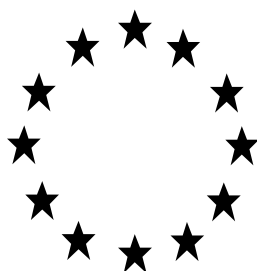


**Regulation (EU) No 528/2012  
concerning the making available on the  
market and use of biocidal products**

*Evaluation of active substances*

Assessment Report



**Polyhexamethylene biguanide  
(Mn = 1600; PDI = 1.8)  
(PHMB)**

Product-type 1: Human hygiene

June 2015

France

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## **1 STATEMENT OF SUBJECT MATTER AND PURPOSE**

### **1.1 PROCEDURE FOLLOWED**

This Assessment Report has been established as a result of the evaluation of the active substance Polyhexamethylene biguanide with a mean number-average molecular weight (Mn) of 1600 and a mean polydispersity (PDI) of 1.8, i.e. PHMB (1600; 1.8), as product-type 1 (human hygiene), carried out in the context of the work programme for the review of existing active substances provided for in Article 89 of Regulation (EU) No 528/2012 concerning the making available on the market and use of biocidal products<sup>1</sup>, with a view to the possible approval of this substance.

PHMB (1600; 1.8) (CAS no. 27083-27-8 and 32289-58-0) was notified as an existing active substance, by Lonza (previously Arch Chemicals Ltd.), hereafter referred to as the applicant, in product-type 1.

Commission Regulation (EC) No 1451/2007 of the 4<sup>th</sup> of December 2007<sup>2</sup> lays down the detailed rules for the evaluation of dossiers and for the decision-making process.

In accordance with the provisions of Article 3 paragraph 2 of that Regulation, France was designated as Rapporteur Member State (RMS, hereafter referred to as the evaluating Competent Authority, eCA) to carry out the assessment on the basis of the dossier submitted by the applicant. The deadline for submission of a complete dossier for PHMB (1600; 1.8) as an active substance in product-type 1 was the 31<sup>st</sup> of July 2007, in accordance with Article 9 paragraph 2 of Regulation (EC) No 1451/2007.

On the 30<sup>th</sup> of July 2007, the French Competent Authority received a dossier from Lonza. The evaluating Competent Authority accepted the dossier as complete for the purpose of the evaluation, taking into account the supported uses, and confirmed the acceptance of the dossier on the 18<sup>th</sup> of February 2008.

On the 5<sup>th</sup> of September 2013, the evaluating Competent Authority submitted to European Chemical Agency (ECHA), hereafter referred to as the Agency, and the applicant a copy of the evaluation report, hereafter referred to as the Competent Authority Report.

In order to review the Competent Authority Report and the comments received on it, consultations of technical experts from all Member States (peer review) were organised by the Agency. Revisions agreed upon were presented at the Biocidal Products Committee and its Working Groups meetings and the Competent Authority Report was amended accordingly.

### **1.2 PURPOSE OF THE ASSESSMENT**

The aim of the Assessment Report is to support a decision on the approval of PHMB (1600; 1.8) for product-type 1, and should it be approved, to facilitate the authorisation of individual biocidal products in product-type 1 that contain PHMB (1600; 1.8). In the evaluation of applications for product-authorisation, the provisions of Regulation (EU) No 528/2012 shall be applied, in particular the provisions of Chapter IV, as well as the common principles laid down in Annex VI.

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<sup>1</sup> 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, OJ L 167/1, 27.6.2012, p1.

<sup>2</sup> OJ L 325, 11.12.2007, p. 3

The conclusions of this report were reached within the framework of the uses that were proposed and supported by the applicant (see Appendix II). For the implementation of the common principles of Annex VI, the content and conclusions of this assessment report shall be taken into account.

However, where conclusions of this assessment report are based on data protected under the provisions of Regulation (EU) No 528/2012, such conclusions may not be used to the benefit of another applicant, unless access to these data has been granted.

## 2 OVERALL SUMMARY AND CONCLUSIONS

### 2.1 GENERAL SUBSTANCE INFORMATION / GENERAL PRODUCT INFORMATION

#### 2.1.1 IDENTITY, PHYSICO-CHEMICAL PROPERTIES & METHODS OF ANALYSIS OF THE ACTIVE SUBSTANCE

##### 2.1.1.1 Identity

**Table 2.1-1: Identification of the active substance**

CAS-No.	CAS-No : 27083-27-8 and 32289-58-0  It must be noted that CAS number 27083-27-8 is not based on characterisation data. In case of a different PHMB (for example with a weigh distribution outside of the specification of the PHMB assessed in this report) the CAS number will not be able to differentiate the PHMB.																
EINECS-No.	Not listed on the EU (EINECS) inventory because PHMB is a polymer. Polymers are exempt from listing on EINECS if the monomers are listed.																
Other No. (CIPAC, ELINCS)	None.																
IUPAC Name	CoPoly(bisiminoimidocarbonyl,hexamethylenehydrochloride),(iminoimidocarbonyl , hexaméthylène hydrochloride) or Copoly(5-imino-7-imino-4,6,8-triazaundecamethylene hydrochloride) (5-imino-4,6-diazanonamethylenehydrochloride)																
Common name, synonym	PHMB (1600; 1.8) i.e. polyhexamethylene biguanide with a mean number-average molecular weight (Mn) of 1600 and a mean polydispersity (PDI) of 1.8; Polyhexamethylene biguanide; Poly(hexamethylene) biguanide hydrochloride; polymeric biguanide hydrochloride; "PHMB"; Polyhexanide (International non-proprietary name); Polyaminopropyl Biguanide (INCI)																
Molecular formula	Terminal function- (CH <sub>2</sub> ) <sub>6</sub> - [C <sub>8</sub> H <sub>18</sub> N <sub>5</sub> Cl] <sub>n</sub> [C <sub>7</sub> H <sub>16</sub> N <sub>3</sub> Cl] <sub>m</sub> - terminal function  Possible terminal functions: - NH <sub>2</sub> (amine) - C <sub>2</sub> H <sub>3</sub> N <sub>4</sub> ( cyanoguanide) - CH <sub>5</sub> N <sub>3</sub> Cl (guanidine)																
		<table border="1"> <thead> <tr> <th></th> <th>range</th> <th>average</th> </tr> </thead> <tbody> <tr> <td>m+n</td> <td>2-40</td> <td>11</td> </tr> <tr> <td>n/(m+n) [biguanide %]</td> <td>90.8 - 91.9%</td> <td>91.3 %</td> </tr> <tr> <td>m/(m+n) [guanide %]</td> <td>8.1 - 9.2 %</td> <td>8.6 %</td> </tr> <tr> <td>↳ ω ↳ amino</td> <td>35% - 46%</td> <td>39%</td> </tr> </tbody> </table>		range	average	m+n	2-40	11	n/(m+n) [biguanide %]	90.8 - 91.9%	91.3 %	m/(m+n) [guanide %]	8.1 - 9.2 %	8.6 %	↳ ω ↳ amino	35% - 46%	39%
	range	average															
m+n	2-40	11															
n/(m+n) [biguanide %]	90.8 - 91.9%	91.3 %															
m/(m+n) [guanide %]	8.1 - 9.2 %	8.6 %															
↳ ω ↳ amino	35% - 46%	39%															

		guanidine	22% - 29%	25%
		cyanoguanide	31 - 39%	35%
Structural formula				
Molecular weight	Number average molecular weight (Mn) = 1610 Mass average molecular weight (Mw) = 2986.			

The active ingredient (a.i.) Poly Hexa Methylene Biguanide (PHMB) is a small size polymer obtained by the polycondensation of two monomers (1,6-hexanemethylenediamine and N,N''-1,6-hexanediybis[N'-cyanoguanidine] (ie. HMBDA)).

As PHMB is a small size polymer, some side reactions that occurred during the manufacturing process could modify significantly the structure of the polymer. The side reaction to obtain the unit guanidine occurred up to 10% in the process. Therefore, it can be considered that the structure of PHMB is not only composed by repetitive unit of guanidine but it is composed by repetitive unit of guanidine and biguanide.

The active substance as manufactured (TK<sup>3</sup>) is a 20% w/w aqueous solution of PHMB. "Purity" is a difficult concept to apply to PHMB which is a mixture of polymers and related substances. Instead, the applicant refers to the "strength" of the polymer which is defined as "% total solids" or "dried material". The typical PHMB strength is 20 %.

However, eCA considers more appropriate to use the term "% of active substance (% a.s.)" or "active substance content" instead of "strength". The active substance content being defined as the sum of PHMB and its impurities contents, it can be considered identical to the % total solids and thus to the strength. However, the terms strength or dried PHMB are also used in identity and physico chemical sections and refer to the same thing.

As the technical material is the 20 % PHMB solution obtained directly from the manufacturing process (active substance as manufactured or TK), characterisation data were generated from the dried technical material (TC<sup>4</sup>) using the technique of freeze drying.

The content of PHMB can be calculated by subtracting the total content of impurities in the dried technical material (without residual water) to 100. This value cannot be considered as a real purity but is the closest available data.

The minimum content of PHMB TC was demonstrated > 95.6%.

Since the active substance is a copolymer, identity characterisation criteria (based on % solid, content of PHMB in dried material, Mw, Mn and the biguanide/guanide ratio) as well as limits or range for each criterion are proposed by eCA in the confidential document IIA to characterise the source of PHMB in order to set reference specifications in case of approval of the active substance and future technical equivalence checks. **eCA proposes to rename PHMB considered for approval in this dossier as "PHMB with a mean number-average molecular weight (Mn) of 1600 and a mean polydispersity (PDI) of 1.8"**

<sup>3</sup> TK: technical concentrate according to GIFAP monograph n°2 nomenclature.

<sup>4</sup> TC: technical material according to GIFAP monograph n°2 nomenclature.

i.e. "PHMB (1600; 1.8)". For convenience, PHMB (1600; 1.8) is referred to hereafter as "PHMB" or "a.s."

There is one relevant impurity, Hexamethylenediamine with a maximal content of 0.4%. All potential impurities have not been looked for and/or quantified. Additional data about impurities and specifications for the active substance and the impurities should be submitted prior to approval.

Quality control data on structural characteristics (2003-2011) are reported in this confidential document to demonstrate that production of TK (liquid form) remained stable during this period of time from a structural point of view. It can be concluded that submitted characterisation data (2011) are representative of current production but also of older production and of active substance material used to perform the toxicological and ecotoxicological studies used to perform the risk assessment (See confidential doc IIA). This statement is only valid for structural data and not for evolution of impurity content in PHMB as no data was submitted to cover this point.

The applicant also manufactures PHMB as a solid material ("Solid PHMB"). Initially the applicant submitted both sources in the dossier. Comparison between liquid and Solid PHMB is discussed in confidential document IIA-02 "Comparison of liquid and solid PHMB". eCA considers that liquid PHMB (VANTOCIL TG) and Solid PHMB are 2 different substances, based on structural considerations. Additional information to demonstrate technical equivalence will be required at product authorisation stage if Applicant claims solid PHMB as a new source. The active substance considered for approval in this dossier is the active substance as manufactured (TK): 20 % w/w aqueous solution of PHMB (VANTOCIL TG) also called liquid PHMB.

#### **Summary of specifications of Lonza PHMB:**

Complete specifications are available in confidential part. The summary is reported here.

- Specifications set by eCA:

**Table 2.1-2: Specifications of PHMB (1600; 1.8) from Lonza**

<b>Characterisation specification</b>	
Strength	18-22%
PHMB in dried material	≥ 95.6%
molecular weight by number (Mn)	1449-1771
molecular weight by mass (Mw)	2687-3285
Polydispersity	1.80-1.91
The biguanide / guanide ratio in chain	90/10 to 92/8
Total fraction <1000 Da	16.6-24.5 %
<b>Impurities</b>	
HMD (relevant impurity)	≤ 0.4%
Other impurities	confidential



- (eco)tox batches: Liquid PHMB used to perform (eco)toxicological key studies and efficacy studies is of the same structure than liquid PHMB characterised in this dossier. However, no data on (eco)toxicity of impurities was provided by the applicant. Additional data about (eco)toxicity of impurities should be submitted for finalisation of specification.
- Criterion data to be used to differentiate PHMB from different origins: All of presented characterisation data are important to differentiate PHMB assessed in this dossier and other PHMB. However, some of those criterion data could be found difficult for control (biguanide / guanide ratio quantified by NMR) or not selective (strength). eCA is of the opinion that Mn and polydispersity would be the most convenient property for the control of the identity of PHMB used in biocidal products.

### **2.1.1.2 Physico-chemical properties**

TC (dried PHMB) is a dusty solid/powder, off white with a strong ammonia smell. It has a glass transition temperature of 90-91°C (non crystalline polymer) and decomposes at 205-210°C before boiling. The TK (PHMB as manufactured, 20% in water) has a boiling point of 100.2°C. The relative density of TC is 1.20 at 20°C and the relative density of the TK is 1.04 at 20°C. As a polymer, PHMB is not considered to be volatile. Henry's Law Constant is not applicable as PHMB is not considered to be volatile and is present in ionic form at neutral pH. It is assumed that PHMB has only slight possibility to go from water to air. It is very soluble in water (426 g/L). It is also soluble in methanol (41%), in ethanol (0.5%) and sparingly soluble in organic solvents (10-3 g/L). The pKa is calculated as approximately 4.4 at 25°C. Log Pow is -2.3 at pH=7.4 and 25°C. TC is not highly flammable, and does not have oxidizing and explosive properties. A surface tension study should be performed but PHMB is not expected to be surface active based on structural considerations.

### **2.1.1.3 Methods of analysis**

It is impossible to determine directly PHMB since it is not a single chemical entity but a polymeric mixture with a range of molecular weight. Adequate methodology exists for the characterisation of the active ingredient and the determination of the known impurities in TC but more validation data are required.

Justifications for non submission of analytical methods for residues of the active substance in soil, water, air and body fluids and tissues, in food or feedstuffs were submitted.

For polymeric substances it may be difficult to develop an adequate residue analytical method. A limited residue definition in form of a marker will be required if PHMB is proposed for approval.

Residue definition: a proposal of residue definition for drinking water, body fluid and tissues and food and feeding stuff is required 6 months before the date of approval.

Monitoring methods:

- Based on the bibliography and the nature of the active ingredient, determination of PHMB in soil is currently not technically feasible. Moreover, eCA considers that if a method could allow to quantify PHMB in soil, this method could probably not be considered as enforcement method.
- The non submission is acceptable for air because occurrence in air is not probable.

- The non submission is acceptable for surface water, as eCA considers that the issue is the same than in soil. However, determination of PHMB in drinking water should be technically feasible. Therefore, a validated method for determination of PHMB would be required.
- The justification for non submission submitted by the applicant is not acceptable for body fluids and tissues as PHMB is classified as very toxic. An analytical method for determination of PHMB in body fluids and tissues or another justification of non submission of data would be required.
- The justification for non submission submitted by the applicant is not acceptable for food and feeding stuff as the justification based on the non exposure of food or feedstuffs is not acceptable, considering that PHMB can be used by operatives in the food processing industry (refer to Document IIIB5). Methods for the determination of PHMB and residues in food and feedstuffs would be required.

## **2.1.2 IDENTITY, PHYSICO-CHEMICAL PROPERTIES & METHODS OF ANALYSIS OF THE BIOCIDAL PRODUCT**

### **2.1.2.1 Identity**

**Table 2.1-3: Identification of the biocidal product**

<b>Trade name</b>	PHMB HANDWASH	
Manufacturer's development code number(s)		
<b>Ingredient of preparation</b>	<b>Function</b>	<b>Content</b> (strength % w/w)
PHMB	Active substance	0.9
Physical state of preparation	Liquid	
Nature of preparation	AL: Other liquids to be applied undiluted	

### **2.1.2.2 Physico-chemical properties**

Applicant claimed that PHMB HANDWASH is a "dummy" product. It is not a commercially available product and as such has not been tested for any physicochemical properties.

eCA considers that a complete dossier is not necessary but a minimum of information should be provided, even if based on theoretical considerations. Complete data on physico-chemical properties should have been submitted and would be required at product authorisation stage.

eCA has estimated some properties from the composition of PHMB HANDWASH. According to the composition of biocidal product:

- No explosive properties are expected for PHMB HANDWASH.
- No oxidising properties are expected for PHMB HANDWASH.
- PHMB HANDWASH is not expected to be highly flammable and is not expected to have a autoflammability point at ambient temperature

Data on other properties would be required at product authorisation stage: appearance, acidity/alkalinity, relative density, accelerated storage stability, long term stability and low temperature stability, reactivity towards container material (as PHMB is known to adsorb on surfaces, a measure of PHMB adsorbed on container after storage would be required), persistent foaming tested on pure material , surface tension and viscosity.

### **2.1.2.3 Methods of analysis**

Methods have not been validated for this specific product because PHMB HANDWASH is a "dummy" product.

Validated analytical methods would be required at product authorisation stage.

## **2.1.3 INTENDED USES AND EFFICACY**

### **2.1.3.1 Field of use envisaged**

This Product Type 01 dossier for PHMB is provided to support the following use:

<i>MG01:</i>	Disinfectants.
<i>Product Type 01:</i>	Human hygiene.
<i>Further specification:</i>	<i>Skin, hair and mucous membranes.</i>

### **2.1.3.2 Function**

Bactericide.

Claims of yeasticidal and virucidal activities have been withdrawn by the applicant respectively on 05/06/2009 and on 08/12/2008.

### **2.1.3.3 Mode of action**

The lethal action of PHMB is an irreversible loss of essential cellular components as a direct consequence of cytoplasmic membrane damage. It is concluded that cytoplasmic precipitation is a secondary event to the death of the bacterial cell.

It has been shown that the lethal sequence consists of a series of cytological and physiological changes - some of which are reversible - which culminate in the death of the cell. The important steps are:

- binding to a receptive site on the surface;
- leakage of low molecular weight cytoplasmic components;
- precipitation of cell contents.

The molecular interaction between PHMB and bacterial membranes has been deduced by overlaying this lethal sequence with the findings of experiments modelling the possible interactions of polymeric biguanides and membrane components - particularly phospholipids.

**2.1.3.4 Objects to be protected, target organisms**

The Table 2.1-4 presents the intended use for which efficacy data support the efficacy of the PHMB for approval. The data are generated from laboratory studies and would have to be consolidated at the product authorisation stage in accordance with the claims and with data generated with real products.

**Table 2.1-4: Intended use for which efficacy data support the efficacy of PHMB**

	<b>Applicati on method</b>	<b>Product</b>	<b>In use concentration / contact time</b> (PHMB in the in-use solution) Product density=1	<b>Activity</b>
Human Hygiene	Hand wash	PHMB HANDWASH formulation HT 330 (4.5 % v/v TK, i.e. 0.9 % w/w a.s.)	0.9 % w/w a.s. 30 seconds/3 ml	Bactericidal

**2.1.3.5 Resistance**

The evaluation of the literature studies provided by the applicant does not show particular resistance to PHMB with bacteria. Nevertheless it is not appropriate to conclude that PHMB resistance is not an issue and that a resistance management strategy is not required. In particular, the description in the literature of:

- cross resistances,
- modifications of the expression of genes as a mechanism of tolerance to sublethal concentrations of PHMB,

should be taken into account in the strategy of resistance management.

Standard methods of measuring resistance brought about by biocide use are not available and should be developed for all type of biocides (Assessment of the Antibiotic Resistance Effects of Biocides, Scenihr 2009).

