ANNEX

In Table 1 of the Annex to Regulation (EU) No 37/2010, the entry for diclazuril is replaced by the following:

Pharmacologically active Substance	**Marker residue**	**Animal Species**	**MRL**	**Target Tissues**	**Other Provisions (*as listed in to Article 14(7) of Regulation (EC) No 470/2009*)**	**Therapeutic classification**
‘Diclazuril	NOT APPLICABLE	All ruminants, porcine	No MRL required	NOT APPLICABLE	For oral use only	NO ENTRY
	Diclazuril	Poultry	500 µg/kg	Muscle	Not for use in animals from which eggs are produced for human consumption	Antiparasitic agents/ Agents acting againsts protozoa’
			500 µg/kg	Skin and fat in natural proportions		
			1 500 µg/kg	Liver		
			1 000 µg/kg	Kidney		
		Rabbit	150 µg/kg	Muscle		
			300 µg/kg	Fat		
			2 500 µg/kg	Liver		
			1 000 µg/kg	Kidney		

Changes to legislation:
There are currently no known outstanding effects for the Commission Implementing Regulation (EU) No 1235/2013, ANNEX.