AUTHORISATION NUMBER: **IE/BPA 70240**

EUROPEAN COMMUNITIES (AUTHORISATION, PLACING ON THE MARKET, USE AND CONTROL OF BIOCIDAL PRODUCTS) REGULATIONS

CERTIFICATE OF AUTHORISATION

The Competent Authority for Biocides in Ireland, pursuant to the provisions of Regulation (EU) No 528/2012 of the European Parliament and of the Council concerning the making available on the market and use of biocidal products, as amended by Regulation (EU) No 334/2014, and European Union (Biocidal Products) Regulations, 2013, (S.I. 427 of 2013), grants authorisation to make available on the market in Ireland, the biocidal product:

Product family name:	**Ant Bait 1R-trans phenothrin**	
Name and address of the authorisation holder	**Name**	Henkel AG & Co. KGaA
	Address	Henkelstrasse 67 Duesseldorf 40589 North-Rhine Westfalia Germany
Authorisation number	IE/BPA 70240	
Authorisation type	National Authorisation	
Date of the authorisation	20th August 2020	
Expiry date of the authorisation	30th April 2029	

subject to the conditions detailed in the Annexes to this certificate.

==========

Authorisation granted on behalf of the Competent Authority for Biocides in Ireland by

Louise Pierce

Mervyn Pa

Pesticide Control Division (PCD)

Official Stamp:

Version: 1.2

ANNEX I

Product Summary and Conditions of Authorisation

Family Name	Ant Bait 1R-trans phenothrin; IE/BPA 70240
Product names/ Trade Names	Vapona Ant Bait C1; IE/BPA 70240-1-001 Vapona Ant Bait C2; IE/BPA 70240-2-001 Vapona Ant Bait Extra; IE/BPA 70240-2-002 Vapona Ant Bait; IE/BPA 70240-003 Vapona Double Ant Bait*; IE/BPA 70240-3-001 Vapona Double Ant Bait*; IE/BPA 70240-3-002 * The Vadona Double Ant Bait contains two chambers with different mixtures in each chamber. Therefore, each product is authorized as a single product with its own authorization number (IE/BPA 70240-3-001 & IE/BPA 70240-3-002). The double bait box will be considered as a special type of package containing two biocidal products. However, only one trade name will appear on the packaging ('Vadona Double Ant Bait').
R4BP asset number	IE-0013719-0000

Active Substance(s) (% w/w):	1R-trans phenothrin 0.065 – 0.093%
Product-Type:	PT18 - Insecticides, acaricides and products to control other arthropods (Pest control)
Product Composition:	Confidential Annex available on R4BP3
Substance(s) of Concern:	Parmetol D11 (BIT) 0.0 – 0.048%
Comparative Assessment Required:	No
Formulation Type:	RB - Bait (ready for use)
Area of Use:	Indoor Outdoor For use in and around buildings.
Statement of Use:	Bait application Ant baits are ready-to-use bait or double bait stations against Lasius niger adult ants. See relevant meta-SPC number and timing of application section - - Meta SPC 1: One or two baits every 10 m2. This corresponds to AS concentrations 0.0052-0.0105g/10m2 for the single bait containing (TP-050-C1), depending on whether 1 or 2 baits /10m2 is used. Replace the ant bait every three months or when the ant bait is empty.

	Meta SPC 2: One or two baits every 10 m2 for product IIRD-08002 which corresponds to a dose rate of 0.0037-0.0073g/10m2 and J-70021 which corresponds to a dose rate of 0.0049-0.0098 g/10m2 depending on whether 1 or 2 bait stations are used per 10m2. Product TP-050-C2 may be marketed as a single bait station and is used at a dose rate of 1 or 2 baits per 10m2 (this correlates with a dose rate of 0.005 to 0.0105 g/10m2). .Replace the ant bait every three months or when the ant bait is empty Meta SPC 3; Double bait: One or two baits every 10 m2. This corresponds to AS concentrations of 0.0059-0.0119g/10m2 for the double bait containing (TP-050-C1b & TP-050-C2) depending on whether 1 or 2 baits /10m2 is used. Replace the ant bait every three months or when the ant bait is empty
User Category:	General Public
Special labelling provisions for Ireland:	In addition to the details recorded on the SPC, the following details shall be recorded on the product label(s). **Use Biocides Safely and Sustainably.** **It is illegal to use this product for uses or in a manner other than that prescribed on this label.** **Bait stations:** Must be labelled with the following information: "Product name or authorisation number(s)"; "Active substance(s)" and "In case of incident, call the National Poisons Information Centre on (01) 809 2166". **Poison Information**: For information or to report a poisoning incident **contact The National Poisons Information Centre, Beaumont Hospital, Dublin (01-809 2166)**, retain the label for reference. **First Aid:** In case of: Oral exposure, rinse mouth carefully with water. Never give anything by mouth to unconscious person. Do not provoke vomiting. If swallowed, seek medical advice immediately and show the product's container or label. Contact a veterinary surgeon in case of ingestion by a pet.

