Status: Point in time view as at 31/12/2020.

Changes to legislation: The Electromagnetic Compatibility Regulations 2016, SCHEDULE 6 is up to date with all changes known to be in force on or before 22 July 2024. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details)

SCHEDULES

SCHEDULE 6

Regulation 50

Operational obligations of [F1notified][F1approved] bodies

Textual Amendments

- F1 Word in Sch. 6 heading substituted (E.W.S.) (31.12.2020) by The Product Safety and Metrology etc. (Amendment etc.) (EU Exit) Regulations 2019 (S.I. 2019/696), reg. 1, Sch. 20 para. 40(a) (with Sch. 20 para. 33) (as amended by S.I. 2020/676, regs. 1(1), 2); 2020 c. 1, Sch. 5 para. 1(1)
- 1. [F2An approved] body must carry out conformity assessments in accordance with the relevant conformity assessment procedures.
- **2.** [F3An approved] body must carry out conformity assessments in a proportionate manner, avoiding unnecessary burdens for economic operators.
- **3.** Conformity assessment bodies must 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 apparatus technology in question and the mass or serial nature of the production process.
- **4.** Conformity assessment bodies must respect the degree of rigour and level of protection required for the compliance of the apparatus with these Regulations.
- **5.** Where [F4an approved] body finds that the essential requirements or the corresponding [F5designated] standards or other technical specifications have not been met by a manufacturer, it must require that manufacturer to take appropriate corrective measures and must not issue a certificate.
- **6.** Where, in the course of the monitoring of the conformity of apparatus following the issue of a certificate, [F6an approved] body finds that apparatus is no longer in conformity, it must require the manufacturer to take appropriate corrective measures and must suspend or withdraw the certificate if necessary.
- 7. Where corrective measures are not taken or do not have the required corrective effect, the [F7approved] body must restrict, suspend or withdraw any certificate as appropriate.
 - 8. Paragraph 9 applies where [F8 an approved] body is minded to—
 - (a) refuse to issue a certificate: or
 - (b) restrict, suspend or withdraw a certificate.
 - 9. Where this paragraph applies, the [F9 approved] body must—
 - (a) give the person applying for the certificate, or the person to whom the certificate was given, a notice in writing giving reasons and specifying the date on which the refusal, restriction, suspension or withdrawal is intended to take effect;
 - (b) give the person applying for the certificate, or the person to whom the certificate was given, an opportunity to make representations within a reasonable period from the date of the notice; and

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- (c) take account of any such representations before taking its decision.
- 10. [F10 An approved] body must inform the Secretary of State of—
 - (a) any refusal, restriction, suspension or withdrawal of a certificate;
 - (b) any circumstances affecting the scope of, or conditions for, notification under regulation 44 (notification);
 - (c) any request for information which it has received from the market surveillance authority regarding conformity assessment activities; and
 - (d) on request, conformity assessment activities performed within the scope of its notifications under regulation 44 and any other activity performed, including cross-border activities and subcontracting.
- 11. [FIIAn approved] body must make provision in its contracts with its clients enabling such clients to appeal against a decision—
 - (a) to refuse to issue a certificate; or
 - (b) to restrict, suspend or withdraw a certificate.
- 12. [F12An approved] body must provide other bodies [F13approved under these Regulations] carrying out similar conformity assessment activities covering the same type of apparatus with relevant information on issues relating to negative and, on request, positive conformity assessment results.
- 13. [F14An approved] body must participate in the work of [F15any approved] body coordination group established [F16by the Secretary of State], directly or by means of its designated representatives.

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