

**Changes to legislation:** There are currently no known outstanding effects for the Commission Implementing Regulation (EU) 2018/1130, ANNEX. (See end of Document for details)

## ANNEX

Common Name	IUPAC Name Identification Numbers	Minimum degree of purity of the active substance <sup>a</sup>	Date of approval	Expiry date of approval	Product type	Specific conditions
Cypermethrin	IUPAC Name: Cypermethrin <i>cis:trans</i> 40:60; (RS)- $\alpha$ -cyano-3-phenoxybenzyl-(1RS)- <i>cis,trans</i> -3-(2,2-dichlorovinyl)-2,2-dimethylcyclopropane carboxylate EC No: 257-842-9 CAS No: 52315-07-8	$\geq 92$ % w/w Isomeric ratio: <i>cis:trans</i> 40:60	1 June 2020	31 May 2030	18	The authorisations of biocidal products are subject to the following conditions: 1. The product assessment shall pay particular attention to the exposures, the risks and the efficacy linked to any uses covered by an application for authorisation, but not

**a** The purity indicated in this column was the minimum degree of purity of the active substance evaluated. The active substance in the product placed on the market can be of equal or different purity if it has been proven to be technically equivalent to the evaluated active substance.

**b** Regulation (EC) No 470/2009 of the European Parliament and of the Council of 6 May 2009 laying down Community procedures for the establishment of residue limits of pharmacologically active substances in foodstuffs of animal origin, repealing Council Regulation (EEC) No 2377/90 and amending Directive 2001/82/EC of the European Parliament and of the Council and Regulation (EC) No 726/2004 of the European Parliament and of the Council (OJ L 152, 16.6.2009, p. 11).

**c** Regulation (EC) No 396/2005 of the European Parliament and of the Council of 23 February 2005 on maximum residue levels of pesticides in or on food and feed of plant and animal origin and amending Council Directive 91/414/EEC (OJ L 70, 16.3.2005, p. 1).









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						applicable MRLs are not exceeded.
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