CORRIGENDA

Corrigendum to Commission Regulation (EC) No 1101/2004 of 10 June 2004 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

(Official Journal of the European Union L 211 of 12 June 2004)

In the Annex, on page 5, point A is replaced as follows:

'A. The following substance(s) is(are) inserted in Annex I (List of pharmacologically active substances for which maximum residue limits have been fixed).

1. Anti-infectious agents

1.2. Antibiotics

1.2.4. Macrolides

Pharmacologically active substance(s)	Marker residue	Animal species	MRLs	Target tissues
"**Tulathromycin**	(2R,3S,4R,5R,8R,10R,11R,12S, 13S,14R)-2-ethyl-3,4,10,13-tetrahydroxy-3,5,8,10,12,14-hexamethyl-11-[[3,4,6-trideoxy-3-(dimethylamino)-ß-D-xylo-hexopy-ranosyl]oxy]-1-oxa-6-azacyclopent-decan-15-one expressed as tulathromycin equivalents	Bovine [1]	100 µg/kg	Fat
			3 000 µg/kg	Liver
			3 000 µg/kg	Kidney
		Porcine	100 µg/kg	Skin + fat
			3 000 µg/kg	Liver
			3 000 µg/kg	Kidney

[1] Not for use in animals from which milk is produced for human consumption."'