AUTHORISATION NUMBER: **IE/BPA 70270**

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:	Quick Bayt WG10	
Name and address of the authorisation holder	**Name**	2022 ENVIRONMENTAL SCIENCE FR SAS
	Address	3, place Giovanni Da Verrazzano, 69009, Lyon, France
Authorisation number	IE/BPA 70270	
Authorisation type	Mutual Recognition in Parallel (NA-MRP)	
Date of the authorisation	24th July 2017	
Expiry date of the authorisation	7th October 2024	

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

Pesticide Control Division (PCD)

Official Stamp:

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

Ver: 1.7

ANNEX I

Product Summary and Conditions of Authorisation

Trade name	Quick Bayt WG10	IE/BPA 70270
Other Trade Names	Quick Bayt WG	IE/BPA 70270-001
	QUICK BAIT WG10	IE/BPA 70270-002
	QuickBait WG	IE/BPA 70270-003
R4BP asset number	IE-0011784-0000	

Active Substance(s) (% w/w):	Imidacloprid (9.99% w/w) Muscalure (Cis-Tricos-9-ene) (0.085% w/w)
Product-Type:	PT 18 (Insecticides, acaricides and products to control other arthropods)
Product Composition:	Confidential PAR on R4BP
Substance(s) of Concern:	Imidacloprid Muscalure
Formulation Type:	Water dispersible granules (WG)
Statement of use:	Quick Bayt WG10 is an insecticidal bait for the control of flies (Musca domestica) for use in animal units or agricultural buildings (e.g. broiler houses, livestock barns, caged layer houses and small animal husbandry e.g. kennels, etc.) as well as in sheltered waste management facilities (animal waste areas).Its method of application is by painting. - For the paint application, 250 g of bait is to be mixed with 200 mL of water providing sufficient paste to treat 100 m² of floor area. Treatments should be distributed across the stable covering a total area of approximately 2 m² for a floor area of 100 m². - Using a paint brush, the solution is applied as stripes or spots onto pieces of timber, or sheets of cardboard or fabric which are hung or attached in areas where flies tend to aggregate. These areas may be walls, on surfaces of pillars and around windows or other structures.
Area of Use:	IV.1. Indoor use only. IV.1.3 To be used in: IV.1.3.4 Animal Houses/Shelters IV.1.3.5 Sheltered waste management facilities (animal waste areas)
User Category:	Professional
Special labelling provisions for Ireland:	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 70270.

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 70270. 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:

Issue	Re-issue	Version	Modifications applied[2]
24/07/2017	-	1.0	Original certificate
-	29/07/2019	1.1	Transfer of National Authorisation
-	14/05/2021	1.2	Update to shelf-life
-	21/10/2022	1.3	Extension of authorisation
-	07/02/2023	1.4	Transfer of authorisation holder - BC-VE075842-28
-	14/03/2023	1.5	Extension of Authorisation to 07/04/2024
-	20/12/2023	1.6	Additional trade names NA-ADC (BC-WL089824-05)
-	25/01/2024	1.7	Extension of Authorisation to 07/10/2024

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
24/07/2017	-	1.0	spc_QUICK BAYT WG10_IE_en_201707201416.xml
-	29/07/2019	1.1	spc_QUICK_BAYT_WG10_IE_en_201901151510
-	14/05/2021	1.2	spc_QUICK BAYT WG10_IE_en_202105041515
-	21/10/2022	1.3	spc_QUICK BAYT WG10_IE_en_202105041515
-	07/02/2023	1.4	spc_QUICK BAYT WG10_IE_en_202105041515
-	14/03/2023	1.5	spc_QUICK BAYT WG10_IE_en_202105041515
-	20/12/2023	1.6	IE_spc_QUICK_BAYT_WG10_IE_en_202310311328
-	25/01/2024	1.7	IE_spc_QUICK_BAYT_WG10_IE_en_202310311328