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**COMMISSION IMPLEMENTING REGULATION (EU) .../...**

**of **XXX****

**approving alkyl (C<sub>12-16</sub>) dimethylbenzyl ammonium chloride as an active substance for  
use in biocidal products of product-types 3 and 4**

(Text with EEA relevance)

# COMMISSION IMPLEMENTING REGULATION (EU) .../...

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## **approving alkyl (C<sub>12-16</sub>) dimethylbenzyl ammonium chloride as an active substance for use in biocidal products of product-types 3 and 4**

(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<sup>1</sup>, and in particular the third subparagraph of Article 89(1) thereof,

Whereas:

- (1) Commission Delegated Regulation (EU) No 1062/2014<sup>2</sup> establishes a list of existing active substances to be evaluated for their possible approval for use in biocidal products. That list includes alkyl (C<sub>12-16</sub>) dimethylbenzyl ammonium chloride (ADBAC/BKC (C<sub>12</sub>-C<sub>16</sub>)) to be renamed for the purposes of this Regulation as alkyl (C<sub>12-16</sub>) dimethylbenzyl ammonium chloride as a result of its evaluation.
- (2) Alkyl (C<sub>12-16</sub>) dimethylbenzyl ammonium chloride has been evaluated for use in biocidal products of product-type 3, veterinary hygiene biocidal products and product-type 4, food and feed area disinfectants, as defined in Annex V to Directive 98/8/EC of the European Parliament and of the Council<sup>3</sup>, which correspond respectively to product-types 3 and 4 as defined in Annex V to Regulation (EU) No 528/2012.
- (3) Italy was designated as the rapporteur Member State and its evaluating competent authority submitted the assessment report together with its conclusions to the Commission on 10 September 2012.
- (4) In accordance with Article 7(2) of Delegated Regulation (EU) No 1062/2014, the Biocidal Products Committee adopted the opinions of the European Chemical Agency<sup>4</sup> ('the Agency') on 6 October 2020, having regard to the conclusions of the evaluating competent authority.
- (5) According to those opinions, biocidal products of product-types 3 and 4 containing alkyl (C<sub>12-16</sub>) dimethylbenzyl ammonium chloride may be expected to satisfy the

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<sup>1</sup> OJ L 167, 27.6.2012, p.1

<sup>2</sup> Commission Delegated Regulation (EU) No 1062/2014 of 4 August 2014 on the work programme for the systematic examination of all existing active substances contained in biocidal products referred to in Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L 294, 10.10.2014, p. 1).

<sup>3</sup> Directive 98/8/EC of the European Parliament and of the Council of 16 February 1998 concerning the placing of biocidal products on the market (OJ L 123, 24.4.1998, p. 1).

<sup>4</sup> Biocidal Products Committee Opinions on the applications for approval of the active substance alkyl(C<sub>12-16</sub>) dimethylbenzyl ammonium chloride; Product types: 3 and 4; ECHA/BPC/267/2020 and ECHA/BPC/268/2020, adopted on 6 October 2020.

requirements laid down in Article 5(1)(b), (c) and (d) of Directive 98/8/EC, provided that certain specifications and conditions concerning their use are complied with.

- (6) Taking into account the opinions of the Agency, it is appropriate to approve alkyl (C<sub>12-16</sub>) dimethylbenzyl ammonium chloride as an active substance for use in biocidal products of product-types 3 and 4, subject to compliance with certain specifications and conditions.
- (7) 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.
- (8) 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*

Alkyl (C<sub>12-16</sub>) dimethylbenzyl ammonium chloride is approved as an active substance for use in biocidal products of product-types 3 and 4 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,

*For the Commission*  
*The President*  
*Ursula VON DER LEYEN*