

Commission Implementing Regulation (EU) 2020/1712 of 16 November
2020 amending Regulation (EU) No 37/2010 to classify the substance
lidocaine as regards its maximum residue limit (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

ANNEX

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

Changes to legislation: There are currently no known outstanding effects for the Commission Implementing Regulation (EU) 2020/1712. (See end of Document for details)

- (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](#)).

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