

CONCLUSION ON PESTICIDE PEER REVIEW

Conclusion on the peer review of the pesticide risk assessment of the active substance potassium hydrogen carbonate¹

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SUMMARY

Potassium hydrogen carbonate is one of the 295 substances of the fourth stage of the review programme covered by Commission Regulation (EC) No 2229/2004³, as amended by Commission Regulation (EC) No 1095/2007⁴.

Potassium hydrogen carbonate was included in Annex I to Directive 91/414/EEC on 1 September 2009 pursuant to Article 24b of the Regulation (EC) No 2229/2004 (hereinafter referred to as 'the Regulation') and has subsequently been deemed to be approved under Regulation (EC) No 1107/2009⁵, in accordance with Commission Implementing Regulation (EU) No 540/2011⁶, as amended by Commission Implementing Regulation (EU) No 541/2011⁷. In accordance with Article 25a of the Regulation, as amended by Commission Regulation (EU) No 114/2010⁸, the European Food Safety Authority (EFSA) is required to deliver by 31 December 2012 its view on the draft review report submitted by the European Commission in accordance with Article 25(1) of the Regulation. This review report was established as a result of the initial evaluation provided by the designated rapporteur Member State in the Draft Assessment Report (DAR). The EFSA therefore organised a peer review of the DAR. The conclusions of the peer review are set out in this report.

Ireland being the designated rapporteur Member State submitted the DAR on potassium hydrogen carbonate in accordance with the provisions of Article 22(1) of the Regulation, which was received by the EFSA on 2 May 2006. The peer review was initiated on 16 May 2008 by dispatching the DAR for consultation of the Member States and the original notifier Brotherton Speciality Products Limited (the notifier subsequently changed and is now Church & Dwight UK Ltd). Following consideration of the comments received on the DAR, it was concluded that there was no need to conduct an expert consultation and EFSA should deliver its conclusions on potassium hydrogen carbonate.

The conclusions laid down in this report were reached on the basis of the evaluation of the representative uses of potassium hydrogen carbonate as a fungicide on apple and grapevines, as

On request from the European Commission, Question No EFSA-Q-2009-00256, approved on 16 December 2011.

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³ OJ L 379, 24.12.2004, p.13

⁴ OJ L 246, 21.9.2007, p.19

⁵ OJ L 309, 24.11.2009, p.1

⁶ OJ L 153, 11.6.2011, p.1

OJ L 153, 11.6.2011, p.187

⁸ OJ L 37, 10.2.2010, p.12

Suggested citation: European Food Safety Authority; Conclusion on the peer review of the pesticide risk assessment of the active substance potassium hydrogen carbonate. EFSA Journal 2012;10(1):2524. [36 pp.] doi:10.2903/j.efsa.2012.2524. Available online: www.efsa.europa.eu/efsajournal



proposed by the notifier. Full details of the representative uses can be found in Appendix A to this report.

Data gaps were identified for the section on physical and chemical properties of the formulation.

No critical areas of concern or data gaps were identified in the toxicology section.

Metabolism and residue studies were considered not necessary and not relevant for the evaluation, due to the nature and properties of the active substance. The setting of MRLs for potassium hydrogen carbonate associated with its use as a plant protection product is considered not necessary and a quantitative consumer risk assessment was not conducted.

Potassium hydrogen carbonate is a naturally occurring inorganic compound that dissociates to K^+ and HCO_3^- in the presence of water. A data gap has been identified for studies or peer reviewed scientific literature to support the naturally occurring levels of potassium proposed by the notifier.

Based on the limited data set available, the risk to birds and to biological methods of sewage treatment was assessed as low for the representative uses. Several data gaps were identified in relation to the assessments for non-target terrestrial vertebrates, for aquatic organisms, for honeybees, for non-target arthropods, for soil non-target organisms and for terrestrial non-target plants. Based on the available data, a high risk for honeybees was identified as a critical area of concern

KEY WORDS

Potassium hydrogen carbonate, peer review, risk assessment, pesticide, fungicide



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BACKGROUND

Potassium hydrogen carbonate is one of the 295 substances of the fourth stage of the review programme covered by Commission Regulation (EC) No 2229/2004⁹, as amended by Commission Regulation (EC) No 1095/2007¹⁰.

Potassium hydrogen carbonate was included in Annex I to Directive 91/414/EEC on 1 September 2009 pursuant to Article 24b of the Regulation (EC) No 2229/2004 (hereinafter referred to as 'the Regulation'), and has subsequently been deemed to be approved under Regulation (EC) No 1107/2009¹¹, in accordance with Commission Implementing Regulation (EU) No 540/2011¹², as amended by Commission Implementing Regulation (EU) No 541/2011¹³. In accordance with Article 25a of the Regulation, as amended by Commission Regulation (EU) No 114/2010¹⁴ the European Food Safety Authority (EFSA) is required to deliver by 31 December 2012 its view on the draft review report submitted by the European Commission in accordance with Article 25(1) of the Regulation (European Commission, 2008). This review report was established as a result of the initial evaluation provided by the designated rapporteur Member State in the Draft Assessment Report (DAR). The EFSA therefore organised a peer review of the DAR. The conclusions of the peer review are set out in this report.

