Commission Regulation (EC) No 1231/2006 of 16 August 2006 amending Annexes I and II to Council Regulation (EEC) No 2377/90 laying down a Community procedure for the establishment of maximum residue limits of veterinary medicinal products in foodstuffs of animal origin, as regards ceftiofur and polyoxyethylene sorbitan monooleate and trioleate (Text with EEA relevance)

COMMISSION REGULATION (EC) No 1231/2006

of 16 August 2006

amending Annexes I and II to Council Regulation (EEC) No 2377/90 laying down a Community procedure for the establishment of maximum residue limits of veterinary medicinal products in foodstuffs of animal origin, as regards ceftiofur and polyoxyethylene sorbitan monooleate and trioleate

(Text with EEA relevance)

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Council Regulation (EEC) No 2377/90 of 26 June 1990 laying down a Community procedure for the establishment of maximum residue limits of veterinary medicinal products in foodstuffs of animal origin[(1)], and in particular Articles 2 and 3 thereof,

Having regard to the opinions of the European Medicines Agency formulated by the Committee for Medicinal Products for Veterinary Use,

Whereas:

(1) All pharmacologically active substances used in the Community in veterinary medicinal products intended for food-producing animals should be evaluated in accordance with Regulation (EEC) No 2377/90.

(2) The substance Ceftiofur is currently included in Annex I to Regulation (EEC) No 2377/90 for bovine and porcine for muscle, fat, liver and kidney and for bovine for milk. The entry for Ceftiofur in that Annex should be modified to include ovine and extended to all mammalian food-producing species for muscle, fat, liver, kidney and milk.

(3) The substance polyoxyethylene sorbitan monooleate is currently included in Annex II to Regulation (EEC) No 2377/90 for all food-producing species. The entry in that Annex for polyoxyethylene sorbitan monooleate should be replaced by polyoxyethylene sorbitan monooleate and trioleate covering polyoxyethylene sorbitan trioleate for all food-producing species.

(4) Regulation (EEC) No 2377/90 should therefore be amended accordingly.

(5) An adequate period should be allowed before the applicability of this Regulation in order to enable Member States to make any adjustment which may be necessary in the light of this Regulation to the authorisations to place the veterinary medicinal products concerned on the market which have been granted in accordance with Directive

2001/82/EC of the European Parliament and of the Council of 6 November 2001 on the Community code relating to veterinary medicinal products[(2)] to take account of the provisions of this Regulation.

(6) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on Veterinary Medicinal Products,

HAS ADOPTED THIS REGULATION:

(**1**) OJ L 224, 18.8.1990, p. 1. Regulation as last amended by Commission Regulation (EC) No 1055/2006 (OJ L 192, 13.7.2006, p. 3).

(**2**) OJ L 311, 28.11.2001, p. 1. Directive as last amended by Directive 2004/28/EC (OJ L 136, 30.4.2004, p. 58).

Status:
Point in time view as at 31/01/2020.

Changes to legislation:
There are currently no known outstanding effects for the Commission Regulation (EC) No 1231/2006, Introductory Text.