AUTHORISATION NUMBER: **IE/BPA 70854**

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 Family Name:	IMIDASECT	
Name and address of the authorisation holder	**Name**	Sharda Cropchem España S.L.
	Address	Edificio Atalayas Business Center, Carril Condomina N°3 Planta 12, 30006, Murcia, Spain
Authorisation number	IE/BPA 70854	
Authorisation type	Mutual recognition in Sequence (NA-MRS)	
Authorisation Details	Article 19(1)	
Date of the authorisation	**24th July 2026**	
Expiry date of the authorisation	**14th December 2026**	

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

Pesticide Control Division (PCD)

Official Stamp:

AN ROINN TALMHAÍOCHTA BIA AGUS MARA
ÉIRE ÉIRE
Dept. of Agriculture, Food and the Marine
IRELAND

Version: 1.0

ANNEX I

Product Family Summary and Conditions of Authorisation

Biocidal Product Name	IMIDASECT (Not Marketed) IE/BPA 70854
Additional Trade names (with suffixes to the Authorisation number)	ROACH BAIT GEL (Not Marketed) IE/BPA 70854-001
R4BP asset number	**IE-0037860-0000**
Marketing Company, Address	To be confirmed

Active Substance(s) (% w/w):	(2E)-1-[(6-chloropyridin-3-yl) methyl]-N-nitroimidazolidin-2-imine (2.194 % w/w)
Product-Type:	PT18 - Insecticides, acaricides and products to control other arthropods
Product Composition:	See Confidential PAR on R4BP3
Substance(s) of Concern:	No
Comparative Assessment	No
Formulation Type:	GD – Gel for direct application
Area of Use:	Indoor Outdoor
Statement of use:	IMIDASECT is an insecticide product containing 2.194 % w/w of the active substance (2E)-1-[(6-chloropyridin-3-yl) methyl]-N-nitroimidazolidin-2-imine, for indoor and outdoor use by professional and non-professional users. See SPC for full details.
User Category:	Professional General Public (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 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. As per Article 47(1) of Regulation (EU) No 528/2012, on becoming aware of information concerning this authorised biocidal product, or the active substance(s) it contains, that may affect this authorisation, the authorisation holder shall without delay notify Irish Competent Authority and the Agency (European Chemical Agency; ECHA).

   In particular, the following shall be notified:

   (a) new data or information on the adverse effects of the active substance or biocidal product for humans, in particular vulnerable groups, animals or the environment;

   (b) any data indicating the potential of the active substance for the development of resistance;

   (c) new data or information indicating that the biocidal product is not sufficiently effective.

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 70854.

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 70854. 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. In accordance with Article 68(1) of the Biocidal Products Regulation 528/2012 *'Authorisation holders shall keep records of the biocidal products they place on the market for at least 10 years after placing on the market, or 10 years after the date on*

which the authorisation was cancelled or expired, whichever is the earlier.' The relevant information contained in these records shall be submitted to the Irish Competent Authority for Biocides if requested. Details of the distributor(s) in Ireland (name and address) may also be requested.

9. Annual Registration Fees (ARF) for the maintenance of the product on the Register of Biocidal Products are payable to the Irish Competent Authority for all trade names listed on this certificate of authorisation. Following authorisation, ARF shall be paid by the 31st December of the following year and each year thereafter. This authorisation will be cancelled if ARF are not paid in due time.

10. In line with Article 48 of Regulation (EU) No 528/2012, the Irish Competent Authority may amend or cancel this authorisation where it is considered:

    (a) the conditions referred to in Article 19 are not satisfied; or

    (b) the authorisation was granted on the basis of false or misleading information; or

    (c) the authorisation holder has failed to comply with the conditions of authorisation or the BPR.

(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]
24/07/2026	-	1.0	Original Certificate NA-MRS (BC-DB073721-60)

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
24/07/2026	-	1.0	8f0b17b4-f831-48ad-bee3-1da4e698f3c0