**2.1.4 CLASSIFICATION AND LABELLING**

**2.1.4.1 Proposed classification for the active substance as manufactured (TK):  
PHMB 20% in water**

<b>Classification according to Regulation (EC) No 1272/2008 (CLP)</b>			
Class of danger	Acute Tox 4	Warning	
	Skin Sens 1B	Warning	
	STOT Rep 1	Danger	
	Carc. 2	Warning	
	Aquatic Acute 1	Danger	
	Aquatic Chronic 1	Danger	

Hazard statement	H332	Harmful if inhaled.
	H317	May cause an allergic skin reaction.
	H372	Causes damage to organs through prolonged or repeated exposure by inhalation.
	H351	Suspected of causing cancer.
	H400	Very toxic to aquatic life.
	H410	Very toxic to aquatic life with long lasting effects.

#### 2.1.4.2 Harmonised classification for the active substance: PHMB

Classification according to Regulation (EC) No 1272/2008 (CLP)		
Class of danger	Acute Tox 4	Warning
	Eye dam 1	Danger
	Skin Sens 1B	Warning
	STOT Rep 1	Danger
	Carc. 2	Warning
	Aquatic Acute 1	Danger
	Aquatic Chronic 1	Danger
Hazard statement	H302	Harmful if swallowed.
	H318	Causes serious eye damage.
	H317	May cause an allergic skin reaction.
	H372	Causes damage to organs through prolonged or repeated exposure by inhalation.
	H351	Suspected of causing cancer.
	H400 (M-factor =10)	Very toxic to aquatic life.
	H410 (M-factor =10)	Very toxic to aquatic life with long lasting effects.

A RAC opinion (March 2014) is also available for the acute inhalation toxicity endpoint:

- Acute Tox. 2; H330: Fatal if inhaled.

#### 2.1.4.3 Proposed classification of PHMB HANDWASH

As already mentioned, PHMB HANDWASH is a “dummy” product which is not produced commercially. Therefore, it has never been formally classified according to the EU requirements (Directive 1995/45/EC and 2006/8/EC).

<b>Classification according to Regulation (EC) No 1272/2008 (CLP)</b>		
	Eye dam 2	Danger
	Aquatic chronic 2	Danger
	H319	Causes serious eye damage.
	H411	Toxic to aquatic life, may cause long-term adverse effects in the aquatic environment

## 2.2 SUMMARY OF THE RISK ASSESSMENT

### 2.2.1 SUMMARY OF HUMAN HEALTH RISK ASSESSMENTS

#### 2.2.1.1 Hazard identification

- **Toxicokinetic:**

Oral absorption of PHMB ranges approximately from 0.3 to 8% but the value of 4% is retained based on the oral absorption of PHMB from diet at the lower dose tested. This value was selected as it corresponds to the closest conditions to the experimental conditions of the study in which the relevant oral NOAEL was determined.

A dermal absorption of PHMB was determined to be 4% by default based on EFSA guidance on dermal absorption (2012), corresponding to the oral absorption value.

Since no information is available on absorption of PHMB by inhalation, an absorption of 100% is retained.

- **Acute toxicity:**

A classification for acute oral or dermal toxicity is not justified for the active substance as manufactured, PHMB 20% in water. For respiratory route, a classification Xn; R20 or Acute Tox 4 – H332 is proposed based on the RAC opinion for PHMB.

- **Irritation/Sensitisation:**

PHMB is not irritant by dermal contact. For eye irritation, classification is not justified based on the data of the PHMB 20% w/w. PHMB is considered as a moderate to strong potency skin sensitizer based on animal data. Human studies indicate that PHMB is a skin sensitizer in humans, although with a rare frequency of sensitisation in the current conditions of consumer uses. Classification Xi; R43 (may cause sensitisation by skin contact) or Skin sens 1 – H317 for CLP, is therefore warranted. Relatively low incidences from human data support classification as CLP Skin Sens **1B** – H317 according to the 2nd ATP to CLP Regulation.

- **Repeated toxicity:**

On the basis of the severity of the effects caused by inhalation of PHMB (mortality and to a lesser extent histopathological changes in the respiratory tract and in the thymus), the absence of reversibility of inflammation in the respiratory tract and the very low doses causing these effects, classification T; R48/23 is warranted (CLP STOT RE 1 - H 372). By inhalation the primary target organ is the respiratory tract and no effect warranting classification are identified by oral and dermal route. The target organs are kidneys and liver via oral route. By dermal contact, local effects are expected.

- **Genotoxicity:**

PHMB is not considered to be mutagenic or genotoxic, according to the results of the *in vitro* (Ames test and chromosomal aberration test) and *in vivo* studies (mouse bone marrow micronucleus test and UDS assay).

- **Carcinogenicity:**

PHMB increases the incidence of benign and malign vascular tumours in female rats by oral route and in male and female mice by oral and dermal route. The tumours are induced mainly in the liver, which is one of the target organ of PHMB and the increase is clearly seen at doses above the MTD. However, it is also observed more equivocally at doses below MTD (mouse oral study at mid-dose and rat oral study at high dose). These increases are not considered incidental when considering the clear induction of vascular tumours at higher doses and they are considered biologically significant and attributed to treatment.

A classification as carcinogenic category 3; R40 or Carc 2 – H351 for CLP, is warranted. In absence of carcinogenicity data by inhalation, it is proposed to allocate the general hazard statement H351 without indication of the route of exposure.

- **Reprotoxicity:**

PHMB has no teratogenic effect and has no effect on fertility or reproductive performance at dose levels up to 2000 ppm.

#### **Determination of AEL/AEC/ADI/ARfD**

- Systemic effects

The lowest NOAEL from any oral studies is 13 mg/kg bw/day from the rat developmental toxicity study (Doc IIIA 6.8.1/01). This value is based on reduced maternal food consumption and body weight (-23% of controls) seen at the next higher dose. The choice of this value is also supported by the rabbit developmental toxicity study, in which increased mortality and reduced bodyweight with associated reduced food consumption were seen at the same level of doses.

The absorption rate following administration in the diet for females is 4%. Hence, internal NOAEL is 0.52 mg a.s./kg bw/day

The default assessment factors are 10 for inter-species variation and 10 for intra-species variation in the case of the systemic effects. The inter-species factor consists of 2.5 for toxicodynamic- and 4.0 for toxicokinetic variability, while the inter-individual factor consists of 3.2 for toxicokinetic and 3.2 for toxicodynamic variability.

Although the selected NOAEL is based on a short duration of exposure (22 days in the rat teratogenicity study), no assessment factor will be applied to take into account the medium and chronic exposure because the NOAEL from teratogenicity is in the same order of magnitude or lower than NOAEL from sub-chronic or chronic studies. Consequently, it means that effects are not more severe with longer exposure of PHMB. The NOAEL from teratogenicity is therefore, sufficiently conservative for these longer exposures and no additional assessment factors to extrapolate NOAEL of the teratogenicity study to longer duration is justified.

The MOE<sub>ref</sub> is therefore 100 for acute-term, medium-term and long-term exposure.

**An acute, medium-term and long-term AEL of  $5.2 \times 10^{-3}$  mg a.s./kg bw/day is proposed.**

- Respiratory exposure, local effects

The relevant study for respiratory exposure is the 28-day inhalation study. The NOAEC from this study is 0.024 mg/m<sup>3</sup> (Document IIIA 6.3.3).



The MOE<sub>ref</sub> is therefore 25, 75, 150 for local effects for acute, medium and long-term respiratory exposure.

An acute respiratory AEC of 0.96 µg/m<sup>3</sup> a.s. is proposed.

A medium-term respiratory AEC of 0.32 µg/m<sup>3</sup> a.s. is proposed.

A long-term respiratory AEC of 0.16 µg/m<sup>3</sup> a.s. is proposed.

According to the TNsG on Annex I inclusion, chapter 4.1: quantitative risk characterisation (2008), ADI and ARfD are usually based on the same NOAEL as the AEL<sub>chronic</sub> and AEL<sub>acute</sub> respectively. They are external reference doses.

A value of 0.13 mg/kg is proposed for ADI and ARfD.

**Table 2.2-1: Summary of the values of AEL and MOE<sub>ref</sub>**

<b>Systemic effects</b>		
	<b>AEL</b>	<b>MOE<sub>ref</sub></b>
acute, medium and long-term	5.2 µg a.s./kg bw/d	100
	<b>ADI - ARfD</b>	<b>MOE<sub>ref</sub></b>
Chronic and acute	0.13 mg a.s./kg bw/d	100
<b>Local effects by inhalation</b>		
	<b>AEC</b>	<b>MOE<sub>ref</sub></b>
acute	0.96 µg/m <sup>3</sup>	25
medium-term	0.32 µg/m <sup>3</sup>	75
long-term	0.16 µg/m <sup>3</sup>	150

### **2.2.1.2 Uses and relevant exposure pathways**

PHMB HANDWASH is a dummy product containing 0.9% w/w of active substance. It is presented to illustrate the use of PHMB as the active substance in PT01 hygienic hand wash products. PHMB HANDWASH would be supplied in a container into which is integrated a small finger operated pump. Operating the pump dispenses 3 mL of product directly onto the hand. The hand washing is followed by a rinsing with water.

PHMB HANDWASH would be used primarily by professional users, but potential use by non-professionals in a domestic setting is also considered.

The relevant routes of exposure for the activities involved in the application method are considered in detail in the appropriate Document IIB section 3. A brief overview is given below:

**Primary exposure**

The route of exposure to PHMB during the use of PHMB HANDWASH is the dermal route. Inhalation is generally not a relevant route because the active substance is non-volatile and there are no high shear operations to generate an aerosol.

**Secondary exposure**

There is a potential for secondary exposure by transfer of residues from washed hands following:

- i) hand to mouth contact
- ii) hand to food contact
- iii) hand to skin contact with others

For points (ii) and (iii) potential exposure are negligible compared with the primary exposure of the hand wash user and are therefore not further assessed. However, ingestion of PHMB residues following hand to mouth contact by a child (ie. mouthing of hands after hand washing) has been considered in this dossier.

Again, inhalation is not a relevant route because the active substance is non-volatile and there are no high shear operations to generate an aerosol.

**Table 2.2-2: Summary of main paths of human exposure**

<b>Exposure path</b>	<b>Industrial use</b>	<b>Professional use</b>	<b>Non-professional</b>	<b>General public (secondary exposure)</b>	<b>Via the environment</b>
Inhalation	NA	No	No	No	No
Dermal	NA	Yes	Yes	Yes	No
Oral	NA	No	No	Yes*	No

\* Worst case of child licking fingers after washing with PHMB HANDWASH

**2.2.1.2.1 Primary exposure and risk characterisation for PHMB in Product Type 01 –Professional and non-professional users**

Considering primary exposure, the risk is characterised for the representative example scenario used to support the use of PHMB as PT 1.

Doc IIB/section 3 describes a “worst” case use pattern for PHMB HANDWASH and identifies the route of exposure and available models for calculating exposure.

The worst case scenarios are detailed in the following table.

**Table 2.2-3: Overview of scenarios of PHMB HANDWASH**

Application method	User	Frequency (applications / day)	Volume used per application	Effective In use concentration (% w/v a.s.)	PPE (personal protective equipment)
Container with integral finger pump dispenser	Professional (60 kg)	10	3 ml (3.0 g)	0.9% (9000 ppm)	None (Not relevant)
	Non-professional (Infant of 10 kg)	5			

As already discussed, the route of exposure relevant to the application method is dermal contamination only.

The primary exposure is considered to be chronic.

The predicted exposures for professional and non-professionals users (as described in Document IIB) are presented below. Exposure estimate is compared to the NOAEL and to the AEL, leading to derive a MOE and a fraction of the AEL (expressed as % AEL). The results are presented in the following tables.

→ **Risk characterisation for systemic effects**

The exposure scenario related to the washing process involves dispensing product onto dry hands by operating the pump then rubbing for 1 minute. The scenario includes rinsing of product with water then drying of hands.

For PHMB, the dermal absorption value of 4% is a default value based on oral absorption value, as no acceptable dermal absorption study was provided. Thereby, it is impossible to establish an absorption rate over the time ( $\mu\text{g}/\text{cm}^2/\text{min}$ , i.e. flux parameter). In consequence, it is not possible to estimate the exposure during 1 minute of hand washing, and this step is not assessed for PHMB: only the calculation of exposure and risk after rinse is presented below.

Even if the hand washing phase is short, its contribution to the total exposure (exposure during hand washing phase and exposure of residues after rinse) is assumed to be not negligible. Thus, it should be noted that risk calculated below is underestimated.

**Table 2.2-4: Risk characterisation concerning systemic effects during washing hands- Professionals and non-professionals**

<u>Hand washing - TP01</u>	<b>Systemic exposure</b> (mg a.s./kg bw/d)	Relevant NOAEL (mg a.s./kg bw/d)	MOE <sub>ref</sub> (sum of AFs)	MOE	AEL (mg a.s./kg bw/d)	%AEL
Exposure of residual PHMB after hands washing per day						
Professionals (10 applications)	$1.80 \times 10^{-3}$	0.52	100	289	$5.20 \times 10^{-3}$	34.6
Infant (5 applications)	$5.40 \times 10^{-3}$	0.52	100	96	$5.20 \times 10^{-3}$	<b>103.8</b>



The risk during exposure of residual PHMB after hands washing is considered to be acceptable for professional users with a MOE (289) higher than the MOE ref (100) and a %AEL (35%) below 100%.

As the assessment is based on an underestimated exposure, at product authorisation level a dermal absorption study allowing derivation of flux parameters is mandatory to confirm the assessment.

Even with an underestimated exposure as explained before, the risk is considered to be unacceptable for non-professional users with a MOE (96) lower than the MOE ref (100) and a %AEL (104) above 100%.

→ **Risk characterisation for local dermal effects**

The content of PHMB in the end use product is below classification limits of skin sensitisation. As the product is not classified for its sensitisation and irritations properties, the risk for dermal local effects is considered to be acceptable.

**2.2.1.2.2 Secondary estimated exposures and risk characterisation for PHMB in Product Type 01 –Secondary exposure of infant**

The model calculates an absolute worst case by assuming 100% additivity of skin residues following each washing, removal of residues from area of the hands.

There is a potential for secondary exposure by transfer of residues from washed hands following:

- hand to mouth contact
- hand to food contact (eg. to food in a food processing environment)
- hand to skin contact with others (eg. dermal transfer to patients in a medical care environment)

This section considers the potential for secondary exposure and identifies hand to mouth transfer as the only activity for which an assessment is required.

**Table 2.2-5: Estimation of residues exposure during hand to mouth transfer**

<u>Hand to mouth transfer- TP01</u>	<b>Infant (5 applications)</b>
Quantity of product	$2.59 \times 10^{-1}$ mg/wash
Systemic exposure via ingestion per body weight	$1.04 \times 10^{-4}$ mg/kg bw for 1 wash
	$5.18 \times 10^{-3}$ mg/kg bw/day (5 washes/day)

The predicted exposures for non-professionals users are presented below. Each exposure estimate is compared to the NOAEL and to the AEL, leading to derive a MOE and a fraction of the AEL (expressed as % AEL), respectively.

→ **Risk characterisation for systemic effects**

**Table 2.2-6: Risk characterisation concerning systemic effects during hand to mouth transfer- Secondary exposure, infant**

<u>Hand to mouth transfer- TP01</u>	<b>Total exposure</b> (mg a.s./kg bw/d)	Relevant NOAEL (mg a.s./kg bw/d)	MOE <sub>ref</sub> (sum of AFs)	MOE	AEL (mg a.s./kg bw/d)	%AEL
Infant (5 applications)	5.18 x 10 <sup>-03</sup>	0.52	100	100.3	5.20 x 10 <sup>-3</sup>	99.7

The risk for systemic effects during hand to mouth transfer is considered to be acceptable for infant with a MOE higher than the MOE ref (100) and a %AEL (99.7) below 100%. As a reminder, the risk of systemic effects during primary exposure is unacceptable for children. Therefore, despite a very high percent of AEL, no refinement is performed for secondary exposure.

→ **Risk characterisation for local oral effects**

The content of PHMB in the end use product is below classification limits of skin sensitisation. As the product is not classified for its sensitisation and irritations properties, the risk for dermal local effects is considered to be acceptable.

**2.2.1.2.3 Combined exposure**

The calculation of combined exposure (primary exposure and secondary exposure) is not relevant because the risk for the primary exposure is considered to be unacceptable for the general public.

**2.2.1.3 Overall conclusion for human health**

The risk when considering primary exposure to PHMB HANDWASH is acceptable for professionals and unacceptable for general public.

The risk when considering possible secondary exposure to PHMB HANDWASH is acceptable.

In conclusion, the human health risk assessment for the use of PHMB in product of Product Type 01 as hand disinfection product demonstrates that there are unacceptable risks for the general public and acceptable risks for professionals.

## **2.2.2 SUMMARY OF ENVIRONMENTAL RISK ASSESSMENTS**

### **2.2.2.1 Fate and distribution in the environment**

#### **2.2.2.1.1 Abiotic degradation**

##### **2.2.2.1.1.1 Hydrolysis as a function of pH**

Hydrolysis study following the OECD guideline 111 was performed. Less than 10% hydrolysis was found after 5 days at 50°C for all pHs (4, 7, 9) tested. Consequently, PHMB is considered to be hydrolytically stable.

##### **2.2.2.1.1.2 Photolysis in water**

According to OECD guideline 316, direct photolysis can be an important dissipation pathway for some chemical pollutants that exhibit significant light absorption above the 290 nm cut-off of solar irradiation at the earth's surface. As PHMB absorption spectra maximum was not found in visible wavelength, PHMB could be considered as not photodegradable.

##### **2.2.2.1.1.3 Photolysis in air**

PHMB degrades quickly in the atmosphere by reaction with OH radicals with a highest DT<sub>50</sub> of 1.351 hours (24H day,  $5 \cdot 10^5$  OH/cm<sup>3</sup>). Nonetheless, considering that PHMB is not volatile, potential photodegradation of PHMB is negligible.

Therefore, the abiotic degradation processes will have a minimal influence on the fate and behaviour of PHMB in the environment.

#### **2.2.2.1.2 Biodegradation**

##### **2.2.2.1.2.1 Ready biodegradation**

A ready biodegradation test is performed on the active substance according to OECD guideline 301B. After 99 days, 3.8% of PHMB is mineralized. Thus this substance is considered as non readily biodegradable.

##### **2.2.2.1.2.2 STP compartment**

A simulation test according to OECD 303A guideline is conducted to investigate PHMB degradation in conditions imitating a domestic sewage treatment plant. During the 144 days-period, less than 1% of PHMB is mineralized. 18% of the applied radioactivity is measured in the aqueous effluent, and the residual 82% is sorbed onto the sludge biomass.

PHMB is very slightly mineralized. The water discharge observed is caused only by a modification of PHMB distribution related to its property of adsorption leading to an accumulation of this active substance in activated sludge.

### **2.2.2.1.2.3 Aquatic compartment**

In seawater, a study performed with OECD 306 guideline demonstrated that after 56 days, at concentrations of 1 and 0.1 mg a.s.L<sup>-1</sup>, 2.6% and 10.1% CO<sub>2</sub> mineralisation was observed respectively. For the highest concentration, some evidence of toxicity was noticed and could explain the lower level of mineralization.

### **2.2.2.1.2.4 Water/sediment system**

A simulation test according to OECD 308 guideline was conducted to investigate PHMB degradation in condition imitating aquatic system. The route and rate of [14C]-PHMB biotransformation was investigated under aerobic condition in two flooded sediment systems (loam and loamy sand) over a period of 101 days. PHMB rapidly dissipated from the water phase, partitioning into the sediment phase where it remained tightly bound for the duration of the study. Less than 3% of PHMB was mineralized to CO<sub>2</sub> after a period of 101 days.

Removal from the water phase has a half-life between 1 to 2.3 days. No half-life from the sediment phase and the whole system were available. In both loam and loamy sand sediments, the main amount (from 77% to 97%) of PHMB in the sediment is fixed in the humin fraction (NER).

