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)

Article 1 Annexes I and II to Regulation (EEC) No 2377/90 are...
Article 2 This Regulation shall enter into force on the third day...
Signature

ANNEX

A. The following substance is inserted in Annex I to Regulation...
   1. Anti-infectious agents
      1.2. Antibiotics
         1.2.2. Cephalosporins

B. The following substance is inserted in Annex II to Regulation...
   3. Substances generally recognised as safe

(**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).

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