Ireland being the designated rapporteur Member State submitted the DAR on potassium hydrogen carbonate in accordance with the provisions of Article 22(1) of the Regulation, which was received by the EFSA on 2 May 2006 (Ireland, 2006). The peer review was initiated on 16 May 2008 by dispatching the DAR to the notifier original Brotherton Speciality Products Limited (the notifier subsequently changed and is now Church & Dwight UK Ltd), and on 24 February 2011 to the Member States, for consultation and comments. In addition, the EFSA conducted a public consultation on the DAR. The comments received were collated by the EFSA and forwarded to the RMS for compilation and evaluation in the format of a Reporting Table. The notifier was invited to respond to the comments in column 3 of the Reporting Table. The comments were evaluated by the RMS in column 3 of the Reporting Table.

The scope of the peer review was considered in a telephone conference between the EFSA, the RMS, and the European Commission on 20 June 2011. On the basis of the comments received and the RMS's evaluation thereof it was concluded that there was no need to conduct an expert consultation.

The outcome of the telephone conference, together with EFSA's further consideration of the comments is reflected in the conclusions set out in column 4 of the Reporting Table. All points that were identified as unresolved at the end of the comment evaluation phase and which required further consideration, and additional information to be submitted by the notifier, were compiled by the EFSA in the format of an Evaluation Table.

The conclusions arising from the consideration by the EFSA, and as appropriate by the RMS, of the points identified in the Evaluation Table, together with the outcome of the expert discussions where these took place, were reported in the final column of the Evaluation Table.

A final consultation on the conclusions arising from the peer review of the risk assessment took place with Member States via a written procedure in November – December 2011.

This conclusion report summarises the outcome of the peer review of the risk assessment on the active substance and the representative formulation evaluated on the basis of the representative uses as a

⁹ OJ L 379, 24.12.2004, p.13

¹⁰ OJ L 246, 21.9.2007, p.19

¹¹ OJ L 309, 24.11.2009, p.1

¹² OJ L 153, 11.6.2011, p.1

¹³ OJ L 153, 11.6.2011, p.187

¹⁴ OJ L 37, 10.2.2010, p.12



fungicide on apple and grapevines, as proposed by the notifier. A list of the relevant end points for the active substance as well as the formulation is provided in Appendix A. In addition, a key supporting document to this conclusion is the Peer Review Report, which is a compilation of the documentation developed to evaluate and address all issues raised in the peer review, from the initial commenting phase to the conclusion. The Peer Review Report (EFSA, 2011) comprises the following documents, in which all views expressed during the course of the peer review, including minority views, can be found:

- the comments received on the DAR,
- the Reporting Table (21 June 2011),
- the Evaluation Table (12 December 2011),
- the comments received on the assessment of the points of clarification,
- the comments received on the draft EFSA conclusion.

Given the importance of the DAR including its addendum (compiled version of December 2011 containing all individually submitted addenda (Ireland, 2011)) and the Peer Review Report, both documents are considered respectively as background documents A and B to this conclusion.



THE ACTIVE SUBSTANCE AND THE FORMULATED PRODUCT

Potassium bicarbonate is the ISO common name for potassium hydrogen carbonate (IUPAC).

The representative formulated product for the evaluation was 'Armicarb 85SP', a water soluble powder (SP), containing 850 g/kg potassium hydrogen carbonate.

The representative uses evaluated comprise spray applications to apple and grapevines as a fungicide against vine powdery mildew and apple scab. Full details of the GAP can be found in the list of end points in Appendix A.

CONCLUSIONS OF THE EVALUATION

1. Identity, physical/chemical/technical properties and methods of analysis

The following guidance documents were followed in the production of this conclusion: SANCO/3030/99 rev.4 (European Commission, 2000) and SANCO/825/00 rev. 7 (European Commission, 2004a).

The minimum purity of potassium hydrogen carbonate is 995 g/kg. No FAO specification exists.

During the peer review it was concluded that lead and arsenic should be considered relevant impurities in potassium hydrogen carbonate used as a plant protection product, with maximum limits of 10 mg/kg of lead and 3 mg/kg of arsenic (see section 2). The assessment of the data package revealed no issues that need to be included as critical areas of concern with respect to the identity, physical, chemical and technical properties of potassium hydrogen carbonate or the representative formulation, however, data gaps were identified for the wettability and foam persistence of the formulation. The main data regarding the identity of potassium hydrogen carbonate and its physical and chemical properties are given in Appendix A.

Adequate analytical methods are available for the determination of potassium hydrogen carbonate in the technical material and in the representative formulation.