### **2.2.2.1.2.5 Soil**

Soil biodegradation was investigated in two reliable studies designed to assess the aerobic degradation in soil.

The first of these studies was conducted according to OECD 304A. Less than 5% mineralization of PHMB is observed during the 64 day study and approximately 90% of applied <sup>14</sup>C-PHMB remained bound to soil. No information on primary degradation of the polymers was provided.

The second study assesses the rate and route of degradation in soil according to the OECD guideline 307. Biodegradation of <sup>14</sup>C-PHMB was investigated in four different soils (loamy sand, silty clay loam, clay loam and sandy loam) under aerobic conditions over a period of 123 days. PHMB was highly adsorbed to four different soils, with <5% being mineralized to <sup>14</sup>CO<sub>2</sub>. The amount of PHMB in non extractable residues was >70%. Therefore, it was not possible to identify any breakdown product, nor to calculate degradation kinetics.

As a conclusion, PHMB was found to be non biodegradable and slight rates of mineralization were found in water/sediment system and soil. Moreover, in the aquatic and terrestrial simulation studies, it seems that more than 90% of PHMB is bound with NER while in the sewage treatment plant more than 80% of PHMB is PHMB forms NER. Therefore, PHMB is adsorbed very quickly and very strongly to organic matter, which induces a very limited bioavailability for biodegradation processes.



### **2.2.2.1.3 Distribution**

Several studies on adsorption/desorption properties according to OECD guidelines 121 and 106 show that PHMB adsorbs rapidly and strongly on any kind of sediments, sewage sludge or soils. PHMB remains practically immobile after adsorption. The Koc values are ranged from 151415 to 428713. The arithmetic mean value of Koc of 276670 is used for the risk assessment.

### **2.2.2.1.4 Accumulation**

The low Kow and the high molecular weight indicate the substance is unlikely to bioaccumulate.

## **2.2.2.2 Effects assessment on environmental organisms (active substance)**

### **2.2.2.2.1 Aquatic organisms**

Acute toxicity data are available for fish and algae. An acute key study with *Daphnia magna* (conducted prior to guideline publications but using a test protocol similar to OECD 202) was submitted. eCA considered this study as invalid due to important waiving and because the validity criteria were not fulfilled. This data gap was accepted by eCA since a chronic study was submitted.

Chronic toxicity data are available for the three trophic levels (fish, algae and invertebrates). The most sensitive endpoint is the NOEC/EC10 value of 7.43 µg.L<sup>-1</sup> of a.s. based on growth rate parameter and on measured concentration from growth inhibition test performed on green algae *Selenastrum capricornutum*.



Hence, the  $PNEC_{\text{surface water}}$  is estimated to be  $0.743 \mu\text{g.L}^{-1}$  of a.s. since a safety factor of 10 according to the TGD Part II (2003)<sup>5</sup> should be applied to the lowest endpoint for aquatic environment when acute and chronic data for three trophic levels are available.

#### **2.2.2.2.2 Inhibition of aquatic microbial activity**

The most sensitive NOEC is the one related to the inhibition of nitrification of activated sludge microorganisms, which gives a NOEC of  $12 \text{ mg.L}^{-1}$  of a.s.. By applying an assessment factor of 1 according to the TGD part II, table 17, the  $PNEC_{\text{microorganisms}}$  is estimated to be  $12 \text{ mg.L}^{-1}$  of a.s.

#### **2.2.2.2.3 Sediment dwelling organisms**

No effects were observed at any concentration in a relevant study performed with sediment dwelling organisms. Therefore, the NOEC, based on mean measured concentrations, derived from this 28-day spiked sediment study is equal to  $196 \text{ mg a.s.}\cdot\text{kg}^{-1}$  wwt sediment on *Chironomus riparius*.

With only one long-term test available, an assessment factor of 100 is applied according to the table 19 of the TGD part II to derive the  $PNEC_{\text{sediment}}$ . Therefore, the  $PNEC_{\text{sediment}}$  for a.s. is  $1.96 \text{ mg}\cdot\text{kg}^{-1}$  wwt.

However, it should be noted that during the exposure period, the organisms were fed with a fish food suspension. About feeding of the organism during the test, the standard guideline OECD218 mentioned that [§31, p.7]:

*"When testing strongly adsorbing substances (e.g. with  $\log K_{ow} > 5$ ), or substances covalently binding to sediment, the amount of food necessary to ensure survival and natural growth of the organisms may be added to the formulated sediment before the stabilisation period."* As a consequence the feeding method applied for the test does not follow the standard guideline, considering the high adsorption properties of the PHMB. Therefore, the results from this study should actually be taken with caution.

As a consequence, it was decided at the Working Group (WG)-I-2015 that  $PNEC_{\text{sediment}}$  should also be calculated via EPM with an additional factor of 10 taking the high adsorption properties of PHMB (TGD part II), and the lowest value should be used for the risk assessment.

The  $PNEC_{\text{sediment}}$  was calculated based on equilibrium partitioning by applying the equation 70 of the TGD, part II. Therefore the  $PNEC_{\text{sediment(EPM)}}$  for a.s. is  $446.94 \mu\text{g a.s.}/\text{kg}$  wwt. This value will be used in the risk assessment for sediment compartment.

#### **2.2.2.2.4 Terrestrial compartment**

No adverse effect was observed in the study carried out on microorganisms, plants and earthworms. Therefore, in all studies the relevant endpoint is considered as the highest test concentration. The standardized EC50 derived from the acute toxicity on earthworms gives the lowest value of  $358.2 \text{ mg a.s.}\cdot\text{kg}^{-1}$  wet weight. This value is used to determine the  $PNEC_{\text{soil}}$ .

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<sup>5</sup> Technical Guidance Document (TGD) on Risk Assessment, [Part II](http://echa.europa.eu/documents/10162/16960216/tgdpart2_2ed_en.pdf), Environmental Risk Assessment. 2003. [http://echa.europa.eu/documents/10162/16960216/tgdpart2\\_2ed\\_en.pdf](http://echa.europa.eu/documents/10162/16960216/tgdpart2_2ed_en.pdf).

For the determination of the assessment factor, as no effects were seen in any of the studies, the issue on the most sensitive species as specified in the MOTA v.5 might not be as relevant. Based on the lack of effects in the studies, it was agreed at WG-I-2015 that an AF of 100 should be sufficient to derive the PNEC<sub>soil</sub>. Consequently, the PNEC<sub>soil</sub> for PHMB is 3.58 mg a.s. kg<sup>-1</sup> wet weight.

### 2.2.2.3 Summary of PNEC values

The table below summarises the PNEC value retained for risk assessment:

**Table 2.2-7: PNEC values for the active substance used for the risk assessment part**

PNEC <sub>water</sub>	0.743 µg.L <sup>-1</sup> of a.s.
PNEC <sub>sediment</sub>	446.94 µg.kg <sup>-1</sup> wwt sediment of a.s.
PNEC <sub>soil</sub>	3.58 mg.kg <sup>-1</sup> wwt soil of a.s.
PNEC <sub>microorganisms</sub>	12 mg a.s..L <sup>-1</sup>

### 2.2.2.4 Environmental effect assessment (product)

No additional data on the environment effects of the biocidal products were submitted. The risk assessment is based on the effect of the active substance PHMB.

### 2.2.2.5 PBT, Endocrine Disrupting (ED) and POP assessment

According to the annex XIII of the REACH regulation EC/1907/2006, substances are classified as PBT when they fulfill the criteria for all three inherent properties Persistent (P), Bioaccumulable (B), Toxic (T), and/or vPvB when they fulfill the criteria the two inherent properties very Persistent (vP), very Bioaccumulable (vB).

#### 2.2.2.5.1 Persistence criteria

According to the annex XIII of the REACH regulation, criteria for substance to be persistent (and very persistent) are fulfilled when:

- T<sub>1/2</sub> in marine water > 60 days (60 days for vP criterion) or,
- T<sub>1/2</sub> in fresh or estuarine water > 40 days (60 days for vP criterion) or,
- T<sub>1/2</sub> in marine sediment > 180 days or,
- T<sub>1/2</sub> in freshwater sediment > 120 days (180 days for vP criterion).

According to study results on biodegradability of active substance PHMB in STP, water/sediment, and soil compartment (*c.f.* section 2.2.2.1.2), **PHMB fulfills the P and vP criteria:**

for soil compartment, DT50/DT90 are greater than 1 year, not extractable residues are > 90% in all tested soils, and mineralization is <5% over the 123 days of incubation .

- for surface water, DT50 in whole system is greater than 6 months at 20°C, non-extractable > 90%, and mineralisation is <3% after 101 days.

#### **2.2.2.5.2 Bioaccumulation criteria**

According to the annex XIII of the REACH regulation, criteria for substance to be bioaccumulable are fulfilled when the bioconcentration factor (BCF) exceeds a value of 2000 L/kg. Moreover, a substance is considered to potentially fulfill the B criteria when log  $K_{ow}$  exceeds a value of 4.5.

The applicant has proposed an estimation of the intrinsic potential for bioconcentration using the octanol/water partition coefficient and the models given in the Technical Guidance Document For Risk Assessment Of New And Existing Substances (Chapter 3 p. 126) (TGD part II, 2003). This linear relation is valid only for a  $K_{ow}$  ranging between 2 and 6 or higher than 6 and could not be used for PHMB. Nevertheless, the low  $K_{ow}$ , the high molecular weight (PHMB >700 g/mol) may indicate the substance is unlikely to bioaccumulate. However, PHMB contains also polymers with short chain of carbons which could penetrate into organisms.

Therefore, Applicant reviewed available data and proposed qualitative explanations based on theoretical consideration. Applicant explained that a quantitative prediction of the solubility of low molecular weight oligomers (*i.e.* the dimer) was not considered possible given the available data. However, given the relationship between water solubility and  $K_{ow}$  then a lower solubility would lead to a higher  $K_{ow}$  and thus a higher BCF. Plus, the smallest oligomers, such as dimers, would be expected to have higher water solubility than larger oligomers. It can therefore expect the dimer to have a lower  $K_{ow}$  and thus a lower BCF. Based on this theoretical consideration, there is no concern over the bioaccumulation potential of low MW oligomers. This view is supported by the measured  $K_{ow}$  value ( $K_{ow}$  = 0.005; log  $K_{ow}$  = -2.29) which reflects the value for a mixture of oligomers. This measured  $K_{ow}$  is extremely low and makes it extremely improbable that the  $K_{ow}$  for any low molecular weight oligomers would even approach the generally accepted trigger limit of 4.5.

Based on the  $K_{ow}$ , the BCF for aquatic organism and for terrestrial organisms is estimated to be 0.002 and 0.0013 L/kg, respectively.

Considering the low log $K_{ow}$  (-2.29), the BCF for aquatic organism (0.002) and for terrestrial organisms (0.0013), PHMB is not considered to fulfill the B criterion.

#### **2.2.2.5.3 Toxicity criteria**

According to the annex XIII of the REACH regulation, the toxicity criterion is fulfilled when the chronic NOEC for aquatic organism is less than 0.01 mg/L or when the substance meets the criteria for classification as carcinogenic (1A or 1B), germ cell mutagenic (1A or 1B) or toxic for reproduction (1A, 1B or 2).

Based on ecotoxicity on the most sensitive species *Selenastrum capricornutum* (*i.e.* NOEC/EC10 = 0.00743 mg/L of a.s.), active substance **PHMB is considered to fulfill T criteria.**

Therefore, PHMB is not considered to fulfill the PBT nor vPvB criterion. Anyhow, as PHMB fulfill the criteria of vP and T, **PHMB should be considered as a candidate for substitution, according to the article 10 of the Biocides Regulation EU/528/2012.**

#### **2.2.2.5.4ED properties**

PHMB is not known as an Endocrine Disruptor with regard to the environment. Considering the mode of action of the substance, observed effects on reproduction on fish and daphnia is not expected to be linked to an ED-mode of action.

#### **2.2.2.5.5POP assessment**

According to the screening criteria described in the Annex D of the Stockholm convention, PHMB is not a POP.

#### **2.2.2.6 Environmental exposure assessment**

PHMB HANDWASH is a dummy product presented to illustrate the use of PHMB as the active substance in PT01 disinfectant hand wash products. Such products are used to disinfect the hands and would contain 0.9% w/w of active substance. PHMB HANDWASH would be supplied in a container into which is integrated a small finger operated pump. Operating the pump dispenses 3 mL of product directly onto the hand. The hand washing is followed by a rinsing with water.

PHMB HANDWASH would be used primarily by professional users, but potential use by non-professionals in a domestic setting is also considered.

The Emission Scenario Document (ESD) for products used as human hygiene biocidal products (PT01)<sup>6</sup> recommends scenarios both for private and professional uses, by applying either an annual tonnage approach, or an average consumption approach. For the environmental exposure assessment of PHMB used for hand disinfection, emission calculations based on the recommended scenarios described in the ESD for PT01 were applied. Moreover, the conclusions from the PHMB discussions at WG-I-2015 were also considered for this assessment.

PHMB HANDWASH used for disinfection of hands will ultimately be discharged to drain and will enter a municipal sewage treatment plant (STP). As a result of this, there will be potential for exposure of both the aquatic (surface water and sediment) and the terrestrial (soil and groundwater) compartments, the latter as a result of contaminated sewage sludge spreading on land.

The environmental exposure assessment is performed by considering four scenarios:

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<sup>6</sup> Environmental Emission Scenarios for biocides used as human hygiene biocidal products (PT01). Royal Haskoning (2004). [http://ihcp.jrc.ec.europa.eu/our\\_activities/public-health/risk\\_assessment\\_of\\_Biocides/doc/ESD/ESD\\_PT/PT\\_1\\_Human\\_Hygiene.pdf/view](http://ihcp.jrc.ec.europa.eu/our_activities/public-health/risk_assessment_of_Biocides/doc/ESD/ESD_PT/PT_1_Human_Hygiene.pdf/view)

- Scenario 1-A: private use of PHMB for hand disinfection based on annual tonnage approach.
- Scenario 1-B: private use of PHMB for hand disinfection based on an average consumption approach.
- Scenario 2-A: professional use of PHMB for hand disinfection based on annual tonnage approach (hospital scenario).
- Scenario 2-B: professional use of PHMB for hand disinfection based on an average consumption approach (hospital scenario). The applied scenario takes into account the conclusion of the WG-V-2014 on the Ad hoc Environmental Exposure WG consultation on default value for professional hand disinfection.

### 2.2.2.7 Risk characterisation

To carry out a quantitative risk assessment for the environment when PHMB in PHMB HANDWASH is used as PT01 for hand disinfection, the PEC values were compared to the respective PNEC values for the different compartments, resulting in the following PEC/PNEC ratios summarised in the table below.

As global tonnage of PHMB used for PT01 is far below the calculated tonnages at the break-even point, it was stated that scenarios based on average consumption of PHMB HANDWASH are more appropriate than scenarios based on annual tonnage for assessing the environmental risk of uses of PHMB HANDWASH for hand disinfection.

Therefore the table below summarises the PEC/PNEC ratios for PHMB for private and professional use considering only the average consumption approach.

**Table 2.2-8: PEC/PNEC ratios for PHMB for the private and professional use scenarios of HANDWASH for hand disinfection**

Scenario	1-B: Private use of PHMB for hand disinfection based on an average consumption approach	2-B: Professional use of PHMB for hand disinfection based on an average consumption approach
Freshwater	0.46	<b>1.04</b>
Sediment	<b>4.62</b>	<b>10.4</b>
STP	4.05E-04	9.11E-04
Soil	0.26	0.58
Groundwater	< 0.1*	< 0.1*
Air	Not relevant	

\*According to groundwater concentrations modeled by FOCUS PEARL 4.4.4 and compared to the maximum permissible concentration set for drinking water by the Directive 98/83/EC of 0.1 µg/L.

#### 2.2.2.7.1 Aquatic compartment

The risk assessment estimates that the predicted PHMB emission levels associated with the private and professional uses of PHMB HANDWASH for hand disinfection will give rise to

adverse effects in organisms present in the water column and the sediment for professional uses and in the sediment for private use and is therefore considered **unacceptable** for the aquatic compartment.

#### **2.2.2.7.2 Sewage treatment plant organisms**

The risk assessment estimates that the predicted PHMB emission levels associated with private and professional uses of PHMB HANDWASH for hand disinfection will not give rise to adverse effects in microorganisms present in STP.

As a consequence the risk following private and professional uses of PHMB in PHMB HANDWASH for hand disinfection is considered **acceptable for the STP**.

#### **2.2.2.7.3 Atmosphere**

No risks are expected, considering that the active substance is not volatile.

#### **2.2.2.7.4 Terrestrial compartment**

The risk assessment estimates that the predicted PHMB emission levels associated with private and professional uses of PHMB HANDWASH for hand disinfection will not give rise to adverse effects in organisms present in soil and is therefore considered **acceptable** for this compartment.

#### **2.2.2.7.5 Groundwater**

With regard to predicted PHMB concentrations in groundwater, these do not exceed the 0.1 g/L limit set by the EU Groundwater Directive following private and professional uses of PHMB HANDWASH.

As a consequence the risk following the use of PHMB in PHMB HANDWASH for hand disinfection is considered **acceptable** for groundwater.

#### **2.2.2.7.6 Non compartment specific effects relevant to the food chain (secondary poisoning)**

It is believed that there is no significant potential for secondary poisoning because the low log octanol/water partition coefficient of -2.29, and the high molecular weight of PHMB.

#### **2.2.2.8 Overall conclusion for the environmental risk assessment**

The environmental risk assessment of PHMB used for hand disinfection is summarised in the table below.



**Table 2.2-9: Summary of the environmental risk assessment of PHMB used for hand disinfection**

Scenario	STP	Aquatic compartment	Terrestrial compartment	Groundwater	Air	Secondary poisoning
Hand disinfection: 3 mL of PHMB HANDWASH containing 0.9% a.s. w/w						
Private uses	Acceptable	Unacceptable	Acceptable	Acceptable	Not relevant	
Professional uses						
Conclusions for private and professional uses: the risk for the environment is considered as <b>unacceptable</b> for the aquatic compartment.						

Considering that:

- PHMB HANDWASH is a dummy product presented to illustrate the use of PHMB as the active substance in hand wash products;
- The efficiency of PHMB used in PHMB HANDWASH is demonstrated for hand disinfection at the application rate of 3 mL of PHMB HANDWASH containing 0.9% w/w of active substance for bactericidal activity;
- PHMB HANDWASH would be used primarily by professional users, but potential use by non-professionals in a domestic setting is also considered;
- PHMB HANDWASH used for disinfection of hands will ultimately be discharged to drain and will enter a municipal sewage treatment plant (STP);
- In accordance with the realistic case scenarios applied for the risk assessment:
  - the derived PEC/PNEC ratios for freshwater and sediment for professional uses and for sediment for private uses are above the trigger value of 1;
  - the derived PEC/PNEC ratios for the terrestrial compartment and the STP are below the trigger value of 1 for private and professional uses;
  - the calculated groundwater concentration are below the maximum permissible concentration set for drinking water by the Directive 98/83/EC of 0.1 µg/L for private and professional uses.

In conclusion, the environmental risk assessment for the use of PHMB in product of Product Type 01 as hand disinfection product demonstrates that there are unacceptable risks for the aquatic compartment (including sediment).

### **2.2.3 ASSESSMENT OF ENDOCRINE DISRUPTOR PROPERTIES**

PHMB (1600; 1.8) is not included in the priority list of substances for further evaluation of their role in endocrine disruption established within the Community Strategy for Endocrine Disruptors (COM (1999) 706, COM (2001) 262). Available evidence at this time indicates that PHMB (1600; 1.8) does not have endocrine-disrupting properties (classification criteria specified in Art. 5(3) of Regulation 528/2012 are not met, no effects on endocrine organs and/or reproduction were observed in standard toxicity studies to raise a concern for potential endocrine disruption).

