Commission Implementing Regulation (EU) 2018/470 of 21 March 2018 on detailed rules on the maximum residue limit to be considered for control purposes for foodstuffs derived from animals which have been treated in the EU under Article 11 of Directive 2001/82/EC (Text with EEA relevance)

Article 4

For pharmacologically active substances included in Table 1 of the Annex to Regulation (EU) No 37/2010 for which no MRL is required, in accordance with Article 14(5) of Regulation (EC) No 470/2009, there shall be no MRL required for control purposes for any target tissue derived from animal species treated in the EU under Article 11 of Directive 2001/82/EC, provided that the restrictions established in Table 1 are complied with.

Changes to legislation:
There are currently no known outstanding effects for the Commission Implementing Regulation (EU) 2018/470, Article 4.