The need for methods of analysis for monitoring this compound in food of plant and animal origin and in the environment have been waived due to the nature of the compound. A method for body fluids and tissues is not required as the active substance is not classified as toxic or very toxic.

2. Mammalian toxicity

The following guidance documents were followed in the production of this conclusion: SANCO/221/2000 rev. 10 - final (European Commission, 2003), SANCO/222/2000 rev. 7 (European Commission, 2004b) and SANCO/10597/2003 – rev. 8.1 (European Commission, 2009).

Lead and arsenic are considered relevant impurities their maximum content being 10 mg/kg and 3 mg/kg respectively.

In experimental animals low acute toxicity of potassium hydrogen carbonate is observed in rats via the oral, dermal and inhalation routes. It is neither a skin nor an eye irritant nor a skin sensitizer. In rats, alkalogenic diets have been associated with adverse effects in adrenals (hyperthrophy of zona glomerulosa) and urinary bladder (hyperplasia, papiloma and carcinoma) through well-recognised mechanisms not considered relevant for humans.

No suitable data are available to set reference values. However, it should be taken into account that potassium hydrogen carbonate is a major constituent of normal human physiology. The recommended



daily intakes for potassium are in the order of 3.5 g/adult/day. Potassium hydrogen carbonate is food grade, as per Food Chemical Codex, and it is used in over-the-counter antacid preparations for humans.

A quantitative risk assessment has been performed by the RMS comparing the non-dietary exposure to potassium hydrogen carbonate arising from the use as a plant protection product with normal dietary intakes of potassium (3.5 g/adult/day, equivalent to 128 mg potassium hydrogen carbonate/kg bw/day) indicating that predicted estimates for operators, workers and bystanders will not impact on potassium balance in the human body.

3. Residues

The assessment in the residue section below is based on the guidance documents listed in the document 1607/VI/97 rev.2 (European Commission, 1999), and the JMPR recommendations on livestock burden calculations stated in the 2004 and 2007 JMPR reports.

Metabolism and residue studies were not considered relevant for the evaluation of the active substance potassium hydrogen carbonate, which dissociates in plants in the presence of water to produce the ions K⁺ and HCO₃⁻, which are naturally occurring in all environmental compartments, including plant tissues. It is therefore not possible to distinguish between the residues arising from the use of potassium hydrogen carbonate as a plant protection product and its natural presence in plants.

Moreover potassium hydrogen carbonate is approved as bicarbonate as a food additive in the EU (E501) and is also registered as an ingredient in pharmaceutical preparations. The intake of potassium hydrogen carbonate through use as a plant protection product will be negligible compared with that through normal consumption of the food additive or pharmaceutical preparations.

Consequently, the setting of MRLs for potassium hydrogen carbonate associated with its use as a plant protection product is considered not necessary and a quantitative consumer risk assessment was not conducted. Potassium hydrogen carbonate could be considered a candidate for the inclusion in Annex IV of Commission Regulation (EC) No 396/2005¹⁵.

4. Environmental fate and behaviour

No studies on the fate and behaviour of potassium bicarbonate in the environment are available in the dossier.

Potassium hydrogen carbonate is a naturally occurring inorganic compound that dissociates to K^+ and HCO_3^- in the presence of water. The notifier proposed a waiver for additional fate studies on the basis of the naturally occurring background levels of K^+ and HCO_3^- in soil and water. Whereas the levels of HCO_3^- have been documented with scientific peer reviewed literature, the levels of K^+ have been provided from an internet link that does not contain references to any scientific peer reviewed study. Therefore, a data gap has been identified for studies or peer reviewed scientific literature to support the naturally occurring levels of potassium proposed by the notifier.

Worst case initial PEC soil and PEC SW have been calculated for the representative uses taking into account the maximum application rate of 5.1 kg a.s./ha. These PECs have been used in the ecotoxicological risk assessment.

¹⁵ OJ L 70, 16.3.2005, p. 16



5. Ecotoxicology

The risk assessment was based on the following documents: European Commission 2002a, 2002b and 2002c.

To support the risk assessment for birds, toxicological endpoints only from study summaries from the open literature could be derived. In these long-term feeding studies, where chickens were fed with high concentrations of sodium hydrogen carbonate or potassium hydrogen carbonate in the diet, no adverse effects were observed. It is noted that poultry are often fed with sodium hydrogen carbonate at a typical dietary concentration of 0.2 %. The available risk assessments for wild birds are based on the endpoint derived from one of the studies conducted with sodium hydrogen carbonate and contain several assumptions and extrapolations, and therefore include some uncertainties. It was however concluded that the risk to birds from the use of potassium hydrogen carbonate as a plant protection product based on the representative uses is low. Only acute endpoints were available for the risk assessment for mammals. These first tier assessments indicated a high risk to wild mammals. No higher tier data and assessments (e.g. refinements of the standard exposure scenarios) were available. However the peer-review considered that the acute risk to non-target vertebrates, such as wild mammals, arising from the representative use of potassium hydrogen carbonate is low. For this conclusion a weight of evidence approach was used considering the nature of the active substance and that its dissociation products are widespread elements of the environment, therefore wildlife will often be exposed to them. Moreover the available toxicity data on vertebrates (mammals and birds) and the risk assessments on birds were also taken into consideration. No risk assessments for long-term scale were available. Since repeated and long-term exposure of wild mammals arising from the representative uses, a data gap was identified for long-term risk assessment for wild mammals.

