

## SCHEDULE 4

Regulation 5(2)(b), 6(2)(c)

### EC TYPE-EXAMINATION

1. This Schedule describes that part of the procedure by which a notified body ascertains and attests that an example, representative of the production envisaged, meets the relevant provisions of these Regulations.

2.—(1) The application for EC type-examination shall be lodged by the manufacturer or his authorised representative established within the Community with a notified body of his choice.

(2) The application shall include—

- (a) the name and address of the manufacturer and, if the application is lodged by the authorised representative, the name and address in addition,
- (b) a written declaration that the same application has not been lodged with any other notified body,
- (c) the technical documents as described in paragraph 3.

(3) The applicant shall place at the disposal of the notified body an example representative of the production envisaged, hereinafter called 'type'. The notified body may request further examples if needed for carrying out the test programme.

3. The technical documents shall enable the conformity of the product with the requirements of these Regulations to be assessed. They shall, as far as is relevant for such assessment, cover the design, manufacture and operation of the product and contain as far as is relevant for assessment —

- (a) a general type-description,
- (b) conceptual design and manufacturing drawings and diagrams of components, sub-assemblies, circuits, etc.,
- (c) descriptions and explanations necessary for the understanding of the drawings and diagrams and the operation of the product,
- (d) a list of the harmonised standards applied in full or in part, and descriptions of the solutions adopted to meet the efficiency requirements where the harmonised standards have not been applied,
- (e) results of design calculations made, examinations carried out, etc.,
- (f) test reports.

4. The notified body shall—

- (a) examine the technical documents, verify that the type has been manufactured in conformity with those documents and identify the elements which have been designed in accordance with the relevant provisions of the harmonised standards as well as the components which have been designed without applying the relevant provisions of those standards;
- (b) perform or have performed the appropriate examinations and necessary tests to check whether, where the harmonised standards have not been applied, the solutions adopted by the manufacturer meet the essential requirements;
- (c) perform or have performed the appropriate examinations and necessary tests to check whether, where the manufacturer has chosen to apply the relevant standards, these have actually been applied;
- (d) agree with the applicant the location where the examinations and necessary tests are to be carried out.

**Status:** This is the original version (as it was originally made). This item of legislation is currently only available in its original format.

**5.—(1)** Where the type meets the relevant provisions of these Regulations, the notified body shall issue an EC type-examination certificate to the applicant. The certificate shall contain the name and address of the manufacturer, the conclusion of the examination and necessary data for identification of the approved type.

(2) A list of the relevant parts of the technical documents shall be annexed to the certificate and a copy kept by the notified body.

(3) If the manufacturer or his authorised representative established in the Community is refused a type certificate, the notified body shall provide detailed reasons for such refusal.

(4) The notified body shall have in place a procedure for considering appeals against refusal of a type certificate given by it, and give details of that procedure to any person refused a type certificate.

**6.** The applicant shall inform the notified body which holds the technical documents concerning the EC type-examination certificate of all modifications to the approved appliance which must receive additional approval where such changes may affect the conformity with the essential requirements or the prescribed conditions for use of the product. This additional approval shall be given in the form of an addition to the original EC type-examination certificate.

**7.** Each notified body shall communicate to the other notified bodies the relevant information concerning the EC type-examination certificates and additions issued and withdrawn.

**8.** The other notified bodies may receive copies of the EC type-examination certificates and their additions. The Annexes to the certificates must be kept at the disposal of the other notified bodies.