EUROPEAN
COMMISSION

Brussels, XXX
SANTE/2574/2022 ANNEX
(POOL/E4/2022/2574/2574-EN
ANNEX.docx)
[…](2023) XXX draft

ANNEX

ANNEX

to the

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

approving (13Z)-Hexadec-13-en-11-yn-1-yl acetate as an active substance for use in biocidal products of product-type 19 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

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
(13Z)-Hexadec-13-en-11-yn-1-yl acetate	IUPAC name: N,N-Didecyl-N,N-dimethylammonium chloride EC No: Not allocated CAS No: 78617-58-0	970 g/kg dry weight	1 June 2023	31 May 2033	19	The authorisation of biocidal products is subject to the following conditions: The product assessment shall pay particular attention to the exposures, the risks and the efficacy linked to any of the uses covered by an application for authorisation, but not addressed in the Union level risk assessment of the active substance. For uses leading to higher exposure of the users, by-standers or the environment compared to the Union level risk assessment of the active substance, applications for product authorisation shall contain all data required for active substances in accordance with Annex II to Regulation (EU) No 528/2012, subject to the possibilities of adaptation of the data requirements in accordance with Annex IV to that Regulation.

[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.