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ANNEX 1

ANNEX

to the

COMMISSION IMPLEMENTING REGULATION (EU) .../...

amending Implementing Regulation (EU) No 540/2011 as regards the conditions of approval of the active substance fenpyrazamine

ANNEX […]

The column 'Purity' of row 25, fenpyrazamine, of Part B of the Annex to Implementing Regulation (EU) No 540/2011 is replaced by the following:

'≥ 960 g/kg

The following manufacturing impurity is of toxicological concern and must not exceed the following amount in the technical material:

Hydrazine: maximum content: < 0.0001% (1 mg/kg)'

The column ‘Specific provisions’ of row 25, fenpyrazamine, of Part B of the Annex to Implementing Regulation (EU) No 540/2011 is replaced by the following:

PART B

'For the implementation of the uniform principles as referred to in Article 29(6) of Regulation (EC) No 1107/2009, the conclusions of the review report on fenpyrazamine, and in particular Appendices I and II thereof, as finalised in the Standing Committee on the Food Chain and Animal Health on 1 June 2012 and the Standing Committee on Plants, Animals, Food and Feed on 18 May 2020 shall be taken into account. The purity given in this entry is based on a commercial plant production.'