Risk assessments for **aquatic organisms** based on the available acute data for fish and daphnia, and considering a worst case approach via spray drift exposure of the aquatic environment, resulted in a low risk. No chronic data and long-term risk assessments were available. It is noted that the background concentrations of the dissociation products of potassium hydrogen carbonate in natural aquatic systems was assumed to be relatively high compared to the predicted concentrations arising from the application of the active substance, although there is a data gap identified for data to support the high background concentrations of K⁺ (see section 4). If the background levels are confirmed to be higher than the exposure from the representative uses then the long-term risk could be considered as low, however, a data gap to re-consider the risk assessment has been identified pending the availability of the necessary data in section 4. No data and no associated risk assessments were available for algae therefore a data gap was identified to address these assessments. No data were available for the representative formulation, which contains relatively toxic co-formulants and is therefore likely to pose a higher risk to aquatic organisms when compared with the active substance. A data gap was therefore identified for appropriate risk assessments for the representative formulation and aquatic organisms.

No data for acute oral toxicity to **honeybees** were available. Considering however that the dissociation products of potassium hydrogen carbonate are common elements of the environment, the peer-review agreed that no risk assessments for the oral exposure of bees are necessary. Risk assessments for the contact route of exposure (e.g. over spraying of the honeybees) resulted in a high risk. No reliable higher tier data were available therefore a data gap was set to further address the risk assessment for honeybees. This was identified as a critical area of concern. No data were available for the representative formulation therefore a data gap was set for appropriate risk assessments for the representative formulation to honeybees.

No data or risk assessments were available for non-target arthropods, earthworms, soil macro- and micro- organisms or for terrestrial non-target plants. It is noted that the background concentrations of the dissociation products of potassium hydrogen carbonate in soil was assumed to be relatively high compared to the predicted concentrations arising from the application of the active substance, although there is a data gap identified for data to support the high background concentrations of K^+ (see section



4). If the background levels are confirmed to be higher than the exposure from the representative uses then the risk could be considered as low, however, a data gap to re-consider the risk assessments has been identified pending the availability of the necessary data in section 4.

The risk to the biological methods for sewage treatments for the representative uses of potassium hydrogen carbonate was considered to be low.



6. Overview of the risk assessment of compounds listed in residue definitions triggering assessment of effects data for the environmental compartments

6.1. Soil

Compound (name and/or code)	Persistence	Ecotoxicology
Potassium hydrogen carbonate	Naturally occurring inorganic compound applied at levels assumed to be in the range of natural occurrence. Data gap identified for the natural background levels of K^+ .	NO data or risk assessment is available for non-target soil organisms. Data gap pending on the information on

6.2. Ground water

Compound (name and/or code)	Mobility in soil	$>$ 0.1 μ g/L 1m depth for the representative uses (at least one FOCUS scenario or relevant lysimeter) ^(a)	Pesticidal activity	Toxicological relevance	Ecotoxicological activity	
Potassium hydrogen carbonate	Not assessed. Naturally occurring inorganic compound applied at levels assumed to be in the range of natural occurrence. Data gap identified for the natural background levels of K ⁺ .	Not assessed. Naturally occurring inorganic compound applied at levels assumed to be in the range of natural occurrence. Data gap identified for the natural background levels of K ⁺ .	Yes	Not assessed.	No data were available for long-term scale for aquatic organisms. Data gap pending on the information on the background level of K ⁺ . The acute risk for fish and daphnia was assessed as low.	

EFSA Journal 2012;10(1):2524



(a): EFSA's reading of the Council Directive 98/83/EC¹⁶ on the quality of drinking water intended for human consumption is that, as an inorganic fungicide, potassium hydrogen carbonate or the relevant ions that are formed from it, are not considered a pesticide under this directive, so the parametric drinking water limit of 0.1µg/L for pesticides, usually used as a decision making criteria regarding groundwater exposure, does not apply. 'Chemical parameters' or 'indicator parameters' levels (as defined in this directive) have not been prescribed for potassium or carbonate ions.

6.3. Surface water and sediment

Compound (name and/or code)	Ecotoxicology
Potassium hydrogen carbonate	No data were available for long-term scale for aquatic organisms. Data gap pending on the information on the background level of K^+ .
	The acute risk for fish and daphnia was assessed as low.