### 2.3 OVERALL CONCLUSION OF THE RISK ASSESSMENT

The outcome of the assessment for PHMB (1600; 1.8) in product-type 1, presented in the Table below, is specified in the BPC opinion following discussions at the 11<sup>th</sup> meeting of the Biocidal Products Committee (BPC). The BPC opinion is available from the ECHA web-site.

#### Substitution/exclusion criteria:

There is no evidence of endocrine effects of PHMB. The substance cannot be considered as carcinogenic, mutagenic and toxic for the reproduction (CMR). PHMB is considered as Toxic for the environment, and very Persistent (vP, T of PBT) and is therefore candidate for substitution.

SCENARIO	Human primary exposure		Human secondary exposure	Environment					
	Professional	Non professional	General public	STP	Aquatic compartment	Terrestrial compartment	Groundwater	Air	Secondary poisoning
<b>Hygienic hand wash</b>									
Bactericide 0.9 % w/w a.s. 30 seconds/3 ml	Acceptable	<b>Not acceptable</b>	Acceptable	Acceptable	<b>Not acceptable</b>	Acceptable	Acceptable	NR	NR

NR: Not relevant.

#### Conditions:

None.



## APPENDICES

### APPENDIX I: LIST OF ENDPOINTS

#### Chapter 1::Identity, Physical and Chemical Properties, Details of Uses, Further Information, and Proposed Classification and Labelling

Active substance (ISO Common Name) PHMB (1600; 1.8) i.e. polyhexamethylene biguanide with a mean number-average molecular weight (Mn) of 1600 and a mean polydispersity (PDI) of 1.8

Function (e.g. fungicide) Bactericide

Rapporteur Member State

France

#### Identity (Annex IIA, point II.)

Chemical name (IUPAC)	CoPoly( bisiminoimidocarbonyl,hexamethylene hydrochloride),(iminoimidocarbonyl, hexame-thylène hydrochloride) or Co poly(5-imino-7-imino-4,6,8-triazaundecamethylene hydrochloride) (5-imino-4,6-diazanonamethylenehydrochloride)
Chemical name (CA)	<ul style="list-style-type: none"> <li>• Guanidine, N,N''-1,6-hexanediylbis[N'-cyano-, polymer with 1,6-hexanediamine, hydrochloride</li> <li>• N,N''-1,6-Hexanediylbis(N'-cyanoguanidine) polymer with 1,6-hexanediamine, hydrochloride</li> <li>• Poly(iminocarbonimidoyliminocarbonimidoylimino-1,6-hexanediyl</li> </ul>
CAS No	27083-27-8 and 32289-58-0
EC No	Not Applicable: the substance is a polymer.
Other substance No.	Not relevant.
Minimum purity of the active substance as manufactured (g/kg or g/l)	The active substance as manufactured (TK) is a 20 % w/w aqueous solution of PHMB plus impurities (total solid) PHMB in dried material $\geq 95.6\%$
Identity of relevant impurities and additives (substances of concern) in the active substance as manufactured (g/kg)	HMD $\leq 4.3$ g/kg

Molecular formula

Terminal function-  $(\text{CH}_2)_6^- [\text{C}_8\text{H}_{18}\text{N}_5\text{Cl}]_n [\text{C}_7\text{H}_{16}\text{N}_3\text{Cl}]_m$  - terminal function

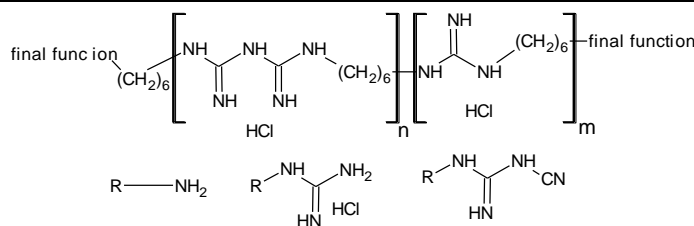
Possible terminal functions:  
 $\text{NH}_2$  (amine)  
 $\text{C}_2\text{H}_3\text{N}_4$  (cyanoguanide)  
 $\text{CH}_5\text{N}_3\text{Cl}$  (guanidine)

		range	average
m+n		2-40	11
n	/(m+n) [biguanide %]	90.8 - 91.9%	91.3 %
m	/(m+n) [guanide %]	8.1 - 9.2 %	8.6 %
Terminal function	amino	35% - 46%	39%
	guanidine	22% - 29%	25%
	cyanoguanide	31 - 39%	35%

Molecular mass

Number average molecular weight (Mn) = 1610  
 Mass average molecular weight (Mw)= 2986.

Structural formula



**Physical and chemical properties** (Annex IIA, point III)

Melting point (state purity)

Glass transition temperature = 90.2-91°C

Boiling point (state purity)

TK : 100.2°C

TC: Decomposition before boiling

Temperature of decomposition

205 to 210°C

Appearance (state purity)

TK : Very pale yellow, Mobile liquid, odourless  
 TC Dusty white solid

Relative density (state purity)

TK : 1.04 at 20°C

TC : 1.20 at 20°C

Surface tension

The active substance is not expected to be surface active based on structural consideration.

Vapour pressure (in Pa, state temperature)

dried PHMB is considered as not volatile

Henry's law constant (Pa m <sup>3</sup> mol <sup>-1</sup> )	Henry's law is not applicable for PHMB. PHMB has only slight possibility to pass from water to air.
Solubility in water (g/l or mg/l, state temperature)	426 g/L at 25°C (41% w/w)
Solubility in organic solvents (in g/l or mg/l, state temperature) (Annex IIIA, point III.1)	Methanol: 41% w/w at 25°C Ethanol: 4.99 g/L (0.5% w/w) Acetone: 2.7 x10 <sup>-3</sup> g/L Dichloromethane: 2.0 x10 <sup>-4</sup> g/L Toluene: 2.0 10 <sup>-4</sup> g/L Ethyl acetate: 1.0 x10 <sup>-4</sup> g/L n-Hexane: 1.0 x10 <sup>-4</sup> g/L Acetonitrile: 8.0 x10 <sup>-4</sup> g/L
Stability in organic solvents used in biocidal products including relevant breakdown products (IIIA, point III.2)	No organic solvent in BP.
Partition coefficient (log P <sub>ow</sub> ) (state temperature)	Log Pow = -2.3 at 25°C ; pH 7.4
Hydrolytic stability (DT <sub>50</sub> ) (state pH and temperature) (point VII.7.6.2.1)	Not calculated: insignificant hydrolysis (<10%) at all pHs after 5 days at 50°C.
Dissociation constant (not stated in Annex IIA or IIIA; additional data requirement from TNsG)	1.2 ± 0.5 x 10 <sup>-1</sup> g equiv/L at 25°C
UV/VIS absorption (max.) (if absorption > 290 nm state ε at wavelength)	Spectrum wavelength maximum: - Distilled water: 236 nm - 0.1M aqueous HCl: 205 nm - 0.1M aqueous NaOH: 234nm
Photostability (DT <sub>50</sub> ) (aqueous, sunlight, state pH) (point VII.7.6.2.2)	Not calculated: Under artificial and natural sunlight, PHMB was not photodegraded in laboratory grade water.
Quantum yield of direct photo-transformation in water at S > 290 nm (point VII.7.6.2.2)	Not relevant. See above.
Flammability	TC: Not Flammable. TC: No ignition below 400°C
Explosive properties	Not Explosive.

**Classification and proposed labelling** (Annex IIA, point IX.)

with regard to physical/chemical data

**Harmonised classification (TC):** None.  
**Proposed classification of PHMB 20 % in water**

with regard to toxicological data

**(TK):** None.  
**Proposed classification of PHMB HANDWASH (0.9% a.s.):** None.

**Harmonised classification (TC):**  
Acute Tox 4; H302: Harmful if swallowed.  
Skin Sens. 1B; H317: May cause an allergic skin reaction.  
Eye Dam. 1; H318: Causes serious eye damage.  
Carc. 2; H351: Suspected of causing cancer.  
STOT RE 1; H372 (respiratory tract) (Inhalation): Causes damage to organs through prolonged or repeated exposure by inhalation.

**Proposed classification of PHMB 20 % in water (TK):**  
Acute Tox 4; H332: Harmful if inhaled.  
Skin Sens. 1B; H317: May cause an allergic skin reaction.  
Carc. 2; H351: Suspected of causing cancer.  
STOT RE 1; H372 (respiratory tract) (Inhalation): Causes damage to organs through prolonged or repeated exposure by inhalation.

**Proposed classification of PHMB HANDWASH (0.9% a.s.):**  
Eye dam.2; H319: Causes serious eye damage.

with regard to fate and behaviour data

**Harmonised classification (TC):** None  
**Proposed classification of PHMB 20 % in water (TK):** None  
**Proposed classification of PHMB HANDWASH (0.9% a.s.) :** None

with regard to ecotoxicological data

**Harmonised classification (TC):**  
Aquatic Acute 1; H400 (M-factor = 10): Very toxic to aquatic life.  
Aquatic Chronic 1; H410 (M-factor = 10): Very toxic to aquatic life with long lasting effects.

**Proposed classification of PHMB 20 % in water (TK):**  
Aquatic Acute 1; H400: Very toxic to aquatic life.  
Aquatic Chronic 1; H410: Very toxic to aquatic life with long lasting effects.

**Proposed classification of PHMB HANDWASH (0.9% a.s.):**  
Aquatic chronic2; H411: Toxic to aquatic life, may

cause long-term adverse effects in the aquatic environment.

**Chapter 2: Methods of Analysis**

**Analytical methods for the active substance**

Technical active substance (principle of method) (Annex IIA, point 4.1)

Gravimetric Analysis: An aliquot of the test substance of known weight is determined gravimetrically after freeze drying until it reaches a constant weight.

Impurities in technical active substance (principle of method) (Annex IIA, point 4.1)

Inorganic salts monitored by determining % w/w sulphated ash.

Residual starting materials monitored by gas chromatography with flame ionisation detection and HPLC with UV detection.

Impurities/related substances, monitored by using size exclusion chromatography (SEC) with UV detection.

Water monitored using Karl Fischer titration.

**Analytical methods for residues**

Soil (principle of method and LOQ) (Annex IIA, point 4.2)

Not technically feasible for an enforcement method

Air (principle of method and LOQ) (Annex IIA, point 4.2)

Occurrence of PHMB in air is not probable.  
No method required

Surface water water (principle of method and LOQ) (Annex IIA, point 4.2)

Not technically feasible for an enforcement method

Drinking water (principle of method and LOQ) (Annex IIA, point 4.2)

Method required

Body fluids and tissues (principle of method and LOQ) (Annex IIA, point 4.2)

Method required

Food/feed of plant origin (principle of method and LOQ for methods for monitoring purposes) (Annex IIIA, point IV.1)

Method required.

Food/feed of animal origin (principle of method and LOQ for methods for monitoring purposes) (Annex IIIA, point IV.1)

**Chapter 3: Impact on Human Health**

**Absorption, distribution, metabolism and excretion in mammals** (Annex IIA, VI.6.2)

Rate and extent of oral absorption:	4% = closest estimate (oral absorption of PHMB ranges approximately from 0.3 to 8%).
Rate and extent of dermal absorption:	4% corresponding to oral absorption, based on default value proposed in the EFSA guidance on dermal absorption.
Distribution:	Uniformly distributed. Target organs: liver and kidneys
Potential for accumulation:	No evidence for bioaccumulation.
Rate and extent of excretion:	Most excreted (>90%) in the faeces.
Toxicologically significant metabolite	-

**Acute toxicity** (Annex IIA, VI.6.1)

Rat LD <sub>50</sub> oral	The oral LD <sub>50</sub> of the 20 % aqueous solution is from 2.5 g (Vantocil P)/kg to > 5g /kg of PHMB 20 % w/w in rat
Rat LD <sub>50</sub> dermal	The dermal LD <sub>50</sub> of the 20 % aqueous solution is > 2000 mg/kg of PHMB 20 % w/w in rabbit.
Rat LC <sub>50</sub> inhalation	No available acute data. Based on RAC opinion: Xn; R20 is warranted.
Skin irritation	Slight to moderate irritant on rabbit. Slight irritant to human skin. But does not meet the criteria for classification.
Eye irritation	20% PHMB in aqueous solution is a moderate irritant but does not meet the criteria for classification.
Skin sensitization (test method used and result)	Moderate to strong potency sensitizer based on animal data. Human studies indicate that PHMB is a skin sensitizer in humans, although with a rare frequency of sensitisation in the current conditions of consumer uses. It meets the classification criteria for an R43, may cause sensitisation by skin contact or Skin Sens. 1B H317 because of low incidences from human data.

**Repeated dose toxicity** (Annex IIA, VI. 6.3, 6.4, and 6.5)

Species/ target / critical effect	Rat/liver and kidney/slight effects to parameters of clinical chemistry, decrease in weight gain, minor histopathological change to the liver and kidneys.  Acute, mid and long-term exposure:  NOAEL = 13 mg/kg/d (Rat - developmental study)
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Lowest relevant inhalation NOAEC

Acute, mid and long-term exposure: Rat – 28 day exposure – 0.024 mg/m<sup>3</sup>

**Genotoxicity** (Annex IIA, VI.6.6)

Not genotoxic *in vitro* or *in vivo*.

**Carcinogenicity** (Annex IIA, VI.6.7)

Species/type of tumour

PHMB increases the incidence of benign and malign vascular tumours in female rats by oral route and in male and female mice by oral and dermal route. The tumours are induced mainly in the liver, which is one of the target organ of PHMB and the increase is clearly seen at doses above the MTD. However, it is also observed more equivocally at doses below MTD (mouse oral study at mid-dose and rat oral study at high dose). These increases are not considered incidental when considering the clear induction of vascular tumours at higher doses and they are considered biologically significant and attributed to treatment.

A classification as carcinogenic category 3; R40 is warranted.

lowest dose with tumours

Rat – via diet - NOAEL for carcinogenicity can be established at 36 mg/kg bw/d in males and 45 mg/kg bw/d in females.

**Reproductive toxicity** (Annex IIA, VI.6.8)

Species/ Reproduction target / critical effect

Rat – lower bodyweights in F0 and F1 animals during the pre-mating period.

Lowest relevant reproductive NOAEL

F0 – 600 ppm (70 – 77 mg/kg bw/d)  
F1 – 600 ppm (70 – 77 mg/kg bw/d)  
F2 – 2000 ppm (239 - 258 mg/kg bw/d)

Species/Developmental target / critical effect

Rabbit – no developmental effects related to treatment.

Rat – increase in extra ribs at maternal toxic doses.

Lowest relevant developmental NOAEL

Rabbit:  
Parental: 20 mg/kg/d  
Developmental: 20 mg/kg/d  
Rat:  
Parental: 13 mg/kg/d  
Developmental: 54 mg/kg/d

**Neurotoxicity** (Annex IIIA, VI.1)

Species/ target/critical effect

Not applicable since no specific studies have been conducted for this endpoint.

Lowest relevant neurotoxicity NOAEL

N/A

**Other toxicological studies** (Annex IIIA, VI/XI)

Neurotoxicity

See section on neurotoxicity.

Toxic effects on livestock and pets

Not relevant, low exposure.

Studies related to the exposure of the a.s. to humans

Studies related to human exposure of the a. s. are not required on the basis of the results of the human health exposure and risk assessments.

Food and feeding stuffs

Transfer of the active substance to food cannot be excluded. However, since the risk is unacceptable for secondary exposure (hand to mouth transfer), the risk related to transfer of the active substance to food was not investigated. .

Other tests related to exposure of the a.s. to human considered to be necessary

Further studies are not necessary for the purpose of a comprehensive evaluation of the a. s.

Tests to assess toxic effects from metabolites of treated plants

Not relevant because PHMB-based products are not used on plants.

Mechanistic studies

No studies are available with data to define the mechanism of action for the toxicity.

Further human health related studies

Not required.

**Medical data** (Annex IIA, VI.6.9)

Medical surveillance data on manufacturing plant personnel

No evidence of adverse effects on workers of manufacturing plants.

Direct observations, e.g. clinical cases, poisoning incidents

No data available.

Health records, both from industry and any other sources

From the data available, no evidence of adverse health effects of PHMB.

Epidemiological studies on the general population

No data available.

Diagnosis of poisoning including specific signs of poisoning and clinical tests

**Skin:** Exposure may cause redness and swelling.  
**Eye:**  
**20% PHMB in aqueous solution:** Exposure may cause eye irritation –redness and swelling.  
**Inhalation:** irritation of the respiratory tract may occur. Exposure may cause coughing.  
**Ingestion:** may cause irritation of the gastrointestinal tract with nausea vomiting or diarrhoea.

Sensitization/allergenicity observations

PHMB is a skin sensitizer in humans, although with a rare frequency of sensitisation in the current conditions of consumer uses.



Specific treatment in case of an accident or poisoning: first aid measures and medical treatment

**Skin:** Remove contaminated clothing. Wash immediately with water followed by soap and water. Obtain medical attention.

Patient may experience an eczematous rash to compound should they have been sensitized by prior exposure. This rash would be expected to respond to removal from exposure and treatment with cortico-steroids.

Contaminated clothing should be laundered before re-issue.

**Eye:**

**20% PHMB in aqueous solution:** Irrigate with eyewash solution or clean water, holding the eyelids apart, for at least 15 minutes. Obtain medical attention as a precaution.

**Inhalation:** Remove patient from exposure. Obtain medical attention if ill effects occur.

**Ingestion:** Provided the patient is conscious, wash out mouth with water and give 200-300 ml (half a pint) of water to drink.

Do not induce vomiting. Obtain medical attention.

Prognosis following poisoning

The prognosis is excellent if First Aid is administered promptly.

**Skin:** Prompt cleansing should minimize irritation to the skin. Patient may be experience sensitization to compound should future exposure occur.

**Eye:** Prompt irrigation should minimize irritation of the eye.

**Inhalation:** Prompt removal from exposure should minimize irritation to the respiratory tract.

**Ingestion:** Prompt treatment should minimize irritation of the gastrointestinal tract.

**Summary** (Annex IIA, VI.6.10)

**Systemic effects**

	<b>AEL</b>	<b>MOE<sub>ref</sub></b>
acute, medium and long-term	5.2 µg a.s./kg bw/d	100
	<b>ADI - ARfD</b>	<b>MOE<sub>ref</sub></b>
Chronic and acute	0.13 mg a.s./kg bw/d	100
<b>Local effects by inhalation</b>		

**Summary** (Annex IIA, VI.6.10)

	<b>AEC</b>	<b>MOE<sub>ref</sub></b>
acute	0.96 µg/m <sup>3</sup>	25
medium-term	0.32 µg/m <sup>3</sup>	75
long-term	0.16 µg/m <sup>3</sup>	150

**Acceptable exposure scenarios** (including method of calculation)

Professional users

The primary exposure of professionals to PHMB HANDWASH is considered to be acceptable.

Non-professional users

The primary exposure of non-professionals to PHMB HANDWASH is considered to be unacceptable.

Indirect exposure as a result of use

The possible secondary exposure to PHMB HANDWASH is considered to be unacceptable.  
The potential combined exposures to PHMB HANDWASH are considered to be unacceptable.

<sup>1)</sup> Technical Notes for Guidance – Human Exposure to Biocidal Products – Guidance on Exposure Estimation (June 2002)

**Chapter 4: Fate and Behaviour in the Environment**

**Route and rate of degradation in water** (Annex point IIA, VII.7.6; Annex point IIIA, XII.2.1, 2.2)

Hydrolysis of active substance and relevant metabolites (DT<sub>50</sub>) (state pH and temperature)

50°C, pH 4, 7 and 9: hydrolytically stable (<10% hydrolysis seen after 5 days).

