EUROPEAN
COMMISSION

Brussels, XXX
SANTE/11868/2017 ANNEX
(POOL/E4/2017/11868/11868-EN
ANNEX.doc)
[…](2017) XXX draft

ANNEX 1

ANNEX

to the

COMMISSION IMPLEMENTING REGULATION (EU) …/...

approving azoxystrobin as an active substance for use in biocidal products of product-types 7, 9 and 10

ANNEX

Common Name	IUPAC Name Identification Numbers	Minimum degree of purity of the active substance[1]	Date of approval	Expiry date of approval	Product type	Specific conditions
Azoxystrobin	IUPAC Name: Methyl(E)-2-{2[6-(2-cyanophenoxy)pyrimidin-4-yloxy]phenyl}-3-methoxyacrylate EC No: not available CAS No: 131860-33-8	965 g/kg	1st November 2018	31 October 2025	7	Azoxystrobin is considered a candidate for substitution in accordance with point d) of Article 10(1) of Regulation (EU) No 528/2012. The authorisations of biocidal products are subject to the following condition: 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 addressed in the Union level risk assessment of the active substance. The placing on the market of treated articles is subject to the following condition: The person responsible for the placing on the market of a treated article treated with or incorporating azoxystrobin shall ensure that the label of that treated article provides the information listed in the second subparagraph of Article 58(3) of Regulation (EU) No 528/2012.
					9	Azoxystrobin is considered a candidate for substitution in accordance with point d) of Article 10(1) of Regulation (EU) No 528/2012. The authorisations of biocidal products are subject to the following condition: 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 addressed in the Union level risk assessment of the active substance.

[1] 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.

						The placing on the market of treated articles is subject to the following condition: The person responsible for the placing on the market of a treated article treated with or incorporating azoxystrobin shall ensure that the label of that treated article provides the information listed in the second subparagraph of Article 58(3) of Regulation (EU) No 528/2012.
					10	Azoxystrobin is considered a candidate for substitution in accordance with Article 10(1)(d) of Regulation (EU) No 528/2012. The authorisations of biocidal products are subject to the following condition: 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 addressed in the Union level risk assessment of the active substance. The placing on the market of treated articles is subject to the following condition: The person responsible for the placing on the market of a treated article treated with or incorporating azoxystrobin shall ensure that the label of that treated article provides the information listed in the second subparagraph of Article 58(3) of Regulation (EU) No 528/2012.