6.4. Air

Compound (name and/or code)	Toxicology
Potassium hydrogen carbonate	LC_{50} inhalation > 4.88 mg/L (whole body, 41/2 hours)

EFSA Journal 2012;10(1):2524

¹⁶ OJ L 330, 5.12.1998, p.32



List of studies to be generated, still ongoing or available but not peer reviewed

This is a complete list of the data gaps identified during the peer review process, including those areas where a study may have been made available during the peer review process but not considered for procedural reasons (without prejudice to the provisions of Article 7 of Directive 91/414/EEC concerning information on potentially harmful effects).

- Wettability and foam persistence of the formulation (relevant for all representative uses evaluated; submission date proposed by the notifier: unknown; see section 1).
- The background level of K⁺ in natural soils and surface waters needs to be reported from a study or a peer reviewed scientific reference (relevant for all representative uses evaluated; submission date proposed by the notifier: unknown; see section 4).
- Appropriate long-term risk assessments for wild mammals (relevant for all representative uses evaluated; submission date proposed by the notifier: unknown; see section 5).
- The long-term risk assessments for aquatic organisms should be re-considered once the information on the background levels of K⁺ is available, see data gap identified in section 4 (relevant for all representative uses evaluated; submission date proposed by the notifier: unknown; see section 5).
- Algal toxicity data and related risk assessments for algae (relevant for all representative uses evaluated; submission date proposed by the notifier: the notifier has indicated that a new study is already available, but not peer-reviewed; see section 5).
- Appropriate risk assessments for the representative formulation and aquatic organisms. For the risk assessment toxicological studies might need to be conducted (relevant for all representative uses evaluated; submission date proposed by the notifier: unknown; see section 5).
- Higher tier risk assessments for honeybees (relevant for all representative uses evaluated; submission date proposed by the notifier: unknown; see section 5).
- Appropriate risk assessments for the representative formulation to honeybees (relevant for all representative uses evaluated; submission date proposed by the notifier: the notifier has indicated that a new study is already available to support the risk assessment, but the study is not peerreviewed; unknown; see section 5).
- Appropriate risk assessments for non-target arthropods to be provided once the information on the background levels of K⁺ is available. For the risk assessments toxicological studies might need to be conducted (relevant for all representative uses evaluated; submission date proposed by the notifier: the notifier has indicated that new studies are already available, but not peer-reviewed; see section 5).
- Appropriate risk assessments for soil non-target organisms to be provided once the information on the background levels of K⁺ is available, see data gap identified in section 4 (relevant for all representative uses evaluated; submission date proposed by the notifier: unknown; see section 5).
- Appropriate risk assessments for non-target terrestrial plants to be provided once the information on the background levels of K⁺ is available, see data gap identified in section 4(relevant for all representative uses evaluated; submission date proposed by the notifier: unknown; see section 5).

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Particular conditions proposed to be taken into account to manage the risk(s) identified None.

9. Concerns

Issues that could not be finalised

An issue is listed as an issue that could not be finalised where there is not enough information available to perform an assessment, even at the lowest tier level, for the representative uses in line with the Uniform Principles of Annex VI to Directive 91/414/EEC and where the issue is of such importance that it could, when finalised, become a concern (which would also be listed as a critical area of concern if it is of relevance to all representative uses).

- The environmental risk assessment for soil and water can not be finalised until the naturally occurring background levels assumed for potassium are confirmed by studies or peer reviewed scientific literature. Consequently, the long-term risk assessment for aquatic organisms, and the risk assessments for non-target arthropods, soil non-target organisms and terrestrial non-target plants could not be finalised.
- There were some indications that the representative formulation is more toxic to aquatic organisms than the active substance. Therefore further data and assessments are necessary to finalise the risk assessments.
- There were some indications that the representative formulation is more toxic to honeybees than the active substance. Therefore further data and assessments are necessary to finalise the risk assessments.

9.2. Critical areas of concern

An issue is listed as a critical area of concern where there is enough information available to perform an assessment for the representative uses in line with the Uniform Principles of Annex VI to Directive 91/414/EEC, and where this assessment does not permit to conclude that for at least one of the representative uses it may be expected that a plant protection product containing the active substance will not have any harmful effect on human or animal health or on groundwater or any unacceptable influence on the environment.

An issue is also listed as a critical area of concern where the assessment at a higher tier level could not be finalised due to a lack of information, and where the assessment performed at the lower tier level does not permit to conclude that for at least one of the representative uses it may be expected that a plant protection product containing the active substance will not have any harmful effect on human or animal health or on groundwater or any unacceptable influence on the environment.

Based on the available data, a high risk to honeybees was identified.



9.3. Overview of the concerns for each representative use considered

(If a particular condition proposed to be taken into account to manage an identified risk, as listed in section 8, has been evaluated as being effective, then 'risk identified' is not indicated in this table.)