No metabolites identified.

Photolytic / photo-oxidative degradation of active substance and resulting relevant metabolites

PHMB absorption spectra maximum was not found in visible wavelength. PHMB is considered as not photodegradable

Readily biodegradable (yes/no)

No.

Inherent biodegradability

No.

Biodegradation in seawater

Up to 10.1% mineralisation after 56 days.

Anaerobic water/sediment study:  
(nonsterile) DT<sub>50</sub> total systems  
(nonsterile) DT<sub>90</sub> total systems

No DT<sub>50 total system</sub> determined

Non-extractable residues

According to a water/sediment degradation study on

Distribution in water / sediment systems (active substance)	PHMB, > 90% of non-extractable residues in sediment after 101 days. According to a water/sediment degradation study on PHMB: - Water = 0.3% after 101 days (DT <sub>50</sub> for removal from the water phase are 1 to 2.3 days); - Sediment > 90% after 101 days; - Mineralisation <3% after 101 days.
Distribution in water / sediment systems (metabolites)	It was not possible to investigate the identity of degradation products due to the sorptive nature of PHMB.

**Route and rate of degradation in soil** (Annex point IIIA, VII.4, XII.1.1, XII.1.4; Annex VI, para. 85)

Mineralisation (aerobic)	Less than 5% mineralisation after 123 days.
Laboratory studies (range or median, with number of measurements, with regression coefficient)	DT <sub>50</sub> lab (25°C, aerobic)- not calculated as <5% mineralisation observed.
Field studies (state location, range or median with number of measurements)	No direct soil exposure expected. Therefore, there is no requirement for terrestrial testing and submission of a field soil dissipation and accumulation study is not required.
Anaerobic degradation	Further studies not required as exposure to anaerobic conditions is not likely where the active substance is to be used.
Soil photolysis	Not required because the degradation of PHMB in soil is primarily microbially mediated.
Non-extractable residues	According to a soil degradation study on PHMB, > 90% of non-extractable residues in soil after 123 days.
Relevant metabolites - name and/or code, % of applied a.s. (range and maximum)	It was not possible to investigate the identity of degradation products due to the sorptive nature of PHMB.
Soil accumulation and plateau concentration	Not required. According to the TNSG this study is required only where the biocide is directly applied or emitted to soil. From the Risk assessment at Doc IIB Chapter 3 and IIC chapter 2, there is no direct soil exposure.

**Adsorption/desorption**

Ka , Kd	Kd (adsorption distribution coefficient): 3172-7614 L/kg (arithmetic mean value of 6177 L/kg)
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Ka <sub>oc</sub> , Kd <sub>oc</sub>	Kom: 88032-244036 L/kg (arithmetic mean value of 160344 L/kg)
pH dependence (yes / no) (if yes type of dependence)	Koc: 151415-428713 L/kg (arithmetic mean value of 276670 L/kg) Adsorption is independent of pH.
K <sub>oc</sub>	276670 L/kg (log K <sub>oc</sub> = 5.44)
Leaching of PHMB	No Leaching studies conducted.

**Fate and behaviour in air** (Annex point IIIA, VII.3, VII.5)

Direct photolysis in air	Not required.
Quantum yield of direct photolysis	Not determined.
Photo-oxidative degradation in air	DT <sub>50</sub> 1.351 – 6.37 hours (24H day, 5 x 10 <sup>5</sup> OH/cm <sup>3</sup> ) derived by the Atkinson method of calculation.
Volatilisation	PHMB is not volatile.

**Monitoring data, if available** (Annex VI, para. 44)

Soil (indicate location and type of study)	No monitoring data has been reported.
Surface water (indicate location and type of study)	No monitoring data has been reported.
Ground water (indicate location and type of study)	No monitoring data has been reported.
Air (indicate location and type of study)	No monitoring data has been reported.

**Chapter 5: Effects on Non-target Species**

**Toxicity data for aquatic species (most sensitive species of each group) for PHMB**

(Annex IIA, VII. 7.1 - 7.4, Annex IIIA, XII. 2.2 and XII 2.4)

Species	Time-scale	Endpoint	Toxicity
<b>Fish</b>			
<i>Oncorhynchus mykiss</i>	96 h (flow through system)	Mortality	LC <sub>50</sub> : 26 µg PHMB.l <sup>-1</sup> (mc) NOEC: 9.8 µg PHMB.l <sup>-1</sup> (mc)
<i>Oncorhynchus mykiss</i>	28 days (flow through system)	Growth	NOEC = 10 µg PHMB.l <sup>-1</sup> (mc)
<b>Invertebrates</b>			

<i>Daphnia magna</i>	21 days (semi static system)	Growth and reproduction	NOEC: 8.4 µg PHMB.l <sup>-1</sup> (mc)
<b>Algae</b>			
<i>Selenastrum capricornutum</i>	72 h (static system)	Rate	ErC <sub>50</sub> = 15 µg.l <sup>-1</sup> (mc) NOEC = 7.43 µg.l <sup>-1</sup> (mc)
<b>Microorganisms</b>			
Activated sludge	4 h	Nitrification inhibition	NOEC: 12 mg PHMB.l <sup>-1</sup> (mc)
Active anaerobic sludge	48 h	Inhibition of CO <sub>2</sub> and CH <sub>4</sub> production	NOEC: 20 mg PHMB.g <sup>-1</sup> MLTS (mc)

(mc: measured concentration)

**Effects on earthworms or other soil non-target organisms**

(Annex IIIA, XIII.3.2)

Acute toxicity to earthworm  
(Annex IIIA, point XIII.3.2)

Mortality after a 14-days exposure:  
LC<sub>50</sub>: > 882 mg PHMB.kg<sup>-1</sup> wet weight soil  
NOEC = 882 mg PHMB.kg<sup>-1</sup> wet weight soil

After normalization at 3.4% of organic matter:  
LC<sub>50\_std</sub>: > 358.2 mg PHMB.kg<sup>-1</sup> wet weight soil  
NOEC<sub>std</sub> = 358.2 mg PHMB.kg<sup>-1</sup> wet weight soil

Reproductive toxicity to other soil non-target macro-organisms, long-term test with terrestrial plants  
(Annex IIIA, point XIII.3.2)

Not required.

**Effects on soil micro-organisms**

(Annex IIA, VII.7.4)

Nitrogen transformation

Inhibition after a 14-days exposure:  
LC<sub>50</sub>: > 882 mg PHMB.kg<sup>-1</sup> wet weight soil  
NOEC = 882 mg PHMB.kg<sup>-1</sup> wet weight soil

After normalization at 3.4% of organic matter:  
LC<sub>50\_std</sub>: > 1609.01 mg PHMB.kg<sup>-1</sup> wet weight soil  
NOEC<sub>std</sub> = 1609.01 mg PHMB.kg<sup>-1</sup> wet weight soil

Carbon mineralisation

Not required

**Effects on sediment dwelling organisms**

(Annex IIIA, XIII.3.4)

Toxicity to *Chironomus riparius*

Emergence of adult midges over to a 28-day period in spiked sediment:

EC<sub>50</sub> > 196 mg PHMB.kg<sup>-1</sup> wet weight sediment (measured concentration)

NOEC = 196 mg PHMB. kg<sup>-1</sup> wet weight sediment (measured concentration)

**Effects on plants**

(Annex IIIA, XIII.3.4)

Toxicity to plants (*Avena sativa*, *Brassica oleracea*, *Phaseolus aureus*)

Seedling emergence after a 28-days exposure:

EC<sub>50</sub>: >1000 mg PHMB.kg<sup>-1</sup> wet weight soil

NOEC: 1000 mg PHMB.kg<sup>-1</sup> wet weight soil

After normalization at 3.4% of organic matter:

LC<sub>50\_std</sub>: > 772.73 mg PHMB.kg<sup>-1</sup> wet weight soil

NOEC<sub>std</sub> = 772.73 mg PHMB.kg<sup>-1</sup> wet weight soil

**Effects on terrestrial vertebrates**

Acute toxicity to mammals (Annex IIIA, point XIII.3.3)

Data submitted in Doc IIIA, Section 6 (Mammalian Toxicity) adequately describes the toxicity to mammals. Additional data/testing on mammals is not appropriate and would be against the spirit of EU legislation on minimising animal testing.

Acute toxicity to birds (Annex IIIA, point XIII.1.1)

Not required

Dietary toxicity to birds (Annex IIIA, point XIII.1.2)

Not required

Reproductive toxicity to birds (Annex IIIA, point XIII.1.3)

Not required

**Effects on honeybees** (Annex IIIA, point XIII.3.1)

Acute oral toxicity

Not required

Acute contact toxicity

Not required

**Effects on other beneficial arthropods** (Annex IIIA, point XIII.3.1)

Acute oral toxicity

Not required

Acute contact toxicity

Not required

Acute toxicity to other beneficial arthropods

Not required

**Bio-concentration** (Annex IIA, point 7.5)

Bio-concentration factor (BCF)

BCF<sub>aquatic organism</sub> calculated from log Kow = 0.002;  
BCF<sub>terrestrial organism</sub> calculated from log Kow = 0.0013;  
therefore no bioaccumulation expected.

Depuration time (DT<sub>50</sub>) / (DT<sub>90</sub>)

Not applicable as no bioaccumulation expected.

Level of metabolites (%) in organisms accounting for > 10 % of residues

Not applicable as no bioaccumulation expected.

**Chapter 6: Other End Points**

Not applicable, no other end points

**APPENDIX II: LIST OF INTENDED USES**

Object and/or situation	Product name	Organisms controlled	Formulation		Application			Applied amount per treatment	Remarks
			Type	Conc [% a.s.]	Method	Number	Interval		
Example 1: Handwash	PHMB HANDWASH	Bacteria	AL*	0.9 % w/w	Finger pump	Use as required**	Use as required**	3 ml Contact time : 30 seconds	Professional and non professional

Note \*: AL: Other liquids to be applied undiluted

Note \*\*: e.g. food industry environment: up to eight applications per day (e.g. before work, after each break, after restroom visits). In a hospital/ medical environment, this may be up to thirty times daily, e.g. between patients. In a domestic setting the number of applications is likely to be much less.



## APPENDIX III: LIST OF STANDARD ABBREVIATIONS

**List of standard terms and abbreviations** (adapted from: (i) Guidelines and criteria for the preparation of PPP dossiers<sup>7</sup>; (ii) TNsG on Data Requirements<sup>8</sup>).

Stand. term/Abbreviation	Explanation
A	ampere
ACh	acetylcholine
AChE	acetylcholinesterase
ADI	acceptable daily intake
ADME	administration distribution metabolism and excretion
ADP	adenosine diphosphate
AE	acid equivalent
AF	assessment factor
AFID	alkali flame-ionisation detector or detection
A/G	albumin/globulin ratio
a.i.	active ingredient
ALT	alanine aminotransferase (SGPT)
<i>Ann.</i>	Annex
AEC	acceptable concentration level
AEL	acceptable exposure level
AMD	automatic multiple development
ANOVA	analysis of variance
AP	alkaline phosphatase
approx	approximate

<sup>7</sup> EU (1998a): European Commission: Guidelines and criteria for the preparation of complete dossiers and of summary dossiers for the inclusion of active substances in Annex I of Directive 91/414/EC (Article 5.3 and 8.2). Document 1663/VI/94 Rev 8, 22 April 1998

<sup>8</sup> European Chemicals Bureau, ECB (1996) Technical Guidance Documents in support of the Commission Directive 93/67/EEC on risk assessment for new notified substances and the Commission Regulation (EC) 1488/94 for existing substances

Stand. term/Abbreviation	Explanation
ARfD	acute reference dose
a.s.	active substance (TC)
AST	aspartate aminotransferase (SGOT)
ASV	air saturation value
ATP	adenosine triphosphate
BAF	bioaccumulation factor
BCF	bioconcentration factor
bfa	body fluid assay
BOD	biological oxygen demand
bp	boiling point
BPD	Biocidal Products Directive
BSAF	biota-sediment accumulation factor
BSP	bromosulphophthalein
Bt	<i>Bacillus thuringiensis</i>
Bti	<i>Bacillus thuringiensis israelensis</i>
Btk	<i>Bacillus thuringiensis kurstaki</i>
Btt	<i>Bacillus thuringiensis tenebrionis</i>
BUN	blood urea nitrogen
bw	body weight
c	centi- ( $\times 10^{-2}$ )
°C	degrees Celsius (centigrade)
CA	controlled atmosphere
CAD	computer aided design
CADDY	computer aided dossier and data supply (an electronic dossier interchange and archiving format)
cd	candela
CDA	controlled drop(let) application
cDNA	complementary DANN
CEC	cation exchange capacity
<i>cf</i>	confer, compare to
CFU	colony forming units
ChE	cholinesterase
CI	confidence interval
CL	confidence limits

<b>Stand. term/Abbr eviation</b>	<b>Explanation</b>
cm	centimetre
CNS	central nervous system
COD	chemical oxygen demand
CPK	creatinine phosphatase
cv	coefficient of variation
Cv	ceiling value
d	day(s)
DCA	Dichloroacetaldehyde
DDVP	Dimethyl Dichloro Vinyl Phosphate
DIS	draft international standard (ISO)
DMSO	dimethylsulfoxide
DNA	deoxyribonucleic acid
dna	designated national authority
DO	dissolved oxygen
DOC	dissolved organic carbon
dpi	days post inoculation
DRP	detailed review paper (OECD)
DT <sub>50(lab)</sub>	period required for 50 percent dissipation (under laboratory conditions) (define method of estimation)
DT <sub>90(field)</sub>	period required for 90 percent dissipation (under field conditions) (define method of estimation)
dw	dry weight
ε	decadic molar extinction coefficient
EC <sub>50</sub>	median effective concentration
ECD	electron capture detector
ED <sub>50</sub>	median effective dose
EINECS	European inventory of existing commercial substances
ELINCS	European list of notified chemical substances
ELISA	enzyme linked immunosorbent assay
e-mail	electronic mail
EMDI	estimated maximum daily intake

<b>Stand. term/Abbr eviation</b>	<b>Explanation</b>
EN	European norm
EPMA	electron probe micro-analysis
ERL	extraneous residue limit
ESPE46/51	evaluation system for pesticides
EUSES	European Union system for the evaluation of substances
F	field
F <sub>0</sub>	parental generation
F <sub>1</sub>	filial generation, first
F <sub>2</sub>	filial generation, second
FBS	full base set
FELS	fish early-life stage
FIA	fluorescence immuno-assay
FID	flame ionisation detector
F <sub>mol</sub>	fractional equivalent of the metabolite 's molecular weight compared to the active substance
FOB	functional observation battery
f <sub>oc</sub>	organic carbon factor (compartment dependent)
fp	freezing point
FPD	flame photometric detector
FPLC	fast protein liquid chromatography
g	gram(s)
GC	gas chromatography
GC-EC	gas chromatography with electron capture detector
GC-FID	gas chromatography with flame ionisation detector
GC-MS	gas chromatography-mass spectrometry
GC-MSD	gas chromatography with mass-selective detection
GEP	good experimental practice
GFP	good field practice
GGT	gamma glutamyl transferase
GI	gastro-intestinal
GIT	gastro-intestinal tract
GL	guideline level

<b>Stand. term/Abbr eviation</b>	<b>Explanation</b>
GLC	gas liquid chromatography
GLP	good laboratory practice
GM	geometric mean
GMO	genetically modified organism
GMM	genetically modified micro-organism
GPC	gel-permeation chromatography
GPMT	guinea pig maximisation test
GPS	global positioning system
GSH	glutathione
GV	granulosevirus
h	hour(s)
H	Henry's Law constant (calculated as a unitless value)
ha	hectare(s)
Hb	haemoglobin
HC5	concentration which will be harmless to at least 95 % of the species present with a given level of confidence (usually 95 %)
HCG	human chorionic gonadotropin
Hct	haematocrit
HDT	highest dose tested
hL	hectolitre
HEED	high energy electron diffraction
HID	helium ionisation detector
HPAEC	high performance anion exchange chromatography
HPLC	high pressure liquid chromatography or high performance liquid chromatography
HPLC-MS	high pressure liquid chromatography - mass spectrometry
HPPLC	high pressure planar liquid chromatography
HPTLC	high performance thin layer chromatography
HRGC	high resolution gas chromatography

<b>Stand. term/Abbr eviation</b>	<b>Explanation</b>
H <sub>s</sub>	Shannon-Weaver index
Ht	haematocrit
HUSS	human and use safety standard
I	indoor
I <sub>50</sub>	inhibitory dose, 50%
IC <sub>50</sub>	median immobilisation concentration or median inhibitory concentration 1
ICM	integrated crop management
ID	ionisation detector
IEDI	international estimated daily intake
IGR	insect growth regulator
im	intramuscular
inh	inhalation
INT	2-p-iodophenyl-3-p-nitrophenyl-5-phenyltetrazoliumchloride testing method
ip	intraperitoneal
IPM	integrated pest management
IR	infrared
IRAC	Insecticide resistance action committee
ISBN	international standard book number
ISSN	international standard serial number
IUCLID	International Uniform Chemical Information Database
iv	intravenous
IVF	<i>in vitro</i> fertilisation
k ( <i>in combination</i> )	kilo
k	rate constant for biodegradation
K	Kelvin
K <sub>a</sub>	acid dissociation constant
K <sub>b</sub>	base dissociation constant
K <sub>ads</sub>	adsorption constant
K <sub>des</sub>	apparent desorption coefficient

<b>Stand. term/Abbr eviation</b>	<b>Explanation</b>
kg	kilogram
K <sub>H</sub>	Henry 's Law constant (in atmosphere per cubic metre per mole)
K <sub>oc</sub>	organic carbon adsorption coefficient
K <sub>om</sub>	organic matter adsorption coefficient
K <sub>ow</sub>	octanol-water partition coefficient
K <sub>p</sub>	solid-water partition coefficient
kPa	kilopascal(s)
l, L	litre
LAN	local area network
LASER	light amplification by stimulated emission of radiation
LBC	loosely bound capacity
LC	liquid chromatography
LC-MS	liquid chromatography- mass spectrometry
LC <sub>50</sub>	lethal concentration, median
LCA	life cycle analysis
LC-MS-MS	liquid chromatography with tandem mass spectrometry
LD <sub>50</sub>	lethal dose, median; dosis letalis media
LDH	lactate dehydrogenase
ln	natural logarithm
LOAEC	lowest observable adverse effect concentration
LOAEL	lowest observable adverse effect level
LOD	limit of detection
LOEC	lowest observable effect concentration
LOEL	lowest observable effect level
log	logarithm to the base 10
LOQ	limit of quantification (determination)
LPLC	low pressure liquid chromatography
LSC	liquid scintillation counting or

<b>Stand. term/Abbr eviation</b>	<b>Explanation</b>
	counter
LSD	least squared denominator multiple range test
LSS	liquid scintillation spectrometry
LT	lethal threshold
m	metre
M	molar
µm	micrometre (micron)
MAC	maximum allowable concentration
MAK	maximum allowable concentration
MC	moisture content
MCH	mean corpuscular haemoglobin
MCHC	mean corpuscular haemoglobin concentration
MCV	mean corpuscular volume
MDL	method detection limit
MFO	mixed function oxidase
µg	microgram
mg	milligram
MHC	moisture holding capacity
MIC	minimum inhibitory concentration
min	minute(s)
MKC	minimum killing concentration
mL	millilitre
MLT	median lethal time
MLD	minimum lethal dose
mm	millimetre
MMAD	mass median aerodynamic diameter
mo	month(s)
MOE	margin of exposure
mol	mole(s)
mp	melting point
MRE	maximum residue expected
MRL	maximum residue level or limit
mRNA	messenger ribonucleic acid