This authorisation may be subject to review in accordance with Regulation (EU) No 528/2012, as amended by Regulation (EU) No 334/2014, or the European Union (Biocidal Products) Regulations, 2013, (S.I. 427 of 2013). The outcome of such a review may lead to amendments to or the revocation of this authorisation.

The following conditions and restrictions apply:

1. Product may **not** be made available on the market or used in the Republic of Ireland unless it complies with the Annexes of this authorisation.

2. The requirements and conditions, specified in the Annexes, of this authorisation may **not** be altered without prior approval of modifications by the Irish Competent Authority for

Biocides in Ireland. Where any amendments are made to the original authorisation in another Member State, the Irish Competent Authority for Biocides in Ireland must be informed by the Authorisation Holder.

3. The holder of this certificate for authorisation must inform or provide the Irish Competent Authority for Biocides with any new or requested information/data, respectively, that shows this biocidal product and/or any of its active substances cause or may cause an adverse effect on human or animal health, ground water or the environment.

4. All product made available on the market in Ireland must comply with the classification, labelling and packaging requirements established in: Article 69 of Regulation (EU) No 528/2012; the Chemicals Act 2008 (as amended) giving further effect to Regulation (EC) No 1272/2008; and the classification, labelling and Safety Data Sheet information detailed in the Annex II to this certificate.

5. All biocidal products advertised must comply with Article 72 of Regulation (EU) No 528/2012.

6. A printed copy of the Irish label(s) in accordance with the Annexes of this authorisation must be submitted to the Irish Competent Authority for Biocides prior to any product being made available on the market in Ireland. All product labels must carry the authorisation number in the form: IE/BPA 70240.

7. Safety Data Sheets (SDS) for the biocidal product(s) shall be prepared and made available in accordance with Article 70 of the Biocidal Products Regulation 528/2012 (as amended). Relevant sections of the SDS must be updated post-authorisation in accordance with Annex II of the authorisation certificate. In particular, Section 15 of the SDS should be updated to contain the authorisation number IE/BPA 70240. The SDS must be submitted to the Irish Competent Authority for Biocides and the National Poisons Information Centre of Ireland http://www.poisons.ie/manufacturers.asp before the product is made available on the market for sale or use.

8. On an annual basis, details of the quantities of this product (by pack size) manufactured in Ireland, imported into Ireland and/or exported from Ireland must be submitted to the Irish Competent Authority for Biocides by 31 January of the following year. Details of the distributor's name and address must also be submitted at this time.

9. Authorisation holders and marketing companies must inform distributors, wholesalers and retailers of their requirements to keep records of goods in and goods out which can be requested by DAFM for inspection.

10. Fees are payable for the maintenance of the product on the Register of Biocidal Products and shall be paid by the 31st December of the following year and each year thereafter.

11. The Irish Competent Authority for Biocides requires Irish specific resistance data to be generated for the active substance contained in this product and submitted at renewal.

(b) Amendments to Authorisation

The following amendments apply to the conditions of authorisation for the biocidal product:

Issue	Re-issue	Version	Modifications applied[2]
30/04/2019	-	1.0	Original certificate
	20/08/2020	1.1	Minor change - extension of shelf-life BC-VV054362-07
	22/03/2023	1.2	NA-ADC – Updated instructions for use BC-SK075580-25

ANNEX II

Summary of Product Characteristics (SPC) for a biocidal product

The following conditions, outlined in the summary of product characteristics (SPC), apply to the authorisation for the biocidal product as provided for in Article 22 of Regulation (EU) No 528/2012 as amended. The authorised biocidal product SPC file is referenced below:

Issue	Re-issue	Version	File Name
30/04/2019	-	1.0	spfbc_Ant Bait 1R-trans phenothrin_IE_en_201905131139
	20/08/2020	1.1	spfbc_Ant Bait 1R-trans phenothrin_IE_en_202009011223
	22/03/2023	1.2	spfbc_Ant_Bait_1R-trans_phenothrin_IE_en_202205161051