Representative use	e	Fungicide on grapevine (8x 5100 g a.s./ha)	Fungicide on apple (8x 5100 g a.s./ha)
Operator risk	Risk identified		
Operator risk	Assessment not finalised		
Worker risk	Risk identified		
WOLKELLISK	Assessment not finalised		
Desetondon viels	Risk identified		
Bystander risk	Assessment not finalised		
Consumer risk	Risk identified		
Consumer risk	Assessment not finalised		
Risk to wild non	Risk identified		
target terrestrial vertebrates	Assessment not finalised		
Risk to wild non target terrestrial	Risk identified	X^4	X^4
organisms other than vertebrates	Assessment not finalised	$X^{1,3}$	$X^{1,3}$
Risk to aquatic	Risk identified		
organisms	Assessment not finalised	X ^{1,2}	$X^{1,2}$
Groundwater	Legal parametric value breached		
exposure active substance	Assessment not finalised		
Committee Ann	Legal parametric value breached		
Groundwater exposure metabolites	Parametric value of 10µg/L ^(a) breached		
metabolites	Assessment not finalised		
Comments/Remar	·ks		
The superscript numbe	rs in this table relate to t	he numbered points indicated in s	sections 9.1 and 9.2. Where there is

The superscript numbers in this table relate to the numbered points indicated in sections 9.1 and 9.2. Where there is no superscript number see sections 2 to 6 for further information. A column is greyed out if there is a concern for that specific use.

⁽a): Value for non-relevant metabolites prescribed in SANCO/221/2000-rev 10-final, European Commission, 2003



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APPENDICES

APPENDIX A - LIST OF END POINTS FOR THE ACTIVE SUBSTANCE AND THE REPRESENTATIVE FORMULATION

Identity, Physical and Chemical Properties, Details of Uses, Further Information

Active substance ((ISO Common Name) ±
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Function (e.g. fungicide)

Rapporteur Member State

Co-rapporteur Member State

Potassium bicarbonate Fungicide

Ireland

Not applicable.

Identity (Annex IIA, point 1)

Chemical name (IUPAC) ‡

Chemical name (CA) ‡

CIPAC No ‡

CAS No ‡

EC No (EINECS or ELINCS) ‡

FAO Specification (including year of publication) +

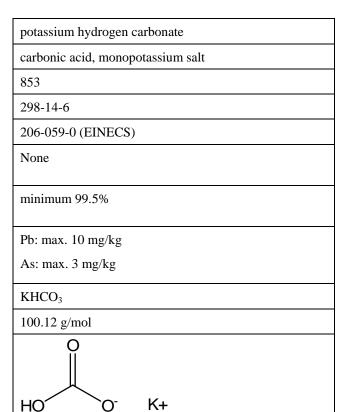
Minimum purity of the active substance as manufactured ‡

Identity of relevant impurities (of toxicological, ecotoxicological and/or environmental concern) in the active substance as manufactured

Molecular formula ‡

Molecular mass ‡

Structural formula ‡





Physical and chemical properties (Annex IIA, point 2)

Melting point (state purity) ‡

Boiling point (state purity) ‡

Temperature of decomposition (state purity)

Appearance (state purity) ‡

Vapour pressure (state temperature, state purity) ‡

Henry's law constant ‡

Solubility in water (state temperature, state purity and pH) ‡

Solubility in organic solvents ‡ (state temperature, state purity)

Surface tension ‡ (state concentration and temperature, state purity)

Partition co-efficient ‡ (state temperature, pH and purity)

Dissociation constant (state purity) ‡

UV/VIS absorption (max.) incl. $\epsilon \ddagger$ (state purity, pH)

Flammability ‡ (state purity)

Explosive properties ‡ (state purity)

Oxidising properties ‡ (state purity)

Decomposes without melting at 156 °C (>99.0%)

Decomposes at 156 °C (>99.0%)

> 156 °C (>99.0%)

Pure material: White crystalline solid (>99.5%)

Technical material: No information.

Not applicable.

No Henry's law constant.

332 g/L at 20°C

Almost insoluble in alcohol.

Not applicable.

No information provided. Not considered relevant.

Not applicable. Potassium hydrogen carbonate completely dissociates to its respective ions when dissolved in water:

$$KHCO_3 \rightarrow K^+ + HCO_3^-$$

HCO₃ is amphoteric and will then naturally participate in natural carbonic acid equilibria:

$$CO_3^{2-} + 2H^+ \iff HCO_3^- + H^+ \text{ (pKa}_1 = 10.377)$$

$$HCO_3^- + H^+ \iff H_2CO_3$$
 (pKa₂ = 6.381)

$$H_2CO_3 = CO_2 + H_2O$$

No UV/Vis, IR, MS or NMR spectra are available.

Not flammable.

Not explosive.

Not oxidizing.