<b>Stand. term/Abbreviation</b>	<b>Explanation</b>
MS	mass spectrometry
MSDS	material safety data sheet
MTD	maximum tolerated dose
MT	material test
MW	molecular weight
n.a.	not applicable
n-	normal (defining isomeric configuration)
n	number of observations
NAEL	no adverse effect level
nd	not detected
NEDI	national estimated daily intake
NEL	no effect level
NERL	no effect residue level
ng	nanogram
nm	nanometre
NMR	nuclear magnetic resonance
no, n°	number
NOAEC	no observed adverse effect concentration
NOAEL	no observed adverse effect level
NOEC	no observed effect concentration
NOED	no observed effect dose
NOEL	no observed effect level
NOIS	notice of intent to suspend
NPD	nitrogen-phosphorus detector or detection
NPV	nuclear polyhedrosis virus
NR	not reported
NTE	neurotoxic target esterase
OC	organic carbon content
OCR	optical character recognition
ODP	ozone-depleting potential
ODS	ozone-depleting substances
OH	hydroxide
OJ	Official Journal
OM	organic matter content

<b>Stand. term/Abbreviation</b>	<b>Explanation</b>
OP	Organophosphate
Pa	pascal
PAD	pulsed amperometric detection
2-PAM	2-pralidoxime
pc	paper chromatography
PC	personal computer
PCV	haematocrit (packed corpuscular volume)
PDI	polydispersity
PEC	predicted environmental concentration
PEC <sub>A</sub>	predicted environmental concentration in air
PEC <sub>S</sub>	predicted environmental concentration in soil
PEC <sub>SW</sub>	predicted environmental concentration in surface water
PEC <sub>GW</sub>	predicted environmental concentration in ground water
PED	plasma-emissions-detector
pH	pH-value
PHED	pesticide handler's exposure data
PIC	prior informed consent
pic	phage inhibitory capacity
PIXE	proton induced X-ray emission
pKa	negative logarithm (to the base 10) of the acid dissociation constant
pKb	negative logarithm (to the base 10) of the base dissociation constant
PND	post natal day
PNEC	predicted no effect concentration (compartment to be added as subscript)
po	by mouth
POP	persistent organic pollutants
ppb	parts per billion (10 <sup>-9</sup> )
PPE	personal protective equipment
ppm	parts per million (10 <sup>-6</sup> )
PPP	plant protection product

<b>Stand. term/Abbr eviation</b>	<b>Explanation</b>
ppq	parts per quadrillion ( $10^{-24}$ )
ppt	parts per trillion ( $10^{-12}$ )
PSP	phenolsulfophthalein
PrT	prothrombin time
PRL	practical residue limit
PT	product type
PT(CEN)	project team CEN
PTT	partial thromboplastin time
QA	quality assurance
QAU	quality assurance unit
(Q)SAR	quantitative structure-activity relationship
r	correlation coefficient
$r^2$	coefficient of determination
RA	risk assessment
RBC	red blood cell
REI	restricted entry interval
RENI	Registry Nomenclature Information System
Rf	retardation factor
RfD	reference dose
RH	relative humidity
RL <sub>50</sub>	median residual lifetime
RNA	ribonucleic acid
RP	reversed phase
rpm	revolutions per minute
rRNA	ribosomal ribonucleic acid
RRT	relative retention time
RSD	relative standard deviation
RTU	ready-to-use
s	second
S	solubility
SAC	strong adsorption capacity
SAP	serum alkaline phosphatase
SAR	structure/activity relationship
SBLC	shallow bed liquid chromatography
sc	subcutaneous

<b>Stand. term/Abbr eviation</b>	<b>Explanation</b>
sce	sister chromatid exchange
SCAS	semi-continous activated sludge
SCTER	smallest chronic toxicity exposure ratio (TER)
SD	standard deviation
se	standard error
SEM	standard error of the mean
SEP	standard evaluation procedure
SF	safety factor
SFC	supercritical fluid chromatography
SFE	supercritical fluid extraction
SIMS	secondary ion mass spectroscopy
S/L	short term to long term ratio
SMEs	small and medium sized enterprises
SOP	standard operating procedures
sp	species (only after a generic name)
SPE	solid phase extraction
SPF	specific pathogen free
spp	subspecies
SSD	sulphur specific detector
SSMS	spark source mass spectrometry
STEL	short term exposure limit
STER	smallest toxicity exposure ratio (TER)
STMR	supervised trials median residue
STP	sewage treatment plant
t	tonne(s) (metric ton)
$t_{1/2}$	half-life (define method of estimation)
T <sub>3</sub>	tri-iodothyroxine
T <sub>4</sub>	thyroxine
T <sub>25</sub>	tumorigenic dose that causes tumours in 25 % of the test animals
TADI	temporary acceptable daily

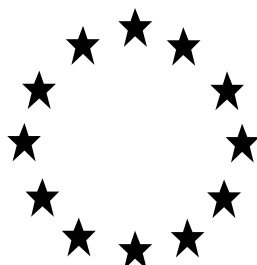
<b>Stand. term/Abbreviation</b>	<b>Explanation</b>
	intake
TBC	tightly bound capacity
TC	technical material according to GIFAP monograph n°2 nomenclature
TCD	thermal conductivity detector
TG	technical guideline, technical group
TGD	Technical guidance document
TID	thermionic detector, alkali flame detector
TDR	time domain reflectometry
TER	toxicity exposure ratio
TER <sub>I</sub>	toxicity exposure ratio for initial exposure
TER <sub>ST</sub>	toxicity exposure ratio following repeated exposure
TER <sub>LT</sub>	toxicity exposure ratio following chronic exposure
tert	tertiary (in a chemical name)
TEP	typical end-use product
TGGE	temperature gradient gel electrophoresis
TIFF	tag image file format
TK	TK: technical concentrate according to GIFAP monograph n°2 nomenclature
TLC	thin layer chromatography
TIm	median tolerance limit
TLV	threshold limit value
TMDI	theoretical maximum daily intake
TMRC	theoretical maximum residue contribution
TMRL	temporary maximum residue limit
TNsG	technical notes for guidance
TOC	total organic carbon
Tremcard	transport emergency card
tRNA	transfer ribonucleic acid
TSH	thyroid stimulating hormone (thyrotropin)

<b>Stand. term/Abbreviation</b>	<b>Explanation</b>
TTC	2,3,5-triphenylterazoliumchloride testing method
TWA	time weighted average
UDS	unscheduled DNA synthesis
UF	uncertainty factor (safety factor)
ULV	ultra low volume
UR	unit risk
UV	ultraviolet
UVC	unknown or variable composition, complex reaction products
UVCB	undefined or variable composition, complex reaction products in biological material
v/v	volume ratio (volume per volume)
vis	visible
WBC	white blood cell
wk	week
wt	weight
w/v	weight per volume
ww	wet weight
w/w	weight per weight
XRFA	X-ray fluorescence analysis
yr	year
<	less than
≤	less than or equal to
>	greater than
≥	greater than or equal to

# **Regulation (EU) No 528/2012 concerning the making available on the market and use of biocidal products**

*Evaluation of active substances*

## **List of References - Part A**



**Polyhexamethylene biguanide**

(Mn = 1600; PDI =1.8)

**(PHMB)**

Applicant: Lonza

Product-types 1, 2, 3, 4, 6, 9, 11

DRAFT FINAL CAR

May 2015

eCA: FRANCE



*This document is a list of all the studies submitted by the Applicant to support the PT1, 2, 3, 4, 6, 9, 11 dossiers. Claims of data protection are proposal from the Applicant.*

*Studies indicated as “Relied on“ are validated studies from which endpoints were established. This corresponds to the list of protected studies.*

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/ IUCLID/ Other	Study relied on
A3_2 (PT1, 3, 4, 6, 11 only)	McGeechan P	2008	Evaluation of the Bactericidal Efficacy of Solid PHMB (EN1276:1997) Arch UK Biocides Microbiology Laboratory, Blackley, Manchester, UK Unpublished; not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-2-05	Other	No
A3_3	Sudworth J	2002	DS6222: Physico-Chemical Data- Project 1270585 Analytical Science Group, Blackley, Manchester, UK Project 1270585 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-3-01	KS	Yes (PT1,2.3.6,9.1 1)
A3_3	Field B.P.	1991	VANTOCIL P: Measurement of selected physical/chemical properties Analytical Science Group, Blackley, Manchester, UK Project 0176 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-3-02	KS	Yes (PT1.2.3.6,9.1 1)
A3_3	Blake J	2003	Product Chemistry and Phys/chemical characteristics study for EPA, Grangemouth solid PHMB. (By analysis of chemical structure and not by experimentation) Analytical Science Group, Blackley, Manchester, UK Project 1273537 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-2-03	KS	Yes (PT1.2.3.6,9.1 1)

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_3	Macnab J.I	2002	Determination of the vapour pressure of poly(hexamethylene)biguanide Syngenta Technology and Projects Process Hazards Section, Huddersfield, UK PC/274 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-3-03	KS	No
A3_3	Bowhill L.	2007	PHMB: Determination of n-Octanol:Water Partition Coefficient InterTek Analytical Science Group, Blackley, Manchester, UK Study 1304881 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-3-04	KS	Yes (PT1.2.3.6,9.1 1)
A3_3	Gillings E, Brown D and Reynolds L F.	1983	The determination of the Octanol-Water Partition Coefficient of Vantocil IB Brixham Environmental Laboratory, Brixham, UK BLS/B/0207 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-3-05	IUCLID	No
A3_3	Schofield D.J	2007	Vantocil 100: Physical Chemical Testing. InterTek Analytical Science Group, Blackley, Manchester, UK Study 1307428 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-3-06	KS	Yes (PT1.2.3.6,9.1 1)
A3_3	Bannon C	2008	Viscosity of VANTOCIL TG Arch Chemicals Inc., Cheshire, USA 112-07B10PHMB Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-3-07	KS	Yes (PT1.2.3.6,9.1 1)
A3_3	Chang S.	2008	Determination of the vapour pressure of Polyhexamethylene Biguanide (PHMB) Arch Chemicals Inc., Cheshire, USA Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-3-08	KS	No

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_3	Bannon C	2008	Melting point of Solid PHMB Arch Chemicals Inc., Cheshire, USA 122-08B10PHMB Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-3-09	KS	No
A3_4	Pickup M.	2002	The extraction and detection of poly(hexamethylenebiguanide) from environmental matrices. Analytical Science Group, Blackley, Manchester, UK Pickup M J Unpublished ; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-4-01	KS	No
A3_4	DeMatteo V A	2008	Validation of the method for determining solution strength for VANTOCIL TG Arch Chemicals Inc, Cheshire, USA 119-08B10PHMB Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-4-02	KS	No
A3_4	Ritter, J.C	2008	INTERIM REPORT: Preliminary Method for the Analysis of PHMB in Drinking Water by Electrochemical Detection with Sample Pre concentration Arch Chemicals Inc, Cheshire, USA  Unpublished; not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-4-03	Other	No
A3_4	Taylor, D.B	2009	Analysis of PHMB in Water by Linear Sweep Stripping Voltammetry, Method Validation. Arch Chemicals Inc, Cheshire, USA  Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-4-04	KS	No

<b>Document/ Section</b>	<b>Author</b>	<b>Year</b>	<b>Description/Title</b>	<b>Owner</b>	<b>Data Protection</b>	<b>Doc IV Code</b>	<b>KS/ IUCLID/ Other</b>	<b>Study relied on</b>
PHMB PT02 B3_5 (PT6 only)	McGeechan P.	2006	Evaluation of the Bacterisostatic and Fungistatic efficacy of VANTOCIL IB. Arch UK Biocides Microbiology Group, Manchester, UK. Report no.004. Not GLP, Unpublished	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I.	PHMB PT02 dossier: ARCH B3-5-04		Yes (PT6)
PT02 IIIB5.10.14	Crane E.	2010	Validation Protocol for Quantitative Suspension Testing for Arch Biocides. MGS Laboratories Ltd., Egham, UK. CVP-2009-014-05 Unpublished, Non-GLP	Arch Chemicals Inc	Yes: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH B3-5-14	KS	Yes (PT2.3.4.9.11)
PT02 IIIB5.10.15	Crane E.	2010	Validation Protocol for Quantitative Suspension Testing for Arch Biocides. MGS Laboratories Ltd., Egham, UK. CVP-2009-014-05 Unpublished, Non-GLP	Arch Chemicals Inc	Yes: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH B3-5-14	KS	Yes (PT2.4.11)
PT02 IIIB5.10.16	Crane E.	2010	Validation Protocol for Quantitative Suspension Testing for Arch Biocides. MGS Laboratories Ltd., Egham, UK. CVP-2009-014-05 Unpublished, Non-GLP	Arch Chemicals Inc	Yes: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH B3-5-14	KS	Yes (PT2,4)
A3_5_02 (B3-5 PT02)	Crane E.	2010	Validation Protocol for Quantitative Suspension Testing for Arch Biocides. MGS Laboratories Ltd., Egham, UK. CVP-2009-014-05 Unpublished, Non-GLP	Arch Chemicals Inc	Yes: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH B3-5-16	KS	Yes (PT3.9)
A3_5	McGeechan P.	2006	PHMB: Mode of Action Arch UK Biocides, Manchester, UK ARCH PHMB 019. Unpublished; not GLP	Arch Chemicals Inc	No	ARCH A3-5-01	Other	Yes (PT1.2.3.11)

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_5	Moore L E.	2004	Evaluation of the risks associated with long term use of cationic antimicrobials Univeristy of Manchester, Manchester, UK ARCH PHMB 020. Unpublished; not GLP	Arch Chemicals Inc	No	ARCH A3-5-02	Other	Yes (PT1.2.3.11)
A3_5	Livermoore D.	2001	MICs of Avecia compounds PUBLIC HEALTH LABORATORY SERVICE CENTRAL PUBLIC HEALTH LABORATORY Antibiotic Resistance Monitoring and Reference Laboratory PHLSCentral Public Health Laboratory 61 Colindale Avenue, London NW9 5HT ARCH PHMB 021. Unpublished; not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-5-03	Other	Yes (PT1.2.3.11)
A3_5	Gilbert P., Moore L.E.	2005	Cationic antiseptics: diversity of action under a common epithet Univeristy of Manchester, Manchester, UK Journal of Applied Microbiology 2005, 99, 703-715 Published; not GLP	Published	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-5-04	Other	Yes (PT1.2.3.4.9.1 1)
A3_5	Moore L.E. <i>et al.</i>	2008	In vitro study of the effect of cationic biocides on bacterial population dynamics and susceptibility Univeristy of Manchester, Manchester, UK Applied and Environmental Microbiology 2008 p. 4825-4834 Published; not GLP	Published	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-5-05	Other	Yes (PT1.2.3.4.9.1 1)

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_5	Tambe S.M. <i>et al.</i>	2001	In vitro evaluation of the risk of developing bacterial resistance to antiseptics and antibiotics used in medical devices Columbia University, New York, USA Journal of Antimicrobial Chemotherapy 2001 47, 589-598 Published; not GLP	Published	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-5-06	Other	Yes (PT1.2.3.4.9.1 1)
A3_5	Turner N.A. <i>et al.</i>	2000	Emergence of resistance to biocides during differentiation of <i>Acanthamoeba castellanii</i> Cardiff University, Cardiff, UK Journal of Antimicrobial Chemotherapy 2000 46, 27-34 Published; not GLP	Published	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-5-07	Other	Yes (PT1.2.3.5.9.1 1)
A3_5	Gilbert P.	No date given	Polyhexamethylene biguanide and infection control Univeristy of Manchester, Manchester, UK www.kendallamd.com Published; not GLP	Published	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-5-08	Other	Yes (PT1.2.3.4.9.1 1)
A3_5	Fraud S. <i>et al.</i>	2008	MexCD-OprJ Multidrug Efflux System of <i>Pseudomonas aeruginosa</i> : Involvement in Chlorhexidine Resistance and Induction by Membrane-Damaging Agents Dependent upon the AlgU Stress Response Sigma Factor Queen's University, Ontario, Canada Antimicrobial Agents and Chemo, Dec 2008, Vol 52, No. 12, p4478-4482 Published; not GLP	Published	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-5-09	Other	Yes (PT1.2.3.4.9.1 1)

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_5	Lakkis C. <i>et al.</i>	2001	Resistance of <i>Pseudomonas aeruginosa</i> Isolates to Hydrogel Contact Disinfection Correlates with Cytotoxicity University of Melbourne, Victoria, Australia Journal of Clinical Microbiology, Apr 2001, Vol 39, No. 4, p1477-1486 Published; not GLP	Published	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-5-10	Other	Yes (PT1.2.3.4.9.1 1)
A3_5	Geraldo I.M. <i>et al.</i>	2008	Rapid antibacterial activity of 2 novel hand soaps: evaluation of the risk of development of bacterial resistance to the antibacterial agents University of Melbourne, Victoria, Australia Infect Control Hosp Epidemiol. 2008 Aug; 29 (8): 736-41 Published; not GLP	Published	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-5-11	Other	Yes (PT1.2.3.4.9.1 1)
A3_5	Allen M.J. <i>et al.</i>	2006	The response of <i>Escherichia coli</i> to exposure to the biocide polyhexamethylene biguanide Cardiff University, Cardiff, UK Microbiology. 2006 Apr; 152 (Pt4): 989-1000 Published; not GLP	Published	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-5-12	Other	Yes (PT1.2.3.4.9.1 1)
A3_5	Khunkitti W. <i>et al.</i>	1998	Biguanide-induced changes in <i>Acanthamoeba castellanii</i> : an electron microscopic study University of Wales Cardiff, Cardiff, UK J Appl Microbiol. 1998 Jan; 84 (1): 53-62 Published; not GLP	Published	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-5-13	Other	Yes (PT1.2.3.4.9.1 1)



Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_5	Turner N.A. <i>et al.</i>	2004	Resistance, biguanide sorption and biguanide-induced pentose leakage during encystment of <i>Acanthamoeba castellanii</i> New York University School of Medicine, New York, USA J Appl Microbiol. 2004; 96 (6): 1287-95 Published; not GLP	Published	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-5-14	Other	Yes (PT1.2.3.4.9.1 1)
A3_5	Pérez-Santonja J.J. <i>et al.</i>	2003	Persistently culture positive <i>Acanthamoeba keratitis</i> : in vivo resistance and in vitro sensitivity Moorfields Eye Hospital, London, UK Ophthalmology. 2003 Aug; 110 (8): 1593-600 Published; not GLP	Published	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-5-15	Other	Yes (PT1.2.3.4.9.1 1)
A3_5	Lloyd D. <i>et al.</i>	2001	Encystation in <i>Acanthamoeba castellanii</i> : development of biocide resistance Cardiff University, Cardiff, UK J Eukaryot Microbiol. 2001 Jan-Feb; 48 (1): 11-6 Published; not GLP	Published	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-5-16	Other	Yes (PT1.2.3.4.9.1 1)
A3_5	Murdoch D. <i>et al.</i>	1998	<i>Acanthamoeba keratitis</i> in New Zealand, including two cases with in vivo resistance to polyhexamethylene biguanide Auckland Hospital, Auckland, New Zealand Aust NZJ Ophthalmol. 1998 Aug; 26 (3): 231-6 Published; not GLP	Published	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-5-17	Other	Yes (PT1.2.3.4.9.1 1)



Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/ IUCLID/ Other	Study relied on
A3_6.1	[REDACTED]	2003	Acute dermal irritation in the rabbit . [REDACTED] Project number: 780/275 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-61-10	KS	No
A3_6.1	[REDACTED]	2003	Acute eye irritation in the rabbit. [REDACTED] Project number: 780/276 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-61-12	KS	No
A3_6.1	[REDACTED]	1993	Polyhexamethylene Biguanide PHMB: Skin sensitisation in the guinea pig of a 20% aqueous solution. [REDACTED] CTL/P/3889. Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-61-16	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.1	Jackson SJ	1979	Vantocil P: Acute Oral and Dermal Toxicity. Central Toxicological Laboratory, Macclesfield, UK CTL/T/1361. Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-61-01	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.1	[REDACTED]	1980	Vantocil P: Skin irritation in the rabbit. [REDACTED] CTL/T/1409 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-61-08	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.1	Jackson SJ	1979	Vantocil P: Skin corrosivity study . Central Toxicological Laboratory, Macclesfield, UK CTL/T/1362 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-61-09	IUCLID	No

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_6.1	[REDACTED]	1980	Vantocil IB: Skin sensitisation studies in the guinea pig [REDACTED] [REDACTED] CTL/T/1423 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-61-17	IUCLID	No
A3_6.1	Jackson SJ	1983	Vantocil IB and Chlorhexidine Gluconate: Potential for cross-reactivity in a skin sensitisation study Central Toxicological Laboratory, Macclesfield, UK CTL/T/1953 Unpublished; not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-61-19	IUCLID	No
A3_6.1	[REDACTED]	1983	Vantocil IB: The effect of variation in induction concentration on skin sensitisation in the guinea pig. [REDACTED] [REDACTED] CTL/T/1952 Unpublished; not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-61-18	IUCLID	No
A3_6.1	Kinch D.A.	1969	The irritant properties of Vantocil IB. Central Toxicological Laboratory, Macclesfield, UK HO/IH/T/704A. Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-61-13	IUCLID	No
A3_6.1	Kinch D.A.	1969	Further Studies on the irritant effects of Vantocil IB. Central Toxicological Laboratory, Macclesfield, UK HO/IH/T/704B. Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-61-14	IUCLID	No

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_6.1	[REDACTED]	1981	Vantocil IB: Eye irritation to the rabbit. [REDACTED] CTL/T/1727. Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-61-11	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.1	[REDACTED]	1993	Baquacil 20% PHMB and Sodium Dichloroisocyanurate: Comparative assessment of sensory irritation potential in the mouse. [REDACTED] CTL/L/5346 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-61-06	KS	No
A3_6.1	Proteau J.	1979	Baquacil SB: Eye irritation French study. Association Pour L'aide Aux Recherches interessant La Medecine Du Travail D8/11 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-61-15	IUCLID	No
A3_6.1	Stevens M.A.	1969	Skin toxicity of Polyhexamethylene biguanide (PHB) solution: Vantocil IB: 20% PHB in water (Antibacterial 9073: 25% PHMB in water) Central Toxicological Laboratory, Macclesfield, UK TR 684 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-61-05	IUCLID	No
A3_6.1	Wnorowski G.	2003	Acute Inhalation Toxicity Feasibility Assessment. Product Safety Laboratories, East Brunswick, New Jersey. OPPTS 870.1300 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-61-07	Other	No

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_6.12	Smith I	1981	Human sensitisation testing of VANTOCIL IB. Ian Smith Consultancy. Project Number 0018; CTL/C/1109. Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-612-01	KS	No
A3_6.12	Hink G, Ison A	1989	Photoreaction patch test using natural sunlight. Hill Top Research, Ohio. Report ref. 76-165-72; CTL/C/2163 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-612-02	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.12	Schnuch A, Geier J, Brasch J et al.	2000	Polyhexamethylene biguanide: A relevant contact allergen? Contact Dermatitis 42:302-3 03 Published; Not GLP	Published	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-612-03	IUCLID	No
A3_6.12	Schnuch A, et al	2007	The biocide polyhexamethylene biguanide remains an uncommon contact allergen. Recent multicentre surveillance data. Contact Dermatitis 2007; 56: 235–239 Published; Not GLP	Published	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-612-04	IUCLID	No
A3_6.12	Geimer P	2007	PHMB: Arch Medical Surveillance Programme Statement from Arch Medical Director dated 23 April 2007 UnPublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-612-05	Other	No
A3_6.14	Sueki H	2001	Polyhexamethylene Biguanide, Cosmocil CQ: Skin Irritation Study in Humans. Dept of Biochemical Toxicology Showa University, Japan. Report APJ-1. Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-614-01	KS	Yes (PT1.2.3.6.9.1 1)

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_6.2	[REDACTED]	1975	Characterisation of the Urinary Polymer-related Material from Rats given Poly[biguanide-1,5-diylhexamethylene hydrochloride] [REDACTED] Makromol. Chem. 177, 2591-2605 Published; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-62-02	IUCLID	No
A3_6.2	Clowes HM	1996	PHMB: In Vitro Absorption through Human Epidermis. Central Toxicological Laboratory, Macclesfield, UK CTL/P/5120. Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-62-03	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.2	Clowes HM	1998	PHMB: In Vitro absorption from a 20% solution through human epidermis at spa temperature. Central Toxicological Laboratory, Macclesfield, UK CTL/P/5916. Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-62-04	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.2	Clowes HM	1995	PHMB: In Vitro Absorption from a 0.5% solution through bovine teat and udder skin . Central Toxicological Laboratory, Macclesfield, UK CTL/P/5683 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-62-06	IUCLID	No

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_6.2	Clowes HM	1997	Development of a method to measure in vitro absorption of chemicals through bovine udder and teat skin. Central Toxicological Laboratory, Macclesfield, UK CTL/L/7823 Unpublished; not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-62-07	Other	No
A3_6.2	Dugard PH, Mawdsley SJ	1982	14C-Polyhexamethylene Biguanide (PHMB): Absorption through human epidermis and rat skin in vitro. Central Toxicological Laboratory, Macclesfield, UK CTL/R/579 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-62-05	IUCLID	Yes (PT1.2.3.6.9.1 1)
A3_6.2	██████████	1976	Studies of Vantocil C14 in Rat and Human Skin. ██████████ D8/35 Unpublished; not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-62-08	IUCLID	No
A3_6.2	██████████	1976	Whole Body Autoradiography of Mice Treated with Vantocil C14. ██████████ Report No 1976_03_03 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-62-09	IUCLID	No
A3_6.2	██████████ ██████████ ██████████	1995	Bioavailability following dietary administration in the rat. ██ ██ CTL/P/4595 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-62-01	KS	Yes (PT1.2.3.6.9.1 1)



Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_6.2	[REDACTED]	1995	PHMB: Absorption, Distribution, Metabolism and Excretion following Single Oral Dosing (20 mg/kg) in the Rat. [REDACTED] [REDACTED] Report No. CTL/P/4537. Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-62-10	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.3	Banham PJ, Marsh DJ	1992	Polyhexamethylene Biguanide: Analysis in dosing solutions. Central Toxicological Laboratory, Macclesfield, UK CTL/I/157 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-63-15	IUCLID	No
A3_6.3	Carney IF	1976	Vantocil IB: Subacute inhalation toxicity. Central Toxicological Laboratory, Macclesfield, UK CTL/T/983 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-63-06	IUCLID	Yes (PT1.2.3.6.9.1 1)
A3_6.3	[REDACTED]	1972	Vantocil IB: Subacute dermal toxicity study in the rabbit. [REDACTED] [REDACTED] CTL/P/22 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-63-04	IUCLID	No
A3_6.3	[REDACTED]	1992	PHMB Polyhexamethylene Biguanide: 28 day drinking water study in the mouse. [REDACTED] [REDACTED] CTL/L/4429 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-63-02	KS	No

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_6.3	[REDACTED]	1992	PHMB: Polyhexamethylene Biguanide: An investigation of its palatability to the mouse in drinking water. [REDACTED] [REDACTED] CTL/L/4843 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-63-13	IUCLID	No
A3_6.3	[REDACTED]	1992	PHMB Polyhexamethylene Biguanide: 28 day drinking water study in the rat. [REDACTED] [REDACTED] CTL/L/4428 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-63-01	KS	No
A3_6.3	[REDACTED]	1993	PHMB: 21 day dermal toxicity study in the rat. [REDACTED] [REDACTED] CTL/P/4200 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-63-03	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.3	Marsh D.L.	1993	PHMB: Gravimetric and homogeneity data to support dietary toxicity studies. Central Toxicological Laboratory, Macclesfield, UK CTL/T/2842 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-63-12	Other	No

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_6.3	[REDACTED]	2006	POLYHEXAMETHYLENE BIGUANIDE: 28 DAY INHALATION STUDY IN RATS WITH RECOVERY [REDACTED] [REDACTED] CTL/MR0219/REGULATORY/REVISION - 001 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-63-05	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.3	[REDACTED]	2006	POLYHEXAMETHYLENE BIGUANIDE: 5 DAY PRELIMINARY INHALATION STUDY IN THE RAT [REDACTED] [REDACTED] MR0218-TEC Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-63-16	IUCLID	No
A3_6.3	[REDACTED]	2006	POLYHEXAMETHYLENE BIGUANIDE: 5 DAY PRELIMINARY INHALATION STUDY IN THE RAT. [REDACTED] [REDACTED] MR0220-TEC Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-63-17	IUCLID	No
A3_6.3	[REDACTED]	1993	6-Week Dietary Toxicity in the Dog [REDACTED] [REDACTED] CTL/L/5227 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-63-10	KS	Yes (PT1.2.3.6.9.1 1)

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/ IUCLID/ Other	Study relied on
A3_6.3	[REDACTED]	1992	Polyhexamethylene Biguanide: Maximum tolerated dose study in the dog. [REDACTED] [REDACTED] CTL/L/4870 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-63-14	IUCLID	No
A3_6.4	[REDACTED] [REDACTED] [REDACTED]	1966	Antibacterial 9073: Ninety-day oral toxicity of antibacterial 9073- Albino rats [REDACTED] [REDACTED] CTL/R/199 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-63-08	IUCLID	No
A3_6.4	[REDACTED] [REDACTED] [REDACTED]	1966	Antibacterial 9073: Ninety-day oral toxicity of antibacterial 9073- beagle dogs [REDACTED] [REDACTED] CTL/R/202 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-63-11	IUCLID	No
A3_6.4	[REDACTED] [REDACTED]	1993	Polyhexamethylene Biguanide PHMB: 90 day oncogenic sighting study in the mouse. [REDACTED] [REDACTED] CTL/T/2825 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-63-09	KS	No
A3_6.4	[REDACTED] [REDACTED]	1993	Polyhexamethylene Biguanide PHMB: 90 day oncogenic sighting study in the rat. [REDACTED] [REDACTED] CTL/T/2824. Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-63-07	KS	Yes (PT1.2.3.6.9.1 1)

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_6.5	[REDACTED]	1977	Baquacil SB: 2-Year Feeding Study in Rats. [REDACTED] [REDACTED] CTL/P/333. Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-65-01	KS	No
A3_6.5	[REDACTED]	1996	Polyhexamethylene Biguanide: Two Year Feeding Study in Rats. Pathology Working Group Peer Review of Proliferative Vascular Lesions in Male & Female Rats. [REDACTED] [REDACTED] CTL/C/3172. Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-65-03	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.5	[REDACTED]	1977	Baquacil SB: Life-Time Feeding Study in the Mouse. [REDACTED] [REDACTED] [REDACTED] CTL/P/332. Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-65-06	KS	No
A3_6.5	[REDACTED]	1996	Polyhexamethylene Biguanide: Two Year Feeding Study in Rats. [REDACTED] [REDACTED] CTL/P/4663. Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-65-02	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.5	[REDACTED]	1993	Polyhexamethylene Biguanide: 2 year drinking water study in the rat. TERMINATED early in week 39 [REDACTED] [REDACTED] CTL/T/2830. Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-65-04	IUCLID	No

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_6.5	[REDACTED]	1995	Polyhexamethylene Biguanide: 1 year dietary toxicity study in the dog. [REDACTED] CTL/P/4488 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-65-07	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.5	Mosinger M.	1973	Prolonged Oral Intake of Vantocil IB Centre D'Explorations et de Recherches Medicales D3/2 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-65-05	IUCLID	No
A3_6.6	[REDACTED]	1981	Vantocil P: Mutation assays using P388 mouse lymphoma cells. [REDACTED] CTL/P/622 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-66-06	KS	No
A3_6.6	Callander R D	1989	Vantocil IB: An evaluation in the Salmonella mutation assay. Central Toxicological Laboratory, Macclesfield, UK CTL/P/2406 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-66-01	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.6	Hastwell RM & McGregor DB.	1979	Testing for mutagenic activity in Salmonella typhimurium Inveresk Research International, Edinburgh, Scotland. IRI 411156 (CTL/C/1720) Unpublished, Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-66-03	IUCLID	No

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_6.6	Howard CA.	1989	Vantocil IB: An evaluation in the in vitro cytogenetic assay in human lymphocytes. Central Toxicological Laboratory, Macclesfield, UK CTL/P/2582 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-66-04	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.6	[REDACTED]	1989	Vantocil IB: An evaluation in the mouse micronucleus test. [REDACTED] [REDACTED] CTL/P/2436 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-66-07	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.6	Richardson CR, Anderson D.	1981	Vantocil P: Cytogenetic study in human lymphocytes in vitro. Central Toxicological Laboratory, Macclesfield, UK CTL/P/613 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-66-05	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.6	Trueman RW	1980	An examination of 'Vantocil' IB for potential carcinogenicity using two in vitro assays. Central Toxicological Laboratory, Macclesfield, UK CTL/P/492	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-66-02	IUCLID	No
A3_6.6	[REDACTED]	1989	Vantocil IB: Assessment for the induction of unscheduled DNA synthesis in rat hepatocytes in vivo. [REDACTED] [REDACTED] CTL/P/2603 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-66-08	KS	Yes (PT1.2.3.6.9.1 1)

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_6.7	[REDACTED]	2002	Historical control data for occurrence of hemangiosarcoma (angiosarcoma) in C57BL/10J/CD-1 Alpk Mice. Supplemental info for CTL/P/4649. [REDACTED] AP-1 Unpublished; not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-67-04	Other	No
A3_6.7	[REDACTED]	2002	Historical control data for occurrence of hemangiosarcoma (angiosarcoma) in Alpk:ApfSD Wistar Rats (re: CTL/P/4663, CTL/C/3172). [REDACTED] AP-5 Unpublished; not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-67-05	Other	No
A3_6.7	[REDACTED]	1996	Polyhexamethylene Biguanide: Two Year Feeding Study in Rats. Pathology Working Group Peer Review of Proliferative Vascular Lesions in Male & Female Rats. [REDACTED] [REDACTED] CTL/C/3172 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-65-03	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.7	[REDACTED] [REDACTED] [REDACTED]	1977	Baquacil SB: 80-week skin painting study in the mouse. [REDACTED] [REDACTED] CTL/P/331 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-67-01	KS	Yes (PT1.2.3.6.9.1 1)



Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_6.7	[REDACTED]	2002	Polyhexamethylene Biguanide (PHMB): Two year Oncogenic Study in Mice. Statistical analysis of the result from the Pathology Working Group peer review of Vascular lesions in male and female mice. Supplemental info for CTL/P/4649. [REDACTED] AP-7 Unpublished; not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-67-06	Other	No
A3_6.7	[REDACTED]	1996	Polyhexamethylene Biguanide: Two Year Feeding Study in Rats. [REDACTED] CTL/P/4663 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-65-02	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.7	[REDACTED]	2002	PHMB 2-year oncogenic study in mice. PWG peer review of vascular proliferative lesions in male and female mice. [REDACTED] EPL Project No 698-001 (= CTL PM0937) Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-67-03	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.7	[REDACTED]	1996	Polyhexamethylene Biguanide: Two year Oncogenic Study in Mice. [REDACTED] CTL/P/4649 Unpublished, GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-67-02	KS	Yes (PT1.2.3.6.9.1 1)

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_6.7	[REDACTED]	2008	Studies to Elucidate the Potential Involvement of the Kupffer Cell in PHMB Mouse Liver Hemangiosarcomas [REDACTED] [REDACTED] 15 Dec 2008 Unpublished, not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-67-07	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.7	Mann P.C, Berry C and Greaves P	2009	Scientific Advisory Panel Review Of Polyhexamethylene Biguanide (Phmb): Carcinogenicity Studies, Pathology Working Groups, Regulatory Responses And Mode-Of-Action Studies  Experimental Pathology Laboratories, Inc. P.O. Box 169, Sterling, VA 20167-0169 EPL STUDY NO. 880-001 5 August 2009 Unpublished, not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-67-08	KS	No
A3_6.8	[REDACTED]	1976	Teratology Evaluation of IL-780 in Rabbits [REDACTED] [REDACTED] FDRL 5022 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-68-04	IUCLID	No
A3_6.8	[REDACTED]	1992	PHMB: Dose range finding study in the rabbit. [REDACTED] [REDACTED] CTL/1/5052 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-68-03	IUCLID	No

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/ IUCLID/ Other	Study relied on
A3_6.8	[REDACTED]	1993	Polyhexamethylene Biguanide PHMB: Dose range finding study in the pregnant rabbit. [REDACTED] [REDACTED] CTL/T/2821 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-68-02	KS	No
A3_6.8	[REDACTED]	1993	PHMB:Developmental toxicity study in the rabbit. [REDACTED] [REDACTED] CTL/P/3997 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-68-01	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.8	Evans DP	1981	Re-evaluation of skeletal variants incorporating historical data. Central Toxicological Laboratory, Macclesfield, UK re: Report CTL/P/335 ReEvaluation Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-68-08	IUCLID	No
A3_6.8	[REDACTED]	1981	Baquacil SB : Mouse Teratology Study (CTL/P/335): Historical control data & clarification of start date. [REDACTED] [REDACTED] re: Report CTL/P/335 Historical Control Data Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-68-09	Other	No
A3_6.8	[REDACTED]	1976	Baquacil SB: A teratology study in the rat by dietary administration. [REDACTED] [REDACTED] [REDACTED] CTL/P/262 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-68-05	KS	Yes (PT1.2.3.6.9.1 1)

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/ IUCLID/ Other	Study relied on
A3_6.8	[REDACTED]	1977	Baquacil SB: Teratogenicity study in the mouse. [REDACTED] [REDACTED] CTL/P/335 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-68-07	IUCLID	No
A3_6.8	[REDACTED]	1995	Polyhexamethylene Biguanide: Multigeneration study in the rat. [REDACTED] [REDACTED] CTL/P/4455 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-68-10	KS	Yes (PT1.2.3.6.9.1 1)
A3_6.8	[REDACTED]	1977	20% PHMB: Three generation reproduction study in the rat CTL/C/2161 Reformatted for EPA 5 July 1990. [REDACTED] Report No. NV-5- L57, Project number 458-119. Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-68-11	IUCLID	No
A3_6.8	[REDACTED]	1988	The Post-natal Fate of Supernumary Ribs in Rat Teratogenicity Studies. [REDACTED] Tox 8 (2) 91-94. Published; GLP unknown	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-68-06	IUCLID	No
A3_7.1.	Brown D., Dowell D.G.	1975	Vantocil IB and sewage treatment Brixham Environmental Laboratory, Brixham, UK BL/B/1649 Unpublished; NOT GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-10	IUCLID	No

