



**AUTHORISATION NUMBER: IE/BPA 70486**

**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:

<b>Biocidal Product Family Name:</b>	XILIX 3000 P	
<b>Name and address of the authorisation holder</b>	<b>Name</b>	BERKEM DEVELOPPEMENT
	<b>Address</b>	Marais Ouest 24680 Gardonne France
<b>Authorisation number</b>	IE/BPA 70486	
<b>Authorisation type</b>	Mutual recognition in parallel (NA-MRP)	
<b>Date of the authorisation</b>	2 <sup>nd</sup> May 2019	
<b>Expiry date of the authorisation</b>	1 <sup>st</sup> July 2023	

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

  
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Felicia Bourne

Pesticide Control Division (PCD)

Official Stamp:



Version: 1.0

**ANNEX I****Product Family Summary and Conditions of Authorisation**

<b>Biocidal Product Name</b>	XILIX 3000 P
<b>Additional Trade names (with suffixes to the Authorisation number)</b>	CORIPHENE IF PRO IE/BPA 70486-001 DEADLINE WOODWORM TREATMENT IE/BPA 70486-002 WOODWORM TREATMENT FLUID IE/BPA 70486-003 WOODWORM TREATMENT SOLUTION IE/BPA 70486-004 XYLOFOR IFACO 3000 IE/BPA 70486-005
<b>R4BP asset number</b>	IE-0020593-0000
<b>Marketing Company, Address</b>	To be confirmed
<b>Active Substance(s) (% w/w):</b>	IPBC (0.073 % w/w) Propiconazole (0.073 % w/w) Tebuconazole (0.073 % w/w) Permethrin (0.35 % w/w)
<b>Product-Type:</b>	PT 08 – Wood Preservatives (Preservatives)
<b>Product Composition:</b>	See Confidential PAR on R4BP3
<b>Substance(s) of Concern:</b>	N/A
<b>Comparative Assessment</b>	YES
<b>Formulation Type:</b>	ME – Micro-emulsion
<b>Area of Use:</b>	Indoor use – Class 1 and class 2
<b>Statement of use:</b>	The XILIX 3000 P product is a preventive and curative treatment effective against wood-destroying insects (including termites) and wood destroying fungi (white rot and brown rot fungi).
<b>User Category:</b>	General public Professional
<b>Special labelling provisions for Ireland:</b>	In addition to the details recorded on the SPC, the following details shall be recorded on the product label(s).  <b>Use Biocides Safely and Sustainably</b> <b>It is illegal to use this product for uses or in a manner other than that prescribed on this label.</b>  <b>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.</b>

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 70486.
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 70486. 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 31<sup>st</sup> 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:

<b>Issue</b>	<b>Re-issue</b>	<b>Version</b>	<b>Modifications applied<sup>2</sup></b>
02/05/2019	-	1.0	Original certificate

**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:

<b>Issue</b>	<b>Re-issue</b>	<b>Version</b>	<b>File Name</b>
02/05/2019	-	1.0	spc_XILIX 3000 P_IE_en_201905021240