Summary of representative uses evaluated (Potassium hydrogen carbonate)

Crop and/	Member State Product		Member G		G Pests or		Application			Application rate per treatment (for explanation see the text in front of this section)			РНІ		
or situation (a)	or Country	name	or I (b)	Group of pests controlled (c)	Type (d-f)	Conc. of as (i)	method kind (f-h)	growth stage & season (j)	number min/ max (k)	interval between applications (min)	kg as/hL min– max (l)	water L/ha min – max	kg as/ha min– max (l)	(days) (m)	Remarks
Vitis vinifera VITVI {Vine}	All EU	Armicarb 85SP	F	Uncinula necator {Vine powdery mildew}	SP	850 g/kg	Broadcas t using air blast orchard sprayer	BBCH 12 to 89	1 to 8	10 days	0.30 - 0.72	200- 600	2.125 to 5.100	1	Volumes and doses will vary according to crop canopy size.
Malus sylvestris MABSD {Apple}	All EU	Armicarb 85SP	F	Venturia inaequalis {Apple SCAB}	SP	850 g/kg	Broadcas t using air blast orchard sprayer	BBCH 10 to 85	1 to 8	10 days	0.34 – 0.51	500- 1000	2.125 to 5.100	1	Volumes and doses will vary according to crop canopy size.

- (a) For crops, the Codex and EU (or other) classifications should be used; where relevant, the use situation should be described (e.g. fumigation of a structure)
- (b) Outdoor or field use (F), glasshouse application (G) or indoor application (I)
- (c) e.g. biting or suckling insects, soil borne insects, foliar fungi, weeds
- (d) e.g. wettable powder (WP), emulsifiable concentrate (EC), granule (GR)
- (e) GCFP Codes GIFAP Technical monograph No 2, 1989
- (f) All abbreviations used must be explained
- (g) Method, e.g. high volume spraying, low volume spraying, spreading, dusting, drenching

- (h) Kind, e.g. overall, broadcast, aerial spraying, row, individual plant, between the plant type of equipment used must be indicated
- (i) g/kg or g/l
- (j) Growth stage at last treatment (BBCH Monograph, Growth Stage of Plants, 1997, Blackwell, ISBN 3-8263-3152-4), including where relevant, information on season at time of application
- (k) Indicate the minimum and maximum number of application possible under practical conditions of use
- (l) PHI minimum pre-harvest interval
- (m) Remarks may include: Extent of use/economic importance/restrictions

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Methods of Analysis

Analytical methods for the active substance (Annex IIA, point 4.1)

Technical as (analytical technique)

Acid-base titration

Impurities in technical as (analytical technique)

Acid-base titration

USP Limit tests 231 & 211

Plant protection product (analytical technique)

Acid-base titration

Analytical methods for residues (Annex IIA, point 4.2)

Residue definitions for monitoring purposes

Food of plant origin

A residue definition is not required and therefore has not been set.

Safety profile of Potassium hydrogen carbonate:

Potassium hydrogen carbonate is approved for food use in Europe and has been assigned a food additive number of E501 (raising agent). Potassium hydrogen carbonate is also listed for food use in the internationally recognized "Food Chemicals Codex"

Food of animal origin

Soil

See statement above

A residue definition is not required

Potassium hydrogen carbonate is a natural component of soil and therefore cannot be distinguished from existing potassium and bicarbonate ions in the soil. Potassium hydrogen carbonate is a natural component of the environment, including aquatic bodies such as streams, rivers, lakes and ponds. A discussion is provided in the environmental section to substantiate that manufactured potassium hydrogen carbonate should not act any differently to the potassium hydrogen carbonate already present in the environment.

Water surface

drinking/ground

Air

See statement above

See statement above

See statement above

Monitoring/Enforcement methods

Food/feed of plant origin (analytical technique and LOQ for methods for monitoring purposes)

Food/feed of animal origin (analytical technique and LOQ for methods for monitoring purposes)

Soil (analytical technique and LOQ)

A waiver for analytical methods for residues is agreed in view of physico-chemical, toxicological, ecotoxicological and environmental fate properties of active substance and formulated material.

See statement above

See statement above



Water (analytical technique and LOQ)	See statement above
Air (analytical technique and LOQ)	See statement above
Body fluids and tissues (analytical technique and LOQ)	Not required

Classification and proposed labelling with regard to physical and chemical data (Annex IIA, point 10)

Active substance

RMS/peer review proposal

Potassium hydrogen carbonate and the P.P.P. Armicarb 85SP will not classify from a physical/chemical viewpoint.



