(7) The maximum residue limits (MRLs) for turkeys and chickens for fattening introduced into the Annex to Regulation (EC) No 1800/2004 by Commission Regulation (EC) No 101/2009 [1] and the trade name 'Robenz 66 G' for turkeys and chickens for fattening introduced into the Annex to Regulation (EC) No 1800/2004 by Commission Regulation (EC) No 214/2009 [2] were, by error, omitted in the Annex to Regulation (EC) No 1800/2004 as amended by Implementing Regulation (EU) No 532/2011. It is therefore necessary to reintroduce these MRLs and the trade name.

(8) Therefore, the Annex to Implementing Regulation (EU) No 532/2011 should be corrected accordingly.

(9) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS REGULATION:

Article 1

Amendment to Regulation (EC) No 2380/2001

In column 2 of the Annex to Regulation (EC) No 2380/2001, the words 'Alpharma Belgium BVBA' are replaced by 'Pfizer Ltd'.

Article 2

Amendment to Regulation (EC) No 1289/2004

In column 2 of the Annex to Regulation (EC) No 1289/2004, the words 'Alpharma (Belgium) BVBA' are replaced by 'Pfizer Ltd'.

Article 3

Amendment to Regulation (EC) No 1455/2004

In column 2 of the Annex to Regulation (EC) No 1455/2004, the words 'Alpharma (Belgium) BVBA' are replaced by 'Pfizer Ltd'.

Article 4

Amendment to Regulation (EC) No 1800/2004

In column 2 of the Annex to Regulation (EC) No 1800/2004, the words 'Alpharma (Belgium) BVBA' are replaced by 'Pfizer Ltd'.

Article 5

Amendment to Regulation (EC) No 600/2005

In column 2 of Annex I to Regulation (EC) No 600/2005, the words 'Alpharma (Belgium) BVBA' are replaced by 'Pfizer Ltd'.

Article 6

Amendment to Regulation (EU) No 874/2010

In column 2 of the Annex to Regulation (EU) No 874/2010, the words 'Alpharma (Belgium) BVBA' are replaced by 'Pfizer Ltd'.

Article 7

Amendment to Implementing Regulation (EU) No 388/2011

In column 2 of the Annex to Implementing Regulation (EU) No 388/2011, the words 'Alpharma (Belgium) BVBA' are replaced by 'Pfizer Ltd'.

Article 8

Amendment to Implementing Regulation (EU) No 532/2011

In column 2 of Annex I to Implementing Regulation (EU) No 532/2011 the words 'Alpharma Belgium BVBA' are replaced by 'Pfizer Ltd'.

Article 9

Amendment to Implementing Regulation (EU) No 900/2011

In column 2 of the Annex to Implementing Regulation (EU) No 900/2011, the words 'Alpharma (Belgium) BVBA' are replaced by 'Pfizer Ltd'.

Article 10

Correction to Implementing Regulation (EU) No 532/2011

Annex II to Implementing Regulation (EU) No 532/2011 is corrected in accordance with the Annex to this Regulation.

Article 11

Transitional measures

Existing stocks which are in conformity with the provisions applying before the date of entry into force of this Regulation may continue to be placed on the market and used until 2 September 2012.

Article 12

Entry into force

This Regulation shall enter into force on the 20th day following its publication in the *Official Journal of the European Union*.

Article 10 and the Annex shall, however, apply from 21 June 2011.

[1] OJ L 34, 4.2.2009, p. 5.
[2] OJ L 73, 19.3.2009, p. 12.

ANNEX

In Annex II to Implementing Regulation (EU) No 532/2011, the Annex to Regulation (EC) No 1800/2004 as amended by Implementing Regulation (EU) No 532/2011 is corrected as follows:

(1) in column 3 the words '(Cycostat 66G)' are replaced by '(Robenz 66 G)';

(2) A new column is added:

'Maximum residue limits (MRLs) in the relevant foodstuffs of animal origin
800 µg robenidine hydrochloride/kg of wet liver. 350 µg robenidine hydrochloride/kg of wet kidney. 200 µg robenidine hydrochloride/kg of wet muscle. 1 300 µg robenidine hydrochloride/kg of wet skin/fat.
400 µg robenidine hydrochloride/kg of skin/fat. 400 µg robenidine hydrochloride/kg of wet liver. 200 µg robenidine hydrochloride/kg of wet kidney. 200 µg robenidine hydrochloride/kg of wet muscle.'