AUTHORISATION NUMBER: **IE/BPA 70384**

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:

Biocidal Product Name:	Detrans HPC3	
Name and address of the authorisation holder	**Name**	Sumitomo Chemical (UK) Plc
	Address	200 Shepherds Bush Road Hythe House W6 7NL London, UK
Authorisation number	IE/BPA 70384	
Authorisation type	Mutual Recognition in Parallel (NA-MRP)	
Date of the authorisation	15th April 2024	
Expiry date of the authorisation	22nd April 2032	

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

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

Louise Pierce

Mervyn Pa

Pesticide Control Division (PCD)

Official Stamp:

Version: 1.0

ANNEX I

Product Family Summary and Conditions of Authorisation

Biocidal Product Name	Detrans HPC3 IE/BPA 70384
Additional Trade names (with suffixes to the Authorisation number)	Raid® Bug Defence IE/BPA 70384-001 Raid® Bug Defense IE/BPA 70384-002 HOME DEFENCE ANT-STOP! GUN IE/BPA 70384-003
R4BP asset number	IE-0011027-0000
Marketing Company, Address	To be confirmed

Active Substance(s) (% w/w):	Deltamethrin (0.03% w/w)
Product-Type:	PT18 Insecticide
Product Composition:	See Confidential PAR on R4BP3
Substance(s) of Concern:	No
Comparative Assessment	No
Formulation Type:	AL-Any other liquid
Area of Use:	Indoor use
Statement of use:	Detrans HPC3 is an insectide product containing the active substance Deltamethrin (0.03% w/w) for indoor use by non-professionals.
User Category:	Non-professional
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.** **Poison Information: For information or to report a poisoning incident contact The National Poisons Information Centre, Beaumont Hospital, Dublin (01-8092166),** retain the label for reference.

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) transposing 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 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 of the form: IE/BPA 70384.

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 70384. 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.

9. 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.

(b) Amendments to Authorisation

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

Issue	Re-issue	Version	Modifications applied[2]
15/04/2024	-	1.0	Original Certificate NA-MRP (BC-YD012751-45)

ANNEX II

Summary of Product Characteristics (SPC) for a biocidal product family

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

Issue	Re-issue	Version	File Name
15/04/2024	-	1.0	04ae9daf-77b1-4a65-a67d-a657ad69ccf6