Chapter 2.3 Impact on Human and Animal Health

Absorption, distribution, excretion and metabolism (toxicokinetics) (Annex IIA, point 5.1)

Rate and extent of oral absorption ‡	Rapidly absorbed, approximately 100%				
Distribution ‡	Widespread				
Potential for accumulation ‡	Not relevant				
Rate and extent of excretion ‡	Not relevant				
Metabolism in animals ‡	Normal homeostasis maintained through well known mechanisms				
Toxicologically relevant compounds ‡ (animals and plants)	K+ ion				
Toxicologically relevant compounds ‡ (environment)	-				

Acute toxicity (Annex IIA, point 5.2)

Rat LD ₅₀ oral ‡	Females: 2064 mg/kg bw/day
Rat LD ₅₀ dermal ‡	>2000 mg/kg bw
Rat LC ₅₀ inhalation ‡	> 4.88 mg/L (whole body, 41/2 hours)
Skin irritation ‡	Non-irritant
Eye irritation ‡	Moderate, reversible
Skin sensitisation ‡	Non-sensitising (M & K)

Short term toxicity (Annex IIA, point 5.3)

Target / critical effect	Altered urinary pH/hypertrophy of adrenal	
	zona glomerulosa/increased potassium	
	exretion. Urinary bladder hyperplasia (rats)	
Lowest relevant oral NOAEL / NOEL	4- and13-week rat-LOAEL: 2%	
Lowest relevant dermal NOAEL / NOEL	No data - not required	
Lowest relevant inhalation NOAEL / NOEL	No data - not required	

Genotoxicity (Annex IIA, point 5.4)

Genotoxicity

Data available of limited validity. No further data required.

Long term toxicity and carcinogenicity (Annex IIA, point 5.5)

Target/critical effect	Growth retardation, ↑ serum potassiumin, ↑urinary potassium, ↑ urinary pH and volume, hypertrophy of
	adrenal zona glomerulosa;
Lowest relevant NOAEL / NOEL	Neoplasia at 4% of diet and pre-neoplastic alterations at 2%.
Carcinogenicity	Hyperplasia, papilloma and carcinoma of urinary bladder in rats through well-recognised mechanism, not considered relevant to humans
	mechanism, not considered relevant to numans



Reproductive toxicity (Annex 11A, point 5.6)		
Reproduction toxicity		
Reproduction target / critical effect	No data available. No data required.	
Relavant parental NOAEL		
Relavant reproductive NOAEL		
Relavant offspring NOAEL		
Developmental toxicity		
Developmental target / critical effect	Data available of limited validity. No further data required.	
Relavant maternal NOAEL	-	
Relavant developmental NOAEL	-	
Neurotoxicity / Delayed neurotoxicity (Annex II	A, point 5.7)	
Acute neurotoxicity ‡	No data available. No data required.	
Repeated neurotoxicity ‡	No data available. No data required.	
Delayed neurotoxicity ‡	No data available. No data required.	



Other toxicological studies	(Annex	IIA,	point	5.8)
Mechanism studies ‡				

No data available. No data required.

Medical data ‡ (Annex IIA, point 5.9)

Overdose: Confusion; irregular or slow heartbeat; numbness or tingling in hands, feet or lips; shortness of breath or difficult breathing; paralysis of arms and legs,

	blood pressure drop; convulsions, coma, cardiac arrest		
Summary (Annex IIA, point 5.10)	Value	Study	Safety factor
ADI ‡	No suitable data available. Not needed.		
AOEL ‡	No suitable data available. Not needed		
ARfD ‡	No suitable data available. Not needed		
Dermal absorption ‡ (Annex IIIA, point 7.	3)		

D

Formulation: Armicarb 85 SP

100% (in the absence of data)

Exposure scenarios (Annex IIIA, point 7.2)

Operator

German and UK POEM models (application rate 5100 g a.s./ha):

Exposure estimates to potassium hydrogen carbonate as a plant protection product are below the normal dietary intake of potassium (3.5 g/adult/day equivalent to 128 mg potassium hydrogen carbonate /kg bw/day) with and without PPE.

Workers

Exposure estimates to potassium hydrogen carbonate as a plant protection product are below the normal dietary intake of potassium (3.5 g/adult/day equivalent to 128 mg potassium hydrogen carbonate /kg bw/day) with and without PPE.

Bystanders

Exposure estimates to potassium hydrogen carbonate as a plant protection product are below the normal dietary intake of potassium (3.5 g/adult/day equivalent to 128 mg potassium hydrogen carbonate /kg bw/day)



C	lassification a	hna	proposed labelling	with i	regard to	toxicological	data	(Annex IIA	point 10)
v	iassincauvii <i>a</i>	anu	DI UDUSCU IADCIIIII		tegaru w	tuaituiugitai	uata	TAILLY HA.	DOING TO I

	peer review proposal
Potassium hydrogen carbonate	None



Residues:

Metabolism in plants (Annex IIA, point 6.1 and 6.7, Annex IIIA, point 8.1 and 8.6)

Plant groups covered	Not required. The residue arising from the use of the plant protection product is indistinguishable from naturally occurring residues present in any treated crop.
Rotational crops	Not provided and not required
Metabolism in rotational crops similar to metabolism in primary crops?	Not applicