





R&TTE directive (1999/5/CE)

European Union Compliances

- The need for CE Marking
- New Approach Directives
 - ✓ Eliminate differences in laws therefore remove technical barriers to trade
 - ✓ Prescribe the Essential requirements for Health, Safety, ...
 - Member states transpose directives and harmonized standards into their national requirements
 - Third party intervention is not mandatory
 - Manufacturer Self conformity Declaration
 - √ Voluntary Use of Standards
 - CE Label as the indication of compliance
 - CE marking process



Essential requirements labs

- the requirements that products must meet to be put on the market.
- They are mandatory.
- They define the results to be attained, or the risks to be dealt with, but do not specify the technical solutions for doing so;
- suppliers are free to choose how the requirements are to be met.



Typical products



Typical products, which are covered by the R&TTE Directive, are:

- Radio terminals: GSM handsets.
- Other radio equipment: GSM base stations, car-door openers and other short range radio devices.
- Fixed network terminal equipment: normal analogue telephones, XDSL terminals, cable and PC modems.

No more applicable in new Radio directive



Manufacturers declaration of Conformity



- Introduction of manufacturers' declaration of conformity
- Assessment of the conformity of a product with the requirements of the Directive is a responsibility for the manufacturer.



No need to obtain an approval certificate from an official body after having passed tests in a legally recognized laboratory.



Minimum Requirements la

- Terminal access requirements have been removed: fixed network terminal equipment therefore only needs to comply with
 - Health and Safety requirements:
 - Health: as per EMF recommandation 1999/519/CE
 - Safety: as per Directive 2006/95/CE (LVD) but with the lower limit removed. (Article 3.1)
 - ✓ EMC requirements: as per Directive 2004/108/CE (Article 3.1)
 - Radio equipment needs to effectively use the spectrum and not cause harmful interference. (Article 3.2)



Example: Bluetooth & Wifi labs

Applied standards

- Radio: EN 300 328 + ERC 70-03 recommandation
- EMC: EN 301 489-17 & EN 301 489-01
- **Safety**: EN 60950
- Health: EN 50364



Equipment Class Identifiers labs

Class 1:

- Radio equipment and telecommunications terminal equipment which can be placed on the market and be put into service without restrictions.
- This class will be referred to as "Class 1". An Equipment Class Identifier is not defined for this class of equipment.



Equipment Class Identifiers abs

Class 2:

- R&TTE equipment for which Member States apply restrictions on the putting into service
- or for which Member States apply restrictions on the placing on the market.
- This class will be referred to as "Class 2".
 The following Equipment Class Identifier or "alert sign" is defined for equipment within

this class:



No more equipment class 2 notifications to national FSA in new Radio directive 10



Putting Into Market



- Comply to essential requirements
- Respect of the national requirements





Radio Equipments RE directive (2014/53/EU)

New EU Legislation in 2014 5

- Electromagnetic Compatibility (EMC) Directive 2014/30/EU
 - OJ L96 29 March 2014
 - Replaces 2004/108/EC
- Low Voltage Directive 2014/35/EU
 - OJ L96 29 March 2014
 - Replaces 2006/95/EC
- Radio Equipment Directive 2014/53/EU
 - OJ L153 22 May 2014
 - Replaces Radio & Telecommunication Terminal Equipment Directive (RTTED) 1999/5/EC



2014/53/EU: New elements

- No need to cover wired terminal equipment, concentrate on radio equipment
- Provisions for Universal Charger
- Increased emphasis on efficient use of spectrum, in particular by improving radio receiver requirements
- Clear scope (but still some boundaries re: purpose of spectrum use)
- Improved provisions for market surveillance & enforcement (in particular between Member States)
 - Simplified marking requirements
 - Product registration can be introduced in cases of extensive noncompliance
 - Clear link with Radio Spectrum Decision
 - New provisions for software-defined radio



major changes (1)



- RED only applies to wireless/radio products; wired Telecom Terminal Equipment (TTE) is not covered anymore
- RED scope includes radio communication and also radio determination (RFID, radar, movement detect, etc.) equipment radio equipment not for communication or determination is not within the scope of the RED
- Broadcast receivers now fall into the scope (actually they weren't in R&TTE)



major changes (2)



- RX only (like GPS) devices remain in scope
- No lower limit of the covered frequency range (for R&TTE lower limit was 9 kHz), upper limit remains at 3000GHz
- Safety requirements now explicitly apply also for animal related equipment (was in R&TTE but not clear to many readers)
- Evaluation kits are now excluded (no approval required under RED)



major changes (3)



- RED requires common / universal chargers
- No more equipment class 2 notifications to national FSA
- No more class 2 labelling (Alert Sign)
- No more Notified Body number with CE mark for single product approvals
- (Notified Body number only applies if the Quality System of the manufacturer was assessed against RED requirements (Full Quality Assurance – Module H))
- No more CE mark in the user manual
- Notified Body Opinion will be replaced by "Type Examination Certificate"



specific requirements



- The new directive also introduces some new specific requirements:
- ensure that software can only be used with radio equipment after the compliance of that particular combination of software and the radio equipment has been demonstrated;
- the Commission will have the possibility to require that mobile phones and other portable devices are compatible with a common charger.



EQUIPMENT NOT COVERED BY THIS DIRECTIVE (1)



- 1. Radio equipment used by radio amateurs, unless the equipment is made available on the market:
 - Unless made available on the market
- 2. Marine equipment.
- 3. Airborne products, parts and appliances.
- 4. Custom-built evaluation kits destined for professionals to be used solely at research and development facilities for such purposes.



Essential requirements for all requirements for all requirement

- Art. 3.1(a): Safety requirements as LVD, but with no lower voltage limit
- Art 3.1(b): EMC requirements as EMCD
- Art 3.2: <u>effective</u> and <u>efficient</u> use of radio spectrum to avoid harmful interference
 - See recitals (10) & (11): aim to increase the resilience of receivers to ensure efficient use of spectrum in adjacent bands.



Art 3.3: Requirements to be labs

- a) Interworking with accessories, in particular common chargers
- b) Interworking via networks
- c) May be connected to interfaces of the appropriate type
- d) Shall not harm the network or misuse network resources
- e) Safeguards personal data & privacy
- f) Protection from fraud
- g) Access to emergency services
- h) Facilitates use by a person with a disability
- i) Can only load compliant software

Commission may invoke the above requirements for certain equipment where via a <u>delegated act</u>



Use of radio frequencies: notified interfaces



- Member states no longer required to notify interfaces which:
- a) Are covered by a Commission Decision under 676/2002/EC (Radio Spectrum Decision), or
- b) Correspond to a class of equivalent interfaces that can be used anywhere in the Union (specified via implementing act)
- Manufacturers no longer required to inform Member State before placing on the market equipment that uses nonharmonised spectrum
 - Manufacturer to check in EFIS if frequencies are available
 - Frequency & powerto be included in user instructions

No 'Alert symbol':∠!\



Access to market



- Art 7: "Member States shall allow the putting into service and use of radio equipment if it complies with this Directive ..."
- Art 16: "Radio equipment which is in conformity with harmonised standards ... shall be presumed to be in conformity with the essential requirements"
- Art 17.3 allows the manufacturer to self-declare conformity ("Internal production control") if he has applied harmonised standards.
- Alternatives are available: "EU-type examination" or "conformity based on full quality assurance" both require use of a Notified Body
- No specific provisions for "essential radio test suites"

