

AUTHORISATION NUMBER: IE/BPA 70268

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:	Celcure C65	
Name and address of the authorisation holder	Name	Koppers Performance Chemicals Denmark ApS
	Address	Avernakke 1, 5800, Nyborg, Denmark
Authorisation number	IE/BPA 70268	
Authorisation type	Mutual recognition in parallel (NA-MRP)	
Date of the authorisation	01st February 2019	
Expiry date of the authorisation	29 th October 2028	

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

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

Pesticide Control Division (PCD)

Official Stamp:

Version: 1.3



ANNEX I Product Family Summary and Conditions of Authorisation

Biocidal Product Name	Celcure C65 IE/BPA 70268			
Additional Trade names (with suffixes to the Authorisation number)	N/A			
R4BP asset number	IE-0014500-0000			
Marketing Company, Address	To be confirmed			
Active Substance(s) (% w/w):	Basic Copper Carbonate (17.27%) DDA Chloride (20.65%) DDA Carbonate (4.345%)			
Product-Type:	PT 08 – Wood Preservatives			
Product Composition:	See Confidential PAR on R4BP3			
Substance(s) of Concern:	Ethanolamine			
Comparative Assessment	No			
Formulation Type:	Soluble concentrate (SL)			
Area of Use:	Indoor use Outdoor use-For use on wood in: -Use Class 1 (situation in which the wood or wood-based product is inside a construction, not exposed to the weather and wetting)Use Class 2 (situation in which the wood or wood-based product is under cover and fully protected from the weather but occasional, non-persistent, wetting may occur. This can include outdoor placement of timber under a roof to prevent any exposure to rain and driven rain)Use Class 3 (situation in which the wood or wood-based product is not covered and not in contact with the ground. It is either continuously exposed to weather or protected from the weather but subject to frequent wetting). Wood treated for use in this use class cannot claim to offer protection against termites -Use Class 4 (situation in which the wood or wood-based product is not covered and in contact with the ground. It is continually exposed to the weather). Treated timber must not be placed over/near surface water bodies.			
Statement of use:	Celcure C65 is a wood preservative applied as a penetrative treatment by vacuum pressure impregnation to control wood rotting fungi and wood destroying insects (including wood boring beetles and termites). It is supplied as two concentrates which are mixed and diluted in water. The product is intended to protect softwood timbers in use classes 1 (wood or wood based product inside a construction, not exposed			

	to weather and wetting), 2 (undercover but exposed to high humidity or occasional wetting), 3 (above ground but exposed to frequent wetting) and 4 (not covered and in contact with the ground. It is continually exposed to the weather).
User Category:	Industrial
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 70268.

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 70268. 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
01/02/2019		1.0	Original certificate
w	12/08/2019	1.1	Transfer of National Authorisation
-	24/11/2021	1.2	Addition of manufacturers to SPC
-	31/05/2023	1.3	Transfer of Authorisation Holder NA-TRS BC-EY080507-08

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
01/02/2019	-	1.0	spc_Celcure C65_IE_en_201901151723
-	12/08/2019	1.1	spc_Celcure C65_IE_en_201901151723
-	24/11/2021	1.2	spc_Celcure C65_IE_en_202111241241
All-	31/05/2023	1.3	spc_Celcure C65_IE_en_202111241241