

Commission Implementing Regulation (EU) No 201/2014 of 3 March 2014 amending the Annex to Regulation (EU) No 37/2010 on pharmacologically active substances and their classification regarding maximum residue limits in foodstuffs of animal origin, as regards the substance tildipirosin (Text with EEA relevance)

- Article 1 The Annex to Regulation (EU) No 37/2010 is amended as...  
Article 2 This Regulation shall enter into force on the twentieth day...  
Signature

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ANNEX

In Table 1 of the Annex to Regulation (EU) No...  
Pharmacologically active Substance Marker residue Animal  
Species MRL Target Tissues...

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**Changes to legislation:** There are currently no known outstanding effects for the Commission Implementing Regulation (EU) No 201/2014. (See end of Document for details)

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- (1) [OJ L 152, 16.6.2009, p. 11.](#)
- (2) Commission Regulation (EU) No 37/2010 of 22 December 2009 on pharmacologically active substances and their classification regarding maximum residue limits in foodstuffs of animal origin ([OJ L 15, 20.1.2010, p. 1.](#))
- (3) Commission Regulation (EU) No 759/2010 of 24 August 2010 amending the Annex to Regulation (EU) No 37/2010 on pharmacologically active substances and their classification regarding maximum residue limits in foodstuffs of animal origin, as regards the substance tildipirosin ([OJ L 223, 25.8.2010, p. 39.](#))

**Changes to legislation:**

There are currently no known outstanding effects for the Commission Implementing Regulation (EU) No 201/2014.