the radio equipment with the essential requirements of *this Directive* or the conditions for validity of *that* certificate. Such modifications shall require additional approval in the form of an addition to the original EU-type examination certificate.

8. Each notified body shall inform its notifying *authority* concerning the EU-type examination certificates and/or any additions thereto which it has issued or withdrawn, and shall, periodically or upon request, make available to its notifying *authority* the list of *such* certificates and/or any additions thereto refused, suspended or otherwise restricted.

Each notified body shall inform the other notified bodies concerning the EU-type examination certificates and/or any additions thereto which it has refused, withdrawn, suspended or otherwise restricted, and, upon request, concerning *such* certificates and/or additions thereto which it has issued.

Each notified body shall inform *the* Member States of EU-type examination certificates it has issued and/or additions thereto in those cases where harmonised standards the references of which have been published in the *Official Journal of the European Union* ■ have **not been applied or** not been fully applied. The Member States, the Commission and the other notified bodies may, on request, obtain a copy of the EU-type examination certificates and/or additions thereto. On request, the Member States and the Commission may obtain a copy of the technical documentation and the results of the examinations carried out by the notified body. The notified body shall keep a copy of the EU-type examination certificate, its annexes and additions, as well as the technical file including the documentation submitted by the manufacturer for ■ 10 years after the radio equipment has been assessed or until the expiry of the validity of *that* certificate.

9. The manufacturer shall keep a copy of the EU-type examination certificate, its annexes and additions together with the technical documentation at the disposal of the national authorities for 10 years after the radio equipment has been placed on the market.

10. The manufacturer's authorised representative may lodge the application referred to in point 3 and fulfil the obligations set out in points 7 and 9, provided that they are specified in the mandate.

## Module C

### Conformity to type based on internal production control

1. Conformity to type based on internal production control is the part of a conformity assessment procedure whereby the manufacturer fulfils the obligations laid down in points 2 and 3, and ensures and declares that the radio equipment concerned is in conformity with the type described in the EU-type examination certificate and satisfy the requirements of *this Directive* that apply to it.

#### 2. Manufacturing

The manufacturer shall take all measures necessary so that the manufacturing process and its monitoring ensure conformity of the manufactured radio equipment with the approved type described in the EU-type examination certificate and with the requirements of *this Directive* that apply to it.

#### 3. CE marking and *EU* declaration of conformity

3.1. The manufacturer shall affix the CE marking in accordance with Articles 19 and 20 to each item of radio equipment that is in conformity with the type described in the EU-type examination certificate and satisfies the applicable requirements of *this Directive*.

3.2. The manufacturer shall draw up a written *EU* declaration of conformity for each radio equipment type and keep it at the disposal of the national authorities for 10 years after the radio equipment has been placed on the market. The EU declaration of conformity shall identify the radio equipment type for which it has been drawn up.

A copy of the EU declaration of conformity shall be made available to the relevant authorities upon request.

#### 4. Authorised representative

The manufacturer's obligations set out in point 3 may be fulfilled by his authorised representative, on his behalf and under his responsibility, provided that they are specified in the mandate.

## ANNEX IV

### CONFORMITY ASSESSMENT MODULE H

#### **CONFORMITY BASED ON FULL QUALITY ASSURANCE**

1. Conformity based on full quality assurance is the conformity assessment procedure whereby the manufacturer fulfils the obligations laid down in points 2 and 5, and ensures and declares on his sole responsibility that the radio equipment concerned satisfies the requirements of *this Directive* that apply to it.

2. Manufacturing

The manufacturer shall operate an approved quality system for design, manufacture, final **radio equipment** inspection and testing of the radio equipment concerned as specified in point 3 and shall be subject to surveillance as specified in point 4.

### 3. Quality system

3.1. The manufacturer shall lodge an application for assessment of his quality system with the notified body of his choice, for the radio equipment concerned.

The application shall include:

- (a) the name and address of the manufacturer and, if the application is lodged by the authorised representative, his name and address as well;
- (b) the technical documentation for **■** each radio equipment type intended to be manufactured. The technical documentation shall contain, wherever applicable, the elements set out in Annex V;
- (c) the documentation concerning the quality system; and
- (d) a written declaration that the same application has not been lodged with any other notified body.

3.2. The quality system shall ensure compliance of the radio equipment with the requirements of *this Directive* that apply to it.

All the elements, requirements and provisions adopted by the manufacturer shall be documented in a systematic and orderly manner in the form of written policies, procedures and instructions. That quality system documentation shall permit a consistent interpretation of the quality programmes, plans, manuals and records.

