Commission Regulation (EC) No 1646/2004 of 20 September 2004 amending Annex I 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 (Text with EEA relevance)

Article 1 Annex I to Regulation (EEC) No 2377/90 is hereby amended...
Article 2 This Regulation shall enter into force on the third day...
Signature

ANNEX

The following substance(s) is(are) inserted in Annex I:
Pharmacologically active substance(s) Marker residue Animal species MRLs Target tissues...
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Further provisions in Commission Directive 98/82/EC are to be observed...
Pharmacologically active substance(s) Marker residue Animal species MRLs Target tissues...

2. Antiparasitic agents

2.1. Agents acting against endoparasites

2.1.3. Benzimidazoles and pro-benzimidazoles

2.1.1. Salicylanilides

2.2. Agents acting against ectoparasites

2.2.2. Formamidines

2.2.3. Pyrethroids

5. Corticoids

5.1. Glucocorticoids

(**1**) OJ L 224, 18.8.1990, p. 1. Regulation as last amended by Commission Regulation (EC) No 1101/2004 (OJ L 211, 12.6.2004, p. 3).

(**2**) Availability of veterinary medicinal products Communication from the Commission to the Council and the European Parliament COM(2000) 806 final.

(**3**) 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).

Changes to legislation:
There are currently no known outstanding effects for the Commission Regulation (EC) No 1646/2004.