Commission Implementing Regulation (EU) 2017/796 of 10 May 2017 approving dichlofluanid as an existing active substance for use in biocidal products of product-type 21 (Text with EEA relevance)

COMMISSION IMPLEMENTING REGULATION (EU) 2017/796

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approving dichlofluanid as an existing active substance for use in biocidal products of product-type 21

(Text with EEA relevance)

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Regulation (EU) No 528/2012 of the European Parliament and of the Council of 22 May 2012 concerning the making available on the market and use of biocidal products[(1)], and in particular the third subparagraph of Article 89(1) thereof,

Whereas:

(1) Commission Delegated Regulation (EU) No 1062/2014[(2)] establishes a list of existing active substances to be evaluated for their possible approval for use in biocidal products. That list includes dichlofluanid.

(2) Dichlofluanid has been evaluated for use in products of product-type 21, antifouling products, as described in Annex V to Regulation (EU) No 528/2012.

(3) The United Kingdom was designated as evaluating competent authority and submitted the assessment report together with its recommendations on 22 October 2015.

(4) In accordance with Article 7(2) of Delegated Regulation (EU) No 1062/2014, the opinion of the European Chemicals Agency was formulated on 11 October 2016 by the Biocidal Products Committee, having regard to the conclusions of the evaluating competent authority.

(5) According to that opinion, biocidal products of product-type 21 containing dichlofluanid may be expected to satisfy the criteria of Article 19(1)(b) of Regulation (EU) No 528/2012, provided that certain specifications and conditions concerning their use are complied with.

(6) The acceptability of the risks related to the use of antifouling products, as well as the suitability of the proposed risk mitigation measures, should however be further confirmed. In order to facilitate, at the time of the renewal of the approvals of existing antifouling active substances, the review and comparison of the risks and benefits of those substances as well as of the risk mitigation measures applied, the expiry date of approval of all those substances should be the same.

Changes to legislation: *There are currently no known outstanding effects for the Commission Implementing Regulation (EU) 2017/796. (See end of Document for details)*

(7) It is therefore appropriate to approve dichlofluanid for use in biocidal products of product-type 21, subject to compliance with certain specifications and conditions.

(8) Since dichlofluanid meets the criteria for classification as skin sensitiser category 1 as defined in Annex I to Regulation (EC) No 1272/2008 of the European Parliament and of the Council[3], treated articles treated with or incorporating dichlofluanid should be appropriately labelled when placed on the market.

(9) A reasonable period should be allowed to elapse before an active substance is approved in order to permit interested parties to take the preparatory measures necessary to meet the new requirements.

(10) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on Biocidal Products,

HAS ADOPTED THIS REGULATION:

Article 1

Dichlofluanid is approved as an active substance for use in biocidal products of product-type 21, subject to the specifications and conditions set out in the Annex.

Article 2

This Regulation shall enter into force on the twentieth day following that of its publication in the *Official Journal of the European Union*.

This Regulation shall be binding in its entirety and directly applicable in all Member States.

Done at Brussels, 10 May 2017.

For the Commission

The President

Jean-Claude JUNCKER

ANNEX

Common Name	**IUPAC NameIdentification Numbers**	**Minimum degree of purity of the active substance**[a]	**Date of approval**	**Expiry date of approval**	**Product type**	**Specific conditions**
Dichlofluanid	IUPAC Name: N-(Dichlorofluoromethylthio)-N′,N′-dimethyl-N-phenylsulfamide EC No: 214-118-7 CAS No: 1085-98-9	96 % w/w	1 November 2018	31 December 2025	21	The product assessment shall pay particular attention to the exposures, the risks and the efficacy linked to any uses covered by an application for authorisation, but not addressed in the Union-level risk assessment of the active substance. In the event that products containing dichlofluanid are subsequently authorised for use by non-professional users, persons making products available on the market

						for non-professional users shall ensure that the products are supplied with appropriate gloves. The authorisations of biocidal products are subject to the following conditions: (1) Products containing dichlofluanid shall not be authorised or used to control the growth and settlement of fouling organisms on freshwater going vessels. (2) For industrial or professional users, safe operational procedures and appropriate organisational measures

							shall be established. Where exposure cannot be reduced to an acceptable level by other means, products shall be used with appropriate personal protective equipment.
						(3)	Labels and, where provided, instructions for use shall indicate that children shall be kept away until treated surfaces are dry.
						(4)	Labels and, where provided, safety data sheets of products