It shall, in particular, contain an adequate description of:

- (a) the quality objectives and the organisational structure, responsibilities and powers of the management with regard to design and product quality;
- (b) the technical design specifications, including standards, that will be applied and, where the relevant harmonised standards ■ will not be applied in full, the means that will be used to ensure that the essential requirements of this Directive that apply to the radio equipment will be met;



- (c) the design control and design verification techniques, processes and systematic actions that will be used when designing radio equipment pertaining to the radio equipment *type* covered;
- (d) the corresponding manufacturing, quality control and quality assurance techniques, processes and systematic actions that will be used;
- (e) the examinations and tests that will be carried out before, during and after manufacture, and the frequency with which they will be carried out;
- (f) the quality records, such as inspection reports and test data, calibration data, ■ reports *concerning the qualifications of* the personnel ■ , etc.;
- (g) the means of monitoring the achievement of the required design and product quality and the effective operation of the quality system.

3.3. The notified body shall assess the quality system to determine whether it satisfies the requirements referred to in point 3.2 **■** .

It shall presume conformity with those requirements in respect of the elements of the quality system that comply with the corresponding specifications of **■** the relevant harmonised standard **■** .

In addition to experience in quality management systems, the auditing team shall have at least one member experienced as an assessor in the relevant radio equipment field and radio equipment technology concerned, and knowledge of the applicable requirements of *this Directive*. The audit shall include an assessment visit to the manufacturer's premises. The auditing team shall review the technical documentation referred to in point 3.1(b) to verify the manufacturer's ability to identify the applicable requirements of *this Directive* and to carry out the necessary examinations with a view to ensuring compliance of the radio equipment with those requirements.

The manufacturer or his authorised representative shall be notified of the decision.

The notification shall contain the conclusions of the audit and the reasoned assessment decision.

3.4. The manufacturer shall undertake to fulfil the obligations arising out of the quality system as approved and to maintain it so that it remains adequate and efficient.

3.5. The manufacturer shall keep the notified body that has approved the quality system informed of any intended change to the quality system.

The notified body shall evaluate any proposed changes and decide whether the modified quality system will continue to satisfy the requirements referred to in point 3.2 or whether a reassessment is necessary.

It shall notify the manufacturer of its decision. The notification shall contain the conclusions of the examination and the reasoned assessment decision.

4. Surveillance under the responsibility of the notified body

4.1. The purpose of surveillance is to make sure that the manufacturer duly fulfils the obligations arising out of the approved quality system.

4.2. The manufacturer shall, for assessment purposes, allow the notified body access to the design, manufacture, inspection, testing and storage sites, and shall provide it with all necessary information, in particular:

- (a) the quality system documentation;
- (b) the quality records as provided for by the design part of the quality system, such as results of analyses, calculations, tests, etc.;
- (c) the quality records as provided for by the manufacturing part of the quality system, such as inspection reports and test data, calibration data, reports concerning the qualifications of the personnel, etc.

4.3. The notified body shall carry out periodic audits to make sure that the manufacturer maintains and applies the quality system and shall provide the manufacturer with an audit report.

4.4. In addition, the notified body may pay unexpected visits to the manufacturer. During such visits, the notified body may, if necessary, carry out radio equipment tests, or have them carried out, in order to check the proper functioning of the quality system. It shall provide the manufacturer with a visit report and, if tests have been carried out, with a test report.

5. CE marking and *EU* declaration of conformity

5.1. The manufacturer shall affix the CE marking in accordance with Articles 19 and 20 and, under the responsibility of the notified body referred to in point 3.1, the latter's identification number to each item of radio equipment that satisfies the applicable requirements set out in Article 3.

5.2. The manufacturer shall draw up a written EU declaration of conformity for each radio equipment type and keep it at the disposal of the national authorities for 10 years after the radio equipment has been placed on the market. The EU declaration of conformity shall identify the radio equipment type for which it has been drawn up.

A copy of the EU declaration of conformity shall be made available to the relevant authorities upon request.

6. The manufacturer shall, for a period ending ■ 10 years after the radio equipment has been placed on the market, keep at the disposal of the national authorities:

- (a) the technical documentation referred to in point 3.1;
- (b) the documentation concerning the quality system referred to in point 3.1;
- (c) the change referred to in point 3.5, as approved;
- (d) the decisions and reports of the notified body referred to in points 3.5, 4.3 and 4.4.

7. Each notified body shall inform its notifying *authority* of quality system approvals issued or withdrawn, and shall, periodically or upon request, make available to its notifying *authority* the list of quality system approvals refused, suspended or otherwise restricted.

Each notified body shall inform the other notified bodies of quality system approvals which it has refused, suspended or withdrawn, and, upon request, of quality system approvals which it has issued.

8. Authorised representative

The manufacturer's obligations set out in points 3.1, 3.5, 5 and 6 may be fulfilled by his authorised representative, on his behalf and under his responsibility, provided that they are specified in the mandate.

