2. The following Article is inserted: 'Article 9a

Undertakings selling and/or distributing the primary raw materials used in the manufacture of substances having a thyrostatic, oestrogenic, androgenic or gestagenic action, and of beta-agonists, shall be required to keep registers detailing, in chronological order, quantities produced or acquired and those sold

- or used for the production of pharmaceutical products or veterinary medicines.'
- 3. The first subparagraph of Article 15 is replaced by the following:

'This Regulation shall enter into force on the day of its publication in the Official Journal of the European Communities.'

Amended proposal for a Council Regulation (EC) on measures to monitor certain substances and residues thereof in live animals and animal products

(94/C 222/09)

(Text with EEA relevance)

COM(94) 294 final

(Submitted by the Commission pursuant to Article 189 a (2) of the EC Treaty on 8 July 1994)

On 14 October 1993 the Comission presented the above proposal to the Council (1). As a result of the opinion delivered by Parliament at its sitting on 19 April 1994 the original proposal is amended as follows:

1. The following recitals are inserted after the fifth recital:

Whereas in its resolution of 26 May 1993 on the Commission communication to the Council and Parliament on inspections for residues in meat (hormones, beta-agonists and other substances) (4a), the European Parliament expressed the view that systems of self-regulation by producer groups can yield very good results in combating the illegal use of growth promoters; whereas it is important for consumers that self-regulation systems of this kind adequately guarantee the absence of hormones and whereas a generic European approach is desirable to safeguard and support self-regulation systems;

Whereas producer groups should be assisted in developing self-regulation systems to ensure that their meat is hormone-free (in accordance with the Commission communication of 21 April 1993 to the Council and European Parliament on inspections for residues in meat);

- (4a) OJ No C 176, 28. 6. 1993, p. 63.'
- 2. Article 16 (7) is replaced by the following:
 - '7. Each year the Commission shall inform the Member States meeting within the Standing

Veterinary Committee and the European Parliament about the implementation of national plans and developments in the situation in the various regions of the European Union.'

3. The first subparagraph of Article 26 (2) is replaced by the following:

Where a Member State considers that, in another Member State, the controls provided for in this Regulation are not being, or have ceased to be, carried out, it shall inform the competent central authority of that Member State accordingly. Following an investigation in accordance with Article 22 (2), in respect of which the financial provisions in the first subparagraph of Article 24 (1) shall not apply, that authority shall take all necessary measures and shall, at the earliest opportunity, notify the competent central authority of the first Member State of the decisions taken and the reasons for those decisions.'

- 4. Article 28 (1) is replaced by the following:
 - 1. Any failure to cooperate with the competent authority and any obstruction of slaughterhouse personnel or the slaughterhouse supervisor or, in the case of private undertakings, of the slaughterhouse owner or owners, and of the owner of the animals or the person having charge of them, during inspection and sampling as required for the implementation of national plans for monitoring residues and during inquiries and checks provided for in this Regulation

⁽¹⁾ OJ No C 302, 9. 11. 1993, p. 12.

shall result in criminal and/or administrative penalties being imposed by the appropriate national authorities.

If it is proven that the slaughterhouse owner or supervisor has been an accessory to the concealment of the illegal use of prohibited substances, he shall be liable to be banned from receiving and applying for Community aid for a period of 12 months.'

- 5. Does not affect English text.
- 6. The first paragraph of Article 37 is replaced by the following:

'This Regulation shall enter into force on the seventh day following its publication in the Official Journal of the European Communities.'

Amended proposal for a Council Directive amending and updating Directive 64/432/EEC on animal health problems affecting intra-Community trade in bovine animals and swine

(94/C 222/10)

(Text with EEA relevance)

COM(94) 295 final

(Submitted by the Commission pursuant to Article 189a (2) of the EC Treaty on 8 July 1994)

On 10 January 1994 the Commission presented the above proposal to the Council (1). As a result of the opinion delivered by Parliament at its sitting on 19 April 1994, the original proposal is amended as follows:

- 1. In Annex I, Article 2 (g) is replaced by the following:
 - '(g) officially brucellosis-free part of a Member State: means a part of a Member State which satisfies the conditions laid down in Annex A.II.7, 8 and 9:'
- 2. In Annex I, Article 2 (k) is replaced by the following:
 - '(k) enzootic bovine leucosis-free Member State or part of a Member State: means a Member State or part of a Member State which meets the requirements laid down in Annex D, Chapter I, sections E, F and G;'
- 3. In Annex I, Article 5 (1) is replaced by the following:
 - '1. Bovine animals and swine covered by this Directive must be accompanied during transportation to their destination by a health certificate conforming to Annex F. The certificate shall consist of a single sheet and shall bear a serial number. It must be drawn up on the day of the health examination, in one of the official languages of the country of origin and in one of the official languages of the country of destination. The certificate shall be valid for 10 days from the date of the health examination. However, where the health examination

- 4. In Annex I, Article 6 (3) is replaced by the following:
 - '3. Bovine animals for slaughter must, in addition to the requirements in Articles 3, 4 and 5, come from holdings that are officially tuberculosis-free, enzootic bovine leucosis-free and, in the case of uncastrated bovines, from holdings that are officially brucellosis-free.'
- 5. In Annex I, Annex A.II the word 'region' in point 7 is replaced by 'part'.
- 6. In Annex I, Annex A.II the word 'region' in point 8 is replaced by 'part of a Member State'.
- 7. In Annex I, Annex A.II the word 'region' in point 9 is replaced by 'part of a Member State'.
- 8. In Annex I, Annex D, Chapter I.E the word 'region' is replaced by 'part' or 'part of a Member State' as appropriate.
- 9. In Annex I, Annex D, Chapter I.F the word 'region' is replaced by 'part' or 'part of a Member State' as appropriate.
- 10. In Annex I, Annex D, Chapter I.G (i) the word 'region' is replaced by 'part' or 'part of a Member State' as appropriate.

takes place after the animals have left the holding of origin, as provided for in paragraph 2, the certificate shall be valid for 10 days after they have left the holding of origin.'

⁽¹) OJ No C 33, 2. 2. 1994, p. 1.