Justification for the selection of a candidate CoRAP substance

Substance Name (Public Name): Isopentyl p-methoxycinnamate

Chemical Group: Organic

EC Number: 275-702-5

CAS Number: 71617-10-2

Submitted by: UK CA

Published: 20/03/2013

NOTE

This document has been prepared by the evaluating Member State given in the CoRAP update.

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1 IDENTITY OF THE SUBSTANCE

1.1 Name and other identifiers of the substance

Table 1: Substance identity

Public Name:	isopentyl p-methoxycinnamate		
EC number:	275-702-5		
EC name:	isopentyl p-methoxycinnamate		
CAS number (in the EC inventory):	71617-10-2		
CAS number:	71617-10-2		
CAS name:	2-Propenoic acid, 3-(4-methoxyphenyl)-, 3-methylbutyl ester		
IUPAC name:	3-methylbutyl 3-(4-methoxyphenyl)acrylate		
Index number in Annex VI of the CLP Regulation	Not listed		
Molecular formula:	C ₁₅ H ₂₀ O ₃		
Molecular weight or molecular weight range:	248.32		
Synonyms:	Neo Heliopan® Galanga Neo Heliopan® E1000 Isoamyl p-methoxycinnamate		

Type of substance		☐ Multi-constituent	
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Structural formula:

2 CLASSIFICATION AND LABELLING

2.1 Harmonised Classification in Annex VI of the CLP

None listed.

2.2 Proposal for Harmonised Classification in Annex VI of the CLP

None proposed.

2.3 Self classification

According to CLP:

Aquatic Acute 1 H400: Very toxic to aquatic life.

According to DSD:

N; R50/53 Dangerous for the environment; Very toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment.

In addition the following has been notified in the Classification and labelling inventory on the ECHA website,

Aquatic Acute 1; H410: Very toxic to aquatic life with long lasting effects.

3 JUSTIFICATION FOR THE SELECTION OF THE CANDIDATE CORAP SUBSTANCE

SOBSTANCE							
3.1 Legal basis for the	oroposal						
\boxtimes Article 44(1) (refined prioritisation criteria for substance evaluation)							
☐ Article 45(5) (Member	State priority)						
3.2 Grounds for concern	n						
☐ (Suspected) CMR	☑ Wide dispersive use	☐ Cumulative exposure					
☐ (Suspected) Sensitiser	⊠ Consumer use	☐ High RCR					
☐ (Suspected) PBT	☐ Exposure of sensitive populations	☐ Aggregated tonnage					
Suspected endocrine disruptor ■	☐ Other (provide further details belo	pw)					
ethylhexyl 4-methoxycinnamate grouping approach. 2-ethylhexyl 4-methoxycinnamate registration; no chronic fish data (48h) of 0.28 mg/l (no chronic s NOEC 0.06 mg/l). Based on the One ready test is available (OEC exposure. Therefore the substar The log Kow is 4.78 (water solutor).	te should be considered for assess for potential environmental endocate is used as a read across for acute are available. Acute toxicity in dastudy is available). The EC50 in algoravailable data the screening criter CD 301F), which showed 70-80% date does not meet screening P and bility 0.8 mg/l), therefore the screen to be a PBT, but should be considered to be a PBT, but should	te fish toxicity in the phnia is high, with an EC50 ae is also high (0.1 mg/l; ion for T is not met. egradation after 28 d vP criteria. ening B criterion is met. ered for investigation of ED					
3.3 Information on agg	regated tonnage and uses						
☐ 1 - 10 tpa	☐ 10 - 100 tpa	☑ 100 – 1000 tpa					
☐ 1000 - 10,000 tpa	☐ 10,000 - 100,000 tpa						
☐ 100,000 - 1000,000 tpa	☐ > 1000,000 tpa						
☐ Confidential							

 $oxed{\boxtimes}$ Industrial use

□ Professional use

☐ Closed System

JUSTIFICATION DOCUMENT FOR THE SELECTION OF A CORAP SUBSTANCE

Industrial use: Manufacture and formulation of cosmetic products.							
Professional use: Formulation of cosmetic products.							
Consumer use: End use of cosmetics							
		_					
-	pleted/ongoing	_		sses that may affect			
Suitability	ioi substance ev	aiua	LIOII				
☐ Compliance check fi	inal decision		☐ Dangerous su	ubstances Directive 67/548/EEC			
☐ Testing proposal			☐ Existing Substances Regulation 793/93/EEC				
☐ Annex VI (CLP)			☐ Plant Protect	ion Products Regulation 91/414/EEC			
☐ Annex XV (SVHC)			☐ Biocidal Prod	ucts Directive 98/8/EEC			
☐ Annex XIV (Authoris	sation)		☐ Other (provide further details below)				
☐ Annex XVII (Restric	tion)						
None that we are aw	are of.						
3.5 Informatio	n to be requeste	d to	clarify the s	uspected risk			
☐ Information on toxic	cological properties		☐ Information on physico-chemical properties				
☐ Information on fate	and behaviour		☐ Information on exposure				
	oxicological properties		☐ Information on uses				
☐ Other (provide furth	ner details below)						
· ·	·						
Information to clarify required.	y the endocrine disrup	tion p	otential of this o	group of substances may be			
required.							
2 6 Detential fo	alland on and link		iek managa				
3.6 Potential follow-up and link to risk management							
Restriction	☐ Harmonised C&L	☐ Au	uthorisation	☐ Other (provide further details)			
This will depend on t	This will depend on the outcome of the evaluation.						