## ANNEX V

### Contents of technical documentation

The technical documentation shall, wherever applicable, contain at least the following elements:

- (a) a general description of the radio equipment including:
  - (i) photographs or illustrations showing external features, marking and internal layout;
  - (ii) versions of software or firmware affecting compliance with essential requirements;
  - (iii) user information and installation instructions;
- (b) conceptual design and manufacturing drawings and schemes of components, sub-assemblies, circuits and other relevant similar elements;
- (c) descriptions and explanations necessary for the understanding of those drawings and schemes and the operation of the radio equipment;

- (d) a list of the harmonised standards ***applied in full or in part*** the references of which have been published in the *Official Journal of the European Union*, ■ and, ***where those harmonised standards have not been applied***, descriptions of the solutions adopted to meet the essential requirements ***set out*** in Article 3, ***including a list of other relevant technical specifications applied***. In the event of partly applied harmonised standards, the technical documentation shall specify the parts which have been applied;
- (e) copy of the EU declaration of conformity;
- (f) where the conformity assessment module in Annex III has been applied, copy of the EU-type examination certificate and its annexes as delivered by the notified body involved;
- (g) results of design calculations made, examinations carried out, and other relevant similar elements;
- (h) test reports;
- (i) ***an explanation of the compliance with the requirement of Article 10(2) and of the inclusion or not of information on the packaging in accordance with Article 10(10)***.



## ANNEX VI

### *EU declaration of conformity (No XXX)<sup>1</sup>*

1. ***Radio equipment (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 of the manufacturer.
4. Object of the declaration (identification of the radio equipment allowing traceability; it may include a ***colour image of sufficient clarity where necessary for the identification of the radio equipment***):
5. The object of the declaration described above is in conformity with the relevant **█** Union harmonisation legislation:  
  
Directive 2014/.../EU\*  
  
Other **█** Union harmonisation legislation where applicable

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<sup>1</sup> ***It is optional for the manufacturer to assign a number to the EU declaration of conformity.***

\* OJ: Please insert the number of this Directive.

6. References to the relevant harmonised standards used or references to the *other technical* specifications in relation to which conformity is declared. References must be listed with their identification number and version and, where applicable, date of issue:

7. Where applicable, the notified body ... (name, number) ... performed ... (description of intervention) ... and issued the EU-type examination certificate: ...

8. **■**

Where applicable, description of accessories and components, including software, which allow the radio equipment to operate as intended and covered by the EU declaration of conformity

9. ***Additional information:***

Signed for and on behalf of: ...

(place and date of issue):

(name, function) (signature):

## ANNEX VII

### Simplified *EU* declaration of conformity

The simplified EU declaration of conformity referred to in Article 10(9) shall be provided as follows:

Hereby, [Name of manufacturer] declares that the radio equipment type [designation of type of radio equipment] is in compliance with Directive 2014/.../EU\*.

***The full text of the EU declaration of conformity is available at the following internet address:***

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\* OJ: Please insert the number of this Directive.

## ANNEX VIII

### Correlation table

Directive 1999/5/EC	This Directive
Article 1	Article 1
Article 2	Article 2
Article 3(1) and (2)	Article 3(1) and (2)
Article 3(3) and Article 15a	Article 3(3), with the exception of Article 3(3)(i), and Article 44
Article 4(1) and Articles 13 to 15	Articles 8 and 45
Article 4(2)	–
Article 5(1)	Article 16
Article 5(2) and (3)	–
Article 6(1)	Article 6
Article 6(2)	–
Article 6(3)	Article 10(8), (9) and (10)
Article 6(4)	–
Article 7(1) and (2)	Article 7
Article 7(3), (4) and (5)	–
Article 8(1) and (2)	Article 9
Article 8(3)	–

Article 9	Articles 39 to 43
Article 10	Article 17
Article 11	Articles 22 to 38
Article 12	Articles 19 and 20 and Article 10(6) and (7)
Article 16	–
Article 17	Article 47
Article 18	Article 48
Article 19	Article 49
Article 20	Article 50
Article 21	Article 51
Article 22	Article 52
Annex I	Annex I
Annex II	Annex II
Annex III	–
Annex IV	Annex III
Annex V	Annex IV
Annex VI	Article 26
Annex VII(1) to (4)	Articles 19 and 20
Annex VII(5)	Article 10(10)

## ANNEX TO THE LEGISLATIVE RESOLUTION

### STATEMENT OF THE EUROPEAN PARLIAMENT

The European Parliament considers that only when and insofar as implementing acts in the sense of Regulation (EU) No 182/2011 are discussed in meetings of committees, can the latter be considered as "comitology committees" within the meaning of Annex I to the Framework Agreement on the relations between the European Parliament and the European Commission. Meetings of committees thus fall within the scope of point 15 of the Framework Agreement when and insofar as other issues are discussed.