



Justification Document for the Selection of a CoRAP Substance

-UPDATE-

Substance Name (public name): 1,4-diisopropylbenzene

EC Number: 202-826-9

CAS Number: 100-18-5

Authority: France (formerly BG MSCA)

Date: 22/03/2016

22/03/2022 (1. update)

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

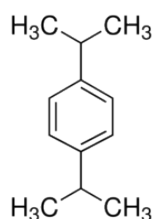
1.1 Other identifiers of the substance

Table 1 : Other Substance identifiers

EC name (public):	1,4-diisopropylbenzene
IUPAC name (public):	1,4-di(propan-2-yl)benzene
Index number in Annex VI of the CLP Regulation:	--
Molecular formula:	C₁₂H₁₈
Molecular weight or molecular weight range:	162.27
Synonyms:	p-Diisopropylbenzene

Type of substance Mono-constituent Multi-constituent UVCB

Structural formula:



Other relevant information about substance composition

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1.2 Similar substances/grouping possibilities

The diisopropylbenzene (DIPB EC 905-459-9), meta-DIPB (EC 202-773-1) and para-DIPB (EC 202-826-9) belongs to a group of three similar substances. The two first members of the group, meta-DIPB (EC 202-773-1) and para-DIPB (EC 202-826-9), are pure isomers while the third member (DIPB) is a reaction mass of the meta- and para-DIPB isomers. France intends to assess the three substances in parallel.

DIPB may contain small amounts of cumene and other aromatic hydrocarbon impurities¹. The three substances, two isomers and the reaction mass are obviously very similar from a structural standpoint as they are all isomers of the same compound and possess nearly identical physical-chemical properties; it has

¹ HPV challenge program, diisopropylbenzene (DIPB) category, test plan, October 3, 2002.

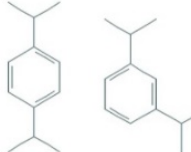
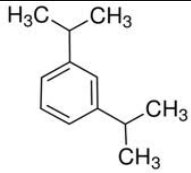
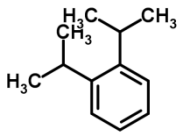
JUSTIFICATION DOCUMENT FOR THE SELECTION OF A CORAP SUBSTANCE

been considered within the HPV Program assessment that data from studies conducted on the mixture itself (DIPB) and each of the individual isomers could be used interchangeably in the evaluation of their environmental fate, ecotoxicity, and mammalian toxicity potentials.

The substance diisopropylbenzene (DIPB), which is a reaction mass of meta-DIPB and para-DIPB was manually screened by France on 27 May 2014 and was then included in the CoRAP.

The registered substance 1,4-diisopropylbenzene (para-DIPB, EC 202-826-9) is the substance of interest for this justification document and is structurally similar to 1,3-diisopropylbenzene (meta-DIPB, EC 202-773-1) and diisopropylbenzene (DIPB) (EC 905-459-9).

Table 2: Similar substances, category approach

EC name	EC and CAS numbers	Structural formula	Molecular formula	Molecular weight
Diisopropylbenzene (DIPB, mixture of para- and meta-DIPB)	EC: 905-459-9		C ₁₂ H ₁₈	162,27
1,3-diisopropylbenzene (meta-DIPB)	EC: 202-773-1 CAS: 99-62-7		C ₁₂ H ₁₈	162,27
1,2-bis(1-methylethyl)benzene (ortho-DIPB)	EC: 209-412-7 CAS: 577-55-9		C ₁₂ H ₁₈	162,27

2 OVERVIEW OF OTHER PROCESSES / EU LEGISLATION

Table 3: Completed or ongoing processes

RMOA	<input type="checkbox"/> Risk Management Option Analysis (RMOA)	
REACH Processes	Evaluation	<input checked="" type="checkbox"/> Compliance check, Final decision
		<input checked="" type="checkbox"/> Testing proposal
		<input type="checkbox"/> CoRAP and Substance Evaluation
	Authorisation	<input type="checkbox"/> Candidate List
		<input type="checkbox"/> Annex XIV
Restri- -ction	<input type="checkbox"/> Annex XVII ²	
Harmonise d C&L	<input type="checkbox"/> Annex VI (CLP) (see section 3.1)	
Processes under other EU legislation	<input type="checkbox"/> Plant Protection Products Regulation Regulation (EC) No 1107/2009	
	<input type="checkbox"/> Biocidal Product Regulation Regulation (EU) 528/2012 and amendments	
Previous legislation	<input type="checkbox"/> Dangerous substances Directive Directive 67/548/EEC (NONS)	
	<input type="checkbox"/> Existing Substances Regulation Regulation 793/93/EEC (RAR/RRS)	
(UNEP) Stockholm convention (POPs Protocol)	<input type="checkbox"/> Assessment	
	<input type="checkbox"/> In relevant Annex	

² Please specify the relevant entry.

Other processes / EU legislation	<input type="checkbox"/> Other (provide further details below)
<p>A CCH was performed for the substance and is considered as concluded. Some additional studies were provided following a TPE.</p>	

3 HAZARD INFORMATION (INCLUDING CLASSIFICATION)

3.1 Classification

3.1.1 Harmonised Classification in Annex VI of the CLP

There is no harmonised classification of the substance under Annex VI of the CLP.

3.1.2 Self classification

- In the registration:

Registrant self-classifies the substance as:

Skin Irrit. 2, H315 Causes skin irritation.

- The following hazard classes are in addition notified among the aggregated self classifications in the C&L Inventory:

Acute Tox. 4,	H302: Harmful if swallowed.
Acute Tox. 4,	H312: Harmful in contact with skin.
Skin Irrit. 2,	H315 Causes skin irritation.
Eye Irrit. 2	H319: Causes serious eye irritation.
Acute Tox. 4,	H332: Harmful if inhaled.
STOT SE 3,	H335: May cause respiratory irritation.
Repr. 2,	H361: Suspected of damaging fertility or the unborn child.
Aquatic Acute 1,	H400: Very toxic to aquatic life
Aquatic Chronic 1,	H410: Very toxic to aquatic life with long lasting effects.
Aquatic Chronic 4,	H413: May cause long lasting harmful effects to aquatic life.

3.1.3 Proposal for Harmonised Classification in Annex VI of the CLP

Not relevant

4 INFORMATION ON (AGGREGATED) TONNAGE AND USES³

4.1 Tonnage and registration status

Table 4: Tonnage and registration status

From ECHA dissemination site*		
<input checked="" type="checkbox"/> Full registration(s) (Art. 10)	<input type="checkbox"/> Intermediate registration(s) (Art. 17 and/or 18)	
Tonnage band (as per dissemination site)		
<input checked="" type="checkbox"/> 1 – 10 tpa	<input type="checkbox"/> 10 – 100 tpa	<input type="checkbox"/> 100 – 1000 tpa
<input type="checkbox"/> 1000 – 10,000 tpa	<input type="checkbox"/> 10,000 – 100,000 tpa	<input type="checkbox"/> 100,000 – 1,000,000 tpa
<input type="checkbox"/> 1,000,000 – 10,000,000 tpa	<input type="checkbox"/> 10,000,000 – 100,000,000 tpa	<input type="checkbox"/> > 100,000,000 tpa
<input type="checkbox"/> <1 >+ tpa (e.g. 10+ ; 100+ ; 10,000+ tpa)		<input type="checkbox"/> Confidential
Joint Submission.		
*the total tonnage band has been calculated by excluding the intermediate uses, for details see the Manual for Dissemination and Confidentiality under REACH Regulation (section 2.6.11): https://echa.europa.eu/documents/10162/22308542/manual_dissemination_en.pdf/7e0b87c2-2681-4380-8389-cd655569d9f0		

4.2 Overview of uses

Table: UsesPart 1:

<input checked="" type="checkbox"/> Manufacture	<input checked="" type="checkbox"/> Formulation	<input checked="" type="checkbox"/> Industrial use	<input checked="" type="checkbox"/> Professional use	<input type="checkbox"/> Consumer use	<input type="checkbox"/> Article service life	<input checked="" type="checkbox"/> Closed system
<p>The identified uses of the substance are:</p> <p><u>Uses at industrial sites:</u></p> <ul style="list-style-type: none"> Industrial uses as Process Solvent for Print Inks Use as an Intermediate <p><u>Uses by Professional Workers:</u></p> <p>Professional Laboratory Use</p>						

³ The dissemination site was accessed Nov 2021

5. JUSTIFICATION FOR THE SELECTION OF THE CANDIDATE CoRAP SUBSTANCE

5.1. Legal basis for the proposal

- Article 44(2) (refined prioritisation criteria for substance evaluation)
- Article 45(5) (Member State priority)

5.2. Selection criteria met (why the substance qualifies for being in CoRAP)

- Fulfils criteria as CMR/ Suspected CMR
- Fulfils criteria as Sensitiser/ Suspected sensitiser
- Fulfils criteria as potential endocrine disrupter
- Fulfils criteria as PBT/vPvB / Suspected PBT/vPvB
- Fulfils criteria high (aggregated) tonnage (*tpa* > 1000)
- Fulfils exposure criteria
- Fulfils MS's (national) priorities

5.3 Initial grounds for concern to be clarified under Substance Evaluation

Hazard based concerns

CMR <input type="checkbox"/> C <input type="checkbox"/> M <input type="checkbox"/> R	Suspected CMR <input type="checkbox"/> C <input type="checkbox"/> M <input checked="" type="checkbox"/> R	<input type="checkbox"/> Potential endocrine disruptor
<input type="checkbox"/> Sensitiser	<input type="checkbox"/> Suspected Sensitiser	
<input type="checkbox"/> PBT/vPvB	<input checked="" type="checkbox"/> Suspected PBT/vPvB ¹	<input checked="" type="checkbox"/> Other (please specify below)

Exposure/risk based concerns

<input type="checkbox"/> Wide dispersive use	<input type="checkbox"/> Consumer use	<input type="checkbox"/> Exposure of sensitive populations
<input checked="" type="checkbox"/> Exposure of environment	<input checked="" type="checkbox"/> Exposure of workers	<input type="checkbox"/> Cumulative exposure
<input type="checkbox"/> High RCR	<input type="checkbox"/> High (aggregated) tonnage	<input type="checkbox"/> Other (please specify below)

Regarding the suspected PBT/vPvB concern

The registered substance is not readily degradable according to the available data in the dossier. However, the biodegradation data are considered insufficient and not fully satisfactory to assess P/vP properties. The registrant stated that no conclusion can be reached based on

available information; however no indication of a testing proposal is provided in the dossier. Based on estimated and experimental data, the substance fulfils the screening criteria for P and leaves the potential for vP. There is a lack of data to fully assess P or vP. Further assessment is considered needed on the P/vP criterion.

Based on the provided experimental Log Kow of 5.23, the substance fulfils the B criterion on screening. The substance is considered stable in the aquatic compartment. Based on a provided read-across bioaccumulation study, the B criterion is clearly fulfilled (with $BCF > 2000 < 5000$) on screening. No bioaccumulation, biota-sediment accumulation, and biomagnification factors (BAF, BSAF, BMF) are provided in the registration dossier. Based on the bioaccumulation estimated data for BCF, the substance indicate a potential for vB (> 5000). Therefore, the substance clearly fulfills the B criterion and is potentially vB on screening based on experimental and estimated data. Further assessment is considered needed on the B/vB criterion.

The substance is presented by the registrant to not fulfill the T criterion, but further information is necessary to conclude on the T properties in the context of the PBT assessment. Based on the estimated chronic aquatic toxicity data, the substance is to be considered as fulfilling the T criterion on screening. Depending on the P/vP and B/vB outcome, the aquatic chronic toxicity could be further investigated. Moreover the notifications of classification as aquatic chronic 1 (H410) should be further assessed.

Regarding the Suspected Reproductive Toxicity concern

In a study on development performed with the m-DiPB (EC 202-773-1) some effects were observed that need to be further investigated. Additionally a notification of classification as Repr. Cat. 2 for the substance and for the m-DiPB are available. . In addition, if reproductive toxicity is verified, workplace exposure scenarios and risk characterization taking into account Council Directive 92/85/EEC and Directive 98/24/EC („Chemical Agents Directive“) would be needed.

Others hazards:

Some concerns (irritation, acute toxicity, effects on liver and kidneys) were identified for the structurally similar substance DiPB (EC 905-459-9) that need to be clarified for this substance also, since DiPB is a reaction mass of two isomers including p-DiPB.

References:

1. EPIWEB 4.1 (US EPA, Nov. 2012). Estimation Programs Interface Suite™ for Microsoft® Windows, v 4.11 or insert version used]. United States Environmental Protection Agency, Washington, DC, USA.
2. PBT profiler (<http://www.pbtprofiler.net/>): Developed by the Environmental Health Analysis Center under contract to the Office of Chemical Safety and Pollution Prevention , U.S. Environmental Protection Agency Computer Resources Donated by SRC, Inc. Ver 2.000 Last Updated September 4, 2012.
3. ECOSAR™ estimation program (<http://www.pbtprofiler.net/ecosarres.asp?I=0&K=4.905>), ECOSAR Version 1.11.
4. www.echemportal.org, OECD SIDS INITIAL ASSESSMENT PROFILE of 1,4-diisopropylbenzene.
5. <http://webnet.oecd.org/CCRWEB/ChemicalDetails.aspx?Key=567a7cdf-4925-4ae3-90da-c8c35a211a30&Idx=0>

5.4 Preliminary indication of information that may need to be requested to clarify the concern

<input checked="" type="checkbox"/> Information on toxicological properties	<input type="checkbox"/> Information on physico-chemical properties
<input checked="" type="checkbox"/> Information on fate and behaviour	<input checked="" type="checkbox"/> Information on exposure
<input checked="" type="checkbox"/> Information on ecotoxicological properties	<input type="checkbox"/> Information on uses
<input type="checkbox"/> Information ED potential	<input type="checkbox"/> Other (provide further details below)

Suspected PBT or vP/vB concern

Additional information may be required to conclude on PBT or vP/vB properties.

Depending on the P/vP and B/vB outcome, the aquatic chronic toxicity (on other species than aquatic invertebrates) might need to be investigated further to conclude on the T properties in the context of the PBT assessment.

Suspected Reproductive Toxicity concern

Some developmental effects are indicated in the provided studies but not yet considered by the registrant, occurring at levels with maternal toxicity. The provided conclusions and explanations are not fully satisfactory and need to be further evaluated. New data may be needed.

5.5 Potential follow-up and link to risk management

<input checked="" type="checkbox"/> Harmonised C&L	<input type="checkbox"/> Restriction	<input type="checkbox"/> Authorisation	<input type="checkbox"/> Other (provide further details)
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Depending on the outcome of the evaluation the substance may result in harmonized classification and labelling and if PBT properties will be confirmed in identification of the substance as SVHC