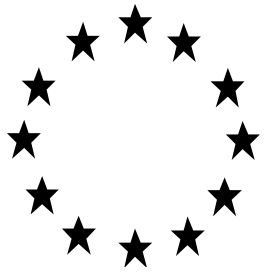
Regulation (EU) No 528/2012 concerning the making available on the market and use of biocidal products

PRODUCT ASSESSMENT REPORT OF A BIOCIDAL PRODUCT FOR NATIONAL AUTHORISATION APPLICATION

(submitted by the evaluating Competent Authority)

Addendum to PAR – access level Applicant



Myrr Q Muurahaisrasia

Product type 18

Imidacloprid as included in the Union list of approved active substances

Case Number in R4BP: BC-FM071504-38

Competent Authority: Finland

Date: [10 February 2023]

Table of Contents

Changes history table

Application type	refMS/eCA	Case number in the refMS	Decision date	Assessment carried out (i.e. first authorisation / amendment / renewal)	Chapter/ page
NA-BBS	FI	BC-AD047760-59	01.07.2019	Authorised as a same product	-
NA-ADC	FI	BC-LY063984-92	03.03.2021	Addition of active substance manufacturer and product formulator	-
NA-AAT	FI	BC-WQ079820-05	23.09.2022	Expiration date extended	-
NA-MIC	FI	BC-FM071504-38	10.2.2023	Extension of shelf life to 5 years.	5
NA-RNL	FI	BC-VQ065403-16	TBD	Renewal	

1 NA-MIC – Minor Change on request (BC-FM071504-38)

A new long term storage stability study was provided to extend the shelf life of the product. The physico-chemical and technical properties of the product remain constant. In addition, the palatability of the product has been demonstrated. Therefore, a shelf life of 5 years at ambient temperature is acceptable.

As the applicant has a Letter of Access to the data concerning the extension of the shelf life but has no permission to see or receive copies of any material about the product data, no detailed results are presented in this document. The evaluation of the competent authority is presented in detail in a separate annex, access level MSCA only.