Council Directive 2008/73/EC of 15 July 2008 simplifying procedures of listing and publishing information in the veterinary and zootechnical fields and amending Directives 64/432/EEC, 77/504/EEC, 88/407/EEC, 88/661/EEC, 89/361/EEC, 89/556/EEC, 90/426/EEC, 90/427/EEC, 90/428/EEC, 90/429/EEC, 90/539/EEC, 91/68/EEC, 91/496/EEC, 92/35/EEC, 92/65/EEC, 92/66/EEC, 92/119/EEC, 94/28/EC, 2000/75/EC, Decision 2000/258/EC and Directives 2001/89/EC, 2002/60/EC and 2005/94/EC (Text with EEA relevance)

Article 15

Amendments to Directive 92/65/EEC

Directive 92/65/EEC is hereby amended as follows:

1. Article 11 shall be replaced by the following:

Article 11

1 The Member States shall ensure that, without prejudice to the decisions to be taken in implementation of Articles 21 and 23, only semen, ova and embryos meeting the conditions laid down in paragraphs 2, 3, 4 and 5 are the subject of trade.

2 Semen of the ovine, caprine and equine species must, without prejudice to any criteria to be complied with for the entry of equids in stud books for certain specific breeds:

— have been collected, processed and stored with a view to artificial insemination in a centre approved from the health point of view in accordance with Annex D(I), or, in the case of ovine and caprine animals by way of derogation from the above, in a holding satisfying the requirements of Directive 91/68/EEC,

— have been collected from animals meeting the conditions laid down in Annex D(II),

— have been collected, processed, preserved, stored and transported in accordance with Annex D(III),

— have been accompanied during transport to another Member State by a health certificate corresponding to a specimen to be determined in accordance with the procedure referred to in Article 26.

3 Ova and embryos of the ovine, caprine, equine and porcine species must:

— have been removed from donor females meeting the conditions laid down in Annex D(IV) by a collection team or have been produced by a production team approved by the competent authority of the Member State and satisfying the conditions to be established in Annex D(I) in accordance with the procedure referred to in Article 26,

— have been collected, processed and preserved in an appropriate laboratory, stored and transported in accordance with Annex D(III),

— be accompanied during transport to another Member State by a health certificate corresponding to a specimen to be determined in accordance with the procedure referred to in Article 26.

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Semen used for the insemination of donor females must comply with the provisions of paragraph 2 in the case of sheep, goats and equids and with the provisions of Directive 90/429/EEC for swine.

Any additional guarantees may be determined in accordance with the procedure referred to in Article 26.

4 The approved centres referred to in the first indent of paragraph 2 and the approved teams referred to in the first indent of paragraph 3 shall be registered by the competent authority of the Member State concerned, each centre and team being given a veterinary registration number.

Each Member State shall draw up and keep up to date a list of those approved centres and teams and their veterinary registration numbers and shall make it available to the other Member States and to the public.

Detailed rules for the uniform application of this paragraph may be adopted in accordance with the procedure referred to in Article 26.

5 The animal health requirements and the specimen health certificates applicable to semen, ova and embryos of species not mentioned in paragraphs 2 and 3 shall be established in accordance with the procedure referred to in Article 26.

Pending the establishment of animal health requirements and specimen health certificates for trade in such semen, ova and embryos, national rules shall continue to apply.;

2. in Article 13(2), point (d) shall be replaced by the following:

(d) All approved bodies, institutes and centres shall be registered and issued with an approval number by the competent authority.

Each Member State shall draw up and keep up to date a list of approved bodies, institutes and centres and their approval numbers and shall make it available to the other Member States and to the public.

Detailed rules for the uniform application of this point may be adopted in accordance with the procedure referred to in Article 26.;

3. in Article 17, paragraph 2 and 3 shall be replaced by the following:

2. Only animals, semen, ova and embryos referred to in Article 1 which satisfy the following requirements may be imported into the Community:

a they must come from a third country on a list to be drawn up in accordance with paragraph 3(a);

b they must be accompanied by the health certificate corresponding to a specimen to be drawn up in accordance with the procedure referred to in Article 26, signed by the competent authority of the exporting country and certifying that,

(i) the animals

— meet the additional conditions or offer the equivalent guarantees referred to in paragraph 4, and

— come from approved centres, bodies, institutes offering guarantees at least equivalent to those in Annex C;

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(ii) semen, ova and embryos come from approved collection and storage centres or collection and production teams offering guarantees at least equivalent to those to be established in Annex D(I) in accordance with the procedure referred to in Article 26.

Pending the establishment of lists of third countries, approved establishments listed in point (b), animal health requirements and specimen health certificates as referred to in paragraphs (a) and (b), national rules shall continue to apply provided they are not more favourable than those laid down in Chapter II.

3 The following shall be established:

a in accordance with the procedure referred to in Article 26, a list of third countries or parts of third countries able to provide Member States and the Commission with guarantees equivalent to those provided for in Chapter II in relation to animals, semen, ova and embryos;

b in accordance with this point, a list of approved centres or teams as referred to in the first indent of paragraph 2 of Article 11 and the first indent of paragraph 3 of that article situated in one of the third countries appearing on the list referred to in point (a) of this paragraph and for which the competent authority is able to give the guarantees provided for in Article 11(2) and (3).

The list of approved centres and teams referred to in the first subparagraph and their veterinary registration numbers shall be communicated to the Commission.

The approval of centres or teams must be immediately suspended or withdrawn by the competent authority of the third country where it no longer complies with the conditions referred to in Article 11(2) and (3) and the Commission must be immediately informed thereof.

The Commission shall provide the Member States with any new and updated lists that it receives from the competent authority of the third country in accordance with the second and third subparagraphs and shall make them available to the public for information purposes.

Detailed rules for the uniform application of this point may be adopted in accordance with the procedure referred to in Article 26;

c in accordance with the procedure referred to in Article 26, the specific animal health requirements, in particular for the protection of the Community from certain exotic diseases, or guarantees equivalent to those provided for in this Directive.

The specific requirements and equivalent guarantees established for third countries may not be more favourable than those provided for in Chapter II.;

4. in Article 20, the first paragraph shall be replaced by the following:

The rules laid down in Directive 97/78/EC shall apply in particular to the organisation of, and follow-up to the checks to be carried out by the Member States and the safeguard measures to be applied in accordance with the procedure referred to in Article 22 of that Directive..