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_7.1.	Brown D., Gillings E.	1983	The determination of the partition of Vantocil IB between a river sediment and water Brixham Environmental Laboratory, Brixham, UK BLS/B/0208 Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-14	IUCLID	No
A3_7.1.	██████████	1980	Vantocil IB: Effect of soil on acute toxicity to rainbow trout. ██ ██████████ BLS/B/0044 Unpublished; not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-19	IUCLID	No
A3_7.1.	Evans K.P., Beaumont G.L., Williams D.G.	1995	PHMB Hydrolysis study for EPA Registration: Project 302, Guideline ref. 161-1 (1995) ASG, Blackley, Manchester, UK Project 302 Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-03	IUCLID	No
A3_7.1.	Gilbert J L	1997	PHMB: Determination of COD Brixham Environmental Laboratory, Brixham, UK BLS 2378 Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-01	IUCLID	No
A3_7.1.	Gilbert JL, Long KWJ, Roberts GC	1995	PHMB: Anaerobic biodegradability Brixham Environmental Laboratory, Brixham, UK BL5342/B Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-12	KS	Yes (PT2.9)

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_7.1.	Gilbert JL, Roberts GC, Woods CB	1993	PHMB: Activated sludge sorption and desorption Brixham Environmental Laboratory, Brixham, UK BL5385/B Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-15	KS	Yes (PT2.9)
A3_7.1.	Habeeb. S.B.	2010	PHMB: Aerobic Transformation in Two Aquatic Sediment Systems ABC Laboratories Inc., Missouri, USA 65393 Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-22		Yes (PT2.9)
A3_7.1.	Jones B.K.	1976	Vantocil IB: microbial degradation studies Central Toxicological Laboratory, Macclesfield, UK CTL/P/289 Unpublished; NOT GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-11	IUCLID	No
A3_7.1.	Leahey J.P., Griggs R.E., Hughes H.E.	1975	Baquacil: Preliminary study of the photodegradation in water. ICI Plant Protection Ltd TMJ 1163B Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-05	KS	Yes (PT2.9)
A3_7.1.	Long K.W.J.	1995	PHMB: Aerobic biodegradation in water (adapted microorganisms). Brixham Environmental Laboratory, Brixham, UK BL1878/B Unpublished; not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-07	IUCLID	No
A3_7.1.	Long K.W.J., Roberts G.C.	1994	PHMB: Aerobic biodegradation in water Brixham Environmental Laboratory, Brixham, UK BL5172/B Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-06	KS	Yes (PT2.9)

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_7.1.	O'Malley et al	2006	Biodegradability of end-groups of the biocide polyhexamethylene biguanide (PHMB) assessed using model compounds J Ind Microbiol Biotechnol (2006) 33: 677–684 Published; not GLP	Published	NO	ARCH A3-71-17	IUCLID	Yes (PT2.9)
A3_7.1.	O'Malley et al	2007	Microbial degradation of the biocide polyhexamethylene biguanide: isolation and characterization of enrichment consortia and determination of degradation by measurement of stable isotope incorporation into DNA. Journal of Applied Microbiology ISSN 1364-5072 Published; not GLP	Published	NO	ARCH A3-71-18	IUCLID	Yes (PT2.9)
A3_7.1.	Oteyza T	2007	PHMB: Toxicity to the green alga <i>Selenastrum capricornutum</i> in the presence of treated sewage effluent. Brixham Environmental Laboratory, Brixham, UK BLS/3377/B Unpublished; not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-20	IUCLID	No
A3_7.1.	Penwell A.J., Roberts G.C., Daniel M.	2003	PHMB: Biodegradation by the ligninolytic fungus <i>Phanerochaete chrysosporium</i> (2003) Brixham Environmental Laboratory, Brixham, UK BL6915/B Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-13	IUCLID	No

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_7.1.	Penwell AJ, MacLean SA, Palmer S, Roberts GC	2005	PHMB: Aerobic sewage treatment simulation and chronic toxicity of treated effluent to Daphnia magna Brixham Environmental Laboratory, Brixham, UK BL7802/B Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-09	KS	No
A3_7.1.	Penwell AJ, MacLean SA, Roberts GC	2005	PHMB: Biodegradability in sea water Brixham Environmental Laboratory, Brixham, UK BL7804/B Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-08	KS	Yes (PT2.9)
A3_7.1.	Peou F., Roberts G.C.	2004	PHMB: Effect of sediment on the acute toxicity to Daphnia magna Brixham Environmental Laboratory, Brixham, UK BL7117/B Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-16	KS	Yes (PT2.9)
A3_7.1.	Sarff P.	2010	PHMB: Estimation of the Adsorption Coefficient (K <sub>oc</sub> ) on Soil and/or Sewage Sludge Using High Performance Liquid Chromatography (HPLC) ABC Laboratories Inc., Missouri, USA 65395 Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-21		Yes (PT1.2.3.6.9.1 1)
A3_7.1.	Sudworth J.	2006	PHMB: Hydrolysis as a function of pH InterTek ASG, Blackley, Manchester, UK Project 1302832 Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-02	KS	Yes (PT1.2.3.6.9.1 1)



Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_7.1.	Turner W.R., Ramaswamy H.N.	1979	Baquacil: Hydrolysis/photodegradation study Source: ICI General Analysis Group, Analytical and Physical Chemistry Section Ref: R5 Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-71-04	IUCLID	No
A3_7.2	Gilbert JL, Gillings EG, Roberts GC	1995	PHMB: Aerobic biodegradation in soil Brixham Environmental Laboratory, Brixham, UK BL5311/B Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-72-01	KS	Yes (PT1.2.3.6.9.1 1)
A3_7.2	Habeeb. S.B.	2010	PHMB: Determination of Adsorption – Desorption Using the Batch Equilibrium Method ABC Laboratories Inc., Missouri, USA 65392 Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-72-05		Yes (PT1.2.3.6.9.1 1)
A3_7.2	Habeeb. S.B.	2010	PHMB: Aerobic Transformation in Four Soils ABC Laboratories Inc., Missouri, USA 65394 Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-72-06		Yes (PT1.2.3.6.9.1 1)
A3_7.2	Hill I.R, Willis J.H	1975	BAQUACIL: Preliminary laboratory studies of the degradation of C14-BAQUACIL in soil Jealott's Hill Research Station, Bracknell, Berkshire, UK TMJ 1165 Unpublished; not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-72-03	IUCLID	No
A3_7.2	Jones-Hughes TL, Penwell A J, Roberts GC	2005	PHMB: Biodegradation in sludge amended soil Brixham Environmental Laboratory, Brixham, UK BL7132/B Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-72-02	KS	Yes (PT1.2.3.6.9.1 1)

Document/ Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/ IUCLID/ Other	Study relied on
A3_7.2	Riley D., Stevens J.E.	1975	Baquacil: Adsorption and leaching in soil. ICI Plant Protection. Report AR 2586A Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-72-04	KS	Yes (PT2.9)
A3_7.3	Ritter, J.C	2006	Estimation of Photochemical Degradation of Polyhexamethylene Biguanide (PHMB) Using the Atkinson Calculation Method Central Analytical Department, Chesire USA CASR-03-2006 Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-73-01	KS	Yes (PT1.2.3.6.9.1 1)
A3_7.4	Brown D	1985	Toxicity to Brown shrimp (Crangon crangon) of Vantocil IB Brixham Environmental Laboratory, Brixham, UK BL/B/2630 Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-13	IUCLID	No
A3_7.4	Brown D	1981	Effect of Vantocil on the reproduction of Daphnia magna Brixham Environmental Laboratory, Brixham, UK BLS/B/0042 Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-27	IUCLID	No
A3_7.4	██████	1981	Determination of the acute toxicity of Vantocil P to Rainbow Trout (Salmo gairdneri) ██ ██ BL/B/2081 Unpublished; Not GLP but QA'd	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-02	IUCLID	No

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/ IUCLID/ Other	Study relied on
A3_7.4	Brown D.	1981	Toxicity to the green alga ( <i>Scenedesmus quadricauda</i> ) of Vantocil IB (1981) summary only Brixham Environmental Laboratory, Brixham, UK BLS/B/0043 Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-19	IUCLID	No
A3_7.4	[REDACTED]	1980	Vantocil P: Acute tox to rainbow trout [REDACTED] Plaice BL/B/2031 Unpublished; Not GLP but QA'd	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-03	IUCLID	No
A3_7.4	[REDACTED]	1977	Acute toxicity of Vantocil IB, mix No 1857, to Bluegill ( <i>Lepomis macrochirus</i> ) and the water flea ( <i>Daphnia magna</i> ) [REDACTED] CTL/C/3039 Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-10	IUCLID	No
A3_7.4	[REDACTED]	1988	Vantocil IB: Acute tox to rainbow trout [REDACTED] BLS/B/0532 Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-04	IUCLID	No
A3_7.4	Gilbert JL, Roberts GC	2002	PHMB: Toxicity to the sediment dwelling larvae <i>Chironomus riparius</i> Brixham Environmental Laboratory, Brixham, UK BL7135/B Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-28	KS	Yes (PT1.2.3.6.9.1 1)

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/ IUCLID/ Other	Study relied on
A3_7.4	Gillings E.	1995	PHMB: Prelim. Investigation of the effects of pH on sorption to glass. Brixham Environmental Laboratory, Brixham, UK BLS1937/B Unpublished; not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-30	IUCLID	No
A3_7.4	[REDACTED]	1975	Determination of the acute toxicity to Rainbow Trout of Vantocil IB in freshwater. [REDACTED] [REDACTED] BL/B/1631 Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-05	IUCLID	No
A3_7.4	Hutchinson T.H.	1993	Vantocil IB: Acute Toxicity to marine polychaete Platynereis dumerilii Brixham Environmental Laboratory, Brixham, UK BL4953/B Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-15	IUCLID	No
A3_7.4	Hutchinson T.H., Jha A.N	1993	Vantocil IB: Effects on fertilisation in marine polychaete Platynereis dumerilii. Brixham Environmental Laboratory, Brixham, UK BL5003/B Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-16	IUCLID	No
A3_7.4	Hutchinson T.H., Jha A.N	1993	Vantocil IB: Effects on embryo development in a polychaete. Brixham Environmental Laboratory, Brixham, UK BL5004/B Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-17	IUCLID	No

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_7.4	[REDACTED]	1991	Vantocil IB: Effects on survival and growth of sheepshead minnow ( <i>Cyprinodon variegatus</i> ) larvae [REDACTED] BL4351/B Unpublished; Not ? GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-25	IUCLID	No
A3_7.4	Maddock B.G.	1983	Vantocil IB: Toxicity to brown shrimp Brixham Environmental Laboratory, Brixham, UK BLS/B/0211 Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-14	IUCLID	No
A3_7.4	Maddock BG	1983	Toxicity to Plaice ( <i>Pleuronectes platessa</i> ) of Vantocil IB Brixham Environmental Laboratory, Brixham, UK BLS/B/0210 Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-07	IUCLID	No
A3_7.4	Mather J.I.	1988	VANTOCIL IB: Bacterial Growth inhibition ( <i>P.putida</i> ) Brixham Environmental Laboratory, Brixham, UK BLS/B/0558 Unpublished; not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-23	IUCLID	No
A3_7.4	Pearson CR	1981	Acute toxicity of Vantocil IB to <i>Daphnia magna</i> (1981) summary only Brixham Environmental Laboratory, Brixham, UK BLS/B/0041 Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-11	KS	Yes (PT2.9)

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_7.4	Penwell A.J.	2006	PHMB: Chronic toxicity to Daphnia magna Brixham Environmental Laboratory, Brixham, UK BL8365/B Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-26	KS	Yes (PT1.2.3.6.9.1 1)
A3_7.4	Penwell A.J., Roberts G.C.	2000	VANTOCIL IB: Inhibition of anaerobic gas production from sewage sludge Brixham Environmental Laboratory, Brixham, UK BL6914/B Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-20	KS	Yes (PT1.2.3.6.9.1 1)
A3_7.4	Penwell A.J., Smyth D.V.	2006	PHMB: Toxicity to the green alga Selenastrum capricornutum Brixham Environmental Laboratory, Brixham, UK BL8161/B Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-18	KS	Yes (PT1.2.3.6.9.1 1)
A3_7.4	██████████ ██████████	1996	PHMB: Acute toxicity to rainbow trout (Oncorhynchus mykiss) ██ ██████████ BL5506/B Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-01	KS	Yes (PT1.2.3.6.9.1 1)
A3_7.4	██████████ ██████████	2004	PHMB: Summary of rangefinding data in Rainbow trout static and flowthrough test systems. ██ ██████████ BL/B/2976 Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-06	IUCLID	No

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_7.4	Penwell AJ, Roberts GC	2000	VANTOCIL IB: Inhibition of nitrification of activated sludge microorganisms Brixham Environmental Laboratory, Brixham, UK BL6913/B Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-21	KS	Yes (PT1.2.3.6.9.1 1)
A3_7.4	Penwell AJ, Roberts GC	2000	VANTOCIL IB: Effect on the respiration rate of activated sludge Brixham Environmental Laboratory, Brixham, UK BL6678/B OECD 209 Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-22	IUCLID	No
A3_7.4	[REDACTED]	2001	PHMB: Effects on growth of juvenile rainbow trout (Oncorhynchus mykiss) [REDACTED] BL7096/B Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-24	KS	Yes (PT1.2.3.6.9.1 1)
A3_7.4	Roberts GC	2004	[14C] PHMB: Evaluation of Sorption to Various Storage Vessels. Brixham Environmental Laboratory, Brixham, UK BLS3110/B Unpublished; not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-31	IUCLID	No
A3_7.4	[REDACTED]	1993	Study X022/B, Vantocil IB: acute toxicity to Bluegill sunfish (Lepomis macrochirus) [REDACTED] BL4778/B Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-09	IUCLID	No

Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_7.4	[REDACTED]	1981	Acute toxicity of Vantocil P to Bluegill (Lepomis macrochirus) [REDACTED] [REDACTED] BW-81-3-847 Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-08	IUCLID	No
A3_7.4	Stewart K.M., Thompson R.S.	1991	Vantocil IB: Acute toxicity to mysid shrimp (Mysidopsis bahia) summary only Brixham Environmental Laboratory, Brixham, UK BL4365/B	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-12	IUCLID	No
A3_7.4	Thompson RS	1983	The effect of Vantocil P on the growth of Lemna minor (Duckweed) Brixham Environmental Laboratory, Brixham, UK BLS/B/0225 Unpublished; Not GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-74-29	IUCLID	No
A3_7.5	[REDACTED]	1979	Baquacil Mix #5889. Acute Oral LD50 - Mallard Duck. MRID No: 27491 + Phase 3 Summary of MRID 27491. Guideline reference 71-1: Acute dietary LD50 test for waterfowl. [REDACTED] Project No 123-131 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-75-09	KS	Yes (PT1.2.3.6.9.1 1)



Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_7.5	██████████	1979	Baquacil Mix #5889. Eight day dietary LC50 Bobwhite Quail MRID No: 41382 + Phase 3 Summary of MRID 41382. Guideline reference 71-2: Acute dietary LC50 test for upland game birds ██ Project No 123-129 Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-75-10	IUCLID	No
A3_7.5	██████████	1979	Baquacil Mix #5889. Eight day dietary LC50 Mallard Duck. Final report. MRID No: 27492 ██ Project No 123-130 Unpublished; Not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-75-11	IUCLID	No
A3_7.5	Gilbert JL, Roberts GC	2002	PHMB: Acute toxicity to the earthworm Eisenia foetida Brixham Environmental Laboratory, Brixham, UK BL7134/B Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-75-02	KS	Yes (PT1.2.3.6.9.1 1)
A3_7.5	Penwell AJ, Roberts GC	2003	PHMB: Effect on nitrogen transformation by soil microorganisms Brixham Environmental Laboratory, Brixham, UK BL7133/B OECD 216 Unpublished; GLP	Arch Chemicals Inc	YES Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-75-01	KS	Yes (PT2.9)
A3_7.5	Penwell AJ, Roberts GC	2002	PHMB: Effect on seedling emergence and growth Brixham Environmental Laboratory, Brixham, UK BL7131/B Unpublished; GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-75-05	KS	Yes (PT1.2.3.6.9.1 1)

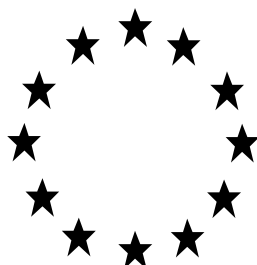
Document/Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/IUCLID/Other	Study relied on
A3_7.5	Stanley R.D.	1983	The effect of Vantocil P on the Earthworm ( <i>Lumbricus terrestris</i> ) Brixham Environmental Laboratory, Brixham, UK BLS/B/0224 Unpublished; not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-75-03	IUCLID	No
A3_7.5	Stanley R.D.	1983	The effect of Vantocil P on the germination and growth of <i>Lepidium sativum</i> (Cress) seeds Brixham Environmental Laboratory, Brixham, UK BLS/B/0222 Unpublished; not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-75-06	IUCLID	No
A3_7.5	Stanley R.D.	1983	The effect of Vantocil P on the germination and growth of <i>Avena sativa</i> (Oat) seeds Brixham Environmental Laboratory, Brixham, UK BLS/B/0223 Unpublished; not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-75-07	IUCLID	No
A3_7.5	Stanley R.D., Tapp J.F.	1981	The effects of Synperonic NP8, Vantocil P, and Chlordane on <i>Lumbricus Terrestris</i> and <i>Allolobophora Caliginosa</i> . Brixham Environmental Laboratory, Brixham, UK BL/A/2111 Unpublished; not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-75-04	IUCLID	No

Document/ Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/ IUCLID/ Other	Study relied on
A3_7.5	Stanley R.D., Tapp J.F.	1981	The Effects of Synperonic NP8, Vantocil P, and Potassium Chlorate on the growth of Avena Sativa. . Brixham Environmental Laboratory, Brixham, UK BL/A/2136 Unpublished; not GLP	Arch Chemicals Inc	YES: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH A3-75-08	IUCLID	No

# **Regulation (EU) No 528/2012 concerning the making available on the market and use of biocidal products**

*Evaluation of active substances*

## **List of References – Part B**



**Polyhexamethylene biguanide**

(Mn = 1600; PDI =1.8)

**(PHMB)**

Applicant: Lonza

**Product-type 1: Human hygiene**

FINAL CAR

June 2015

eCA: FRANCE

*This document is a list of all the studies submitted by the Applicant to support the PT01 dossier. Claims of data protection are proposal from the Applicant.*

*Studies indicated as “Relied on“ are validated studies from which endpoints were established. This corresponds to the list of protected studies.*

Document/ Section	Author	Year	Description/Title	Owner	Data Protection	Doc IV Code	KS/ IUCLID/ Other	Study relied on
B3_5	McGeechan P.	2007 Revised 2008	Evaluation of the Bactericidal Efficacy of VANTOCIL TG. Arch UK Biocides Microbiology Group, Manchester, UK. Report no. 024. Unpublished, Not GLP	Arch Chemicals Inc	Yes: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH B3-5-01		No
B3_5	McGeechan P.	2005 Revised 2008	Evaluation of the Bactericidal Efficacy of an Antibacterial Handwash Formulation. Arch UK Biocides Microbiology Group, Manchester, UK. Report no. 026. Unpublished, Not GLP	Arch Chemicals Inc	Yes: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH B3-5-02		No
B3_5	McGeechan P.	2007 Revised 2008	Evaluation of the Bactericidal Efficacy of VANTOCIL TG. Arch UK Biocides Microbiology Group, Manchester, UK. Report no. 027. Unpublished, Not GLP	Arch Chemicals Inc	Yes: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH B3-5-03		No
B3_5	Hall K.	2003	Investigation into the effectiveness of DS6259 handwash formulation in accordance with EN1499:1997. Arch UK Biocides Microbiology Group, Manchester, UK. Report no. 028. Unpublished, Not GLP	Arch Chemicals Inc	Yes: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH B3-5-04	KS	Yes
B3_5	Crane E.	2009	Validation Protocol for Quantitative Disinfectant Suspension Testing for Arch Biocides. Doc No. CVP-2009-014-05	Arch Chemicals Inc	Yes: Data on existing a.s. submitted for the first time for entry into Annex I	ARCH B3-5-14	KS	Yes