Commission Implementing Regulation (EU) 2020/1771 of 26 November 2020 approving reaction mass of peracetic acid (PAA) and peroxyoctanoic acid (POOA) as an existing active substance for use in biocidal products of product-types 2, 3 and 4 (Text with EEA relevance)

COMMISSION IMPLEMENTING REGULATION (EU) 2020/1771

of 26 November 2020

approving reaction mass of peracetic acid (PAA) and peroxyoctanoic acid (POOA) as an existing active substance for use in biocidal products of product-types 2, 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⁽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 peroxyoctanoic acid, to be renamed reaction mass of peracetic acid and peroxyoctanoic acid, as the result of its evaluation.
- (2) Reaction mass of peracetic acid and peroxyoctanoic acid has been evaluated for use in biocidal products of product-type 2, disinfectants and algaecides not intended for direct application to humans or animals, product-type 3, veterinary hygiene, and product-type 4, food and feed area, as described in Annex V to Regulation (EU) No 528/2012.
- (3) France was designated as the rapporteur Member State and its evaluating competent authority submitted the assessment report together with its conclusions to the European Chemicals Agency ('the Agency') on 2 January 2019.
- (4) In accordance with Article 7(2) of Delegated Regulation (EU) No 1062/2014, the Biocidal Products Committee adopted the opinions of the Agency⁽³⁾ on 4 March 2020, having regard to the conclusions of the evaluating competent authority.
- (5) According to those opinions, biocidal products of product-types 2, 3 and 4 containing reaction mass of peracetic acid and peroxyoctanoic acid may be expected to meet the criteria laid down in point (b) of Article 19(1) of Regulation (EU) No 528/2012, provided that certain specifications and conditions concerning their use are complied with.

Status: Point in time view as at 26/11/2020.

Changes to legislation: There are currently no known outstanding effects for the Commission Implementing Regulation (EU) 2020/1771. (See end of Document for details)

- (6) Taking into account the opinions of the Agency, it is appropriate to approve reaction mass of peracetic acid and peroxyoctanoic acid for use in biocidal products of product-types 2, 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

Reaction mass of peracetic acid and peroxyoctanoic acid is approved as an active substance for use in biocidal products of product-types 2, 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, 26 November 2020.

For the Commission
The President

Ursula VON DER LEYEN

Status: Point in time view as at 26/11/2020.

Changes to legislation: There are currently no known outstanding effects for the Commission Implementing Regulation (EU) 2020/1771. (See end of Document for details)

ANNEX

Common Name		IUPAC Minimum degree of purity NameIdenfffinetienive substance ^a Numbers							ons	
Reaction mass of peracetic acid (PAA) and peroxyoct acid (POOA)	name: Reaction mass of peracetic acid talkia) and peroxyoc acid (POOA) EC No: 201-186-3 and 450-280-7 CAS No: 79-21-0	the active not releva substance equilibriu peroxide, octanoic a materials. auriespon concentra	is a doubl m using hy acetic acid acid as star The speci d to a rang tion.	is ctive e ydrogen I and ting fications se of	1 April 2022	31 March 2032	2	The authorisations of biocidal products are subject to the following conditions: (a) The		
			nts	Specificat range content (%w/w)	tions				product assessment shall pay particular attention	
		Active substance	Peracetic acid	1,8–13,9						
		Active Substance	Peroxyocacid	tandse- 2,42					to the exposures,	
		3734-57 Substance Relevant impurity								the risks and the efficacy linked to any uses covered by an application for authorisation, but not addressed in the Union level

a The purity indicated in this column was the minimum degree of purity of the active substance evaluated. The active substance in the product placed on the market can be of equal or different purity if it has been proven to be technically equivalent to the evaluated active substance.

b Regulation (EC) No 1935/2004 of the European Parliament and of the Council of 27 October 2004 on materials and articles intended to come into contact with food and repealing Directives 80/590/EEC and 89/109/EEC (OJ L 338, 13.11.2004, p. 4).

Status: Point in time view as at 26/11/2020.

						(b)	assessment of the active substance. In view of the risks identified for the uses assessed, the product assessment shall pay particular attention to professional users.
	Relevant impurity		5,74–51		3	The authorisa	tions
	Relevant impurity	Octanoic acid	1,63- 9,03			of biocidal products are subject to the following condition (a)	s: The product assessment shall pay particular attention to the exposures, the risks

a The purity indicated in this column was the minimum degree of purity of the active substance evaluated. The active substance in the product placed on the market can be of equal or different purity if it has been proven to be technically equivalent to the evaluated active substance.

b Regulation (EC) No 1935/2004 of the European Parliament and of the Council of 27 October 2004 on materials and articles intended to come into contact with food and repealing Directives 80/590/EEC and 89/109/EEC (OJ L 338, 13.11.2004, p. 4).

Status: Point in time view as at 26/11/2020.

		4	(b)	and the efficacy linked to any uses covered by an application for authorisation, but not addressed in the Union level assessment of the active substance. In view of the risks identified for the uses assessed, the product assessment shall pay particular attention to professional users.
		4	The authorisa	ations

a The purity indicated in this column was the minimum degree of purity of the active substance evaluated. The active substance in the product placed on the market can be of equal or different purity if it has been proven to be technically equivalent to the evaluated active substance.

b Regulation (EC) No 1935/2004 of the European Parliament and of the Council of 27 October 2004 on materials and articles intended to come into contact with food and repealing Directives 80/590/EEC and 89/109/EEC (OJ L 338, 13.11.2004, p. 4).

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			of	
			biocidal	
			products	
			are	
			subject	
			to the	
			following	g
			condition	ns:
			(a)	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
				assessment
				of
				the
				active
			(1)	substance.
			(b)	Products
				_containing

a The purity indicated in this column was the minimum degree of purity of the active substance evaluated. The active substance in the product placed on the market can be of equal or different purity if it has been proven to be technically equivalent to the evaluated active substance.

b Regulation (EC) No 1935/2004 of the European Parliament and of the Council of 27 October 2004 on materials and articles intended to come into contact with food and repealing Directives 80/590/EEC and 89/109/EEC (OJ L 338, 13.11.2004, p. 4).

Status: Point in time view as at 26/11/2020.

			reaction
			mass
			of
			peracetic
			acid
			and
			peroxyoctanoic acid
			shall
			not
			be
			incorporated
			in
			materials
			and
			articles
			intended
			to
			come
			into
			contact
			with
			food
			within
			the
			meaning
			of
			Article
			1(1)
			1(1) of
			01
			Regulation
			(EC)
			No
			1935/2004
			of
			the
			European
			Parliament
			and
			of
			the
			Council ^b ,
			unless
			the
			Commission
			has
			established

a The purity indicated in this column was the minimum degree of purity of the active substance evaluated. The active substance in the product placed on the market can be of equal or different purity if it has been proven to be technically equivalent to the evaluated active substance.

B Regulation (EC) No 1935/2004 of the European Parliament and of the Council of 27 October 2004 on materials and articles intended to come into contact with food and repealing Directives 80/590/EEC and 89/109/EEC (OJ L 338, 13.11.2004, p. 4).

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						specific
						limits
						on
						the
						migration
						of
						reaction
						mass
						of
						peracetic
						acid
						and
						peroxyoctanoic
						acid
						into
						food
						or
						it
						has
						established]
						in
						accordance
						with
						that
						Regulation
						that
						such
						limits
						are
						not
					(a)	necessary.
						In
						view
						of the
						risks
						identified
						for
						the
						uses
						assessed,
						the
						product
						assessment
						shall
						pay
						particular
Thomus	itri indicata 4 :	- 4hih	 ·	l		Fartioniai

a The purity indicated in this column was the minimum degree of purity of the active substance evaluated. The active substance in the product placed on the market can be of equal or different purity if it has been proven to be technically equivalent to the evaluated active substance.

b Regulation (EC) No 1935/2004 of the European Parliament and of the Council of 27 October 2004 on materials and articles intended to come into contact with food and repealing Directives 80/590/EEC and 89/109/EEC (OJ L 338, 13.11.2004, p. 4).

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			attention
			to
			professional
			users.

The purity indicated in this column was the minimum degree of purity of the active substance evaluated. The active substance in the product placed on the market can be of equal or different purity if it has been proven to be technically equivalent to the evaluated active substance.

b Regulation (EC) No 1935/2004 of the European Parliament and of the Council of 27 October 2004 on materials and articles intended to come into contact with food and repealing Directives 80/590/EEC and 89/109/EEC (OJ L 338, 13.11.2004, p. 4).

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Changes to legislation: There are currently no known outstanding effects for the
Commission Implementing Regulation (EU) 2020/1771. (See end of Document for details)

- **(1)** OJ L 167, 27.6.2012, p. 1.
- (2) 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).
- (3) Biocidal Products Committee Opinions on the application for approval of the active substance reaction mass of peracetic acid (PAA) and peroxyoctanoic acid (POOA); Product type: 2, 3 and 4; ECHA/BPC/242, 243 and 244, adopted on 4 March 2020.

Status:

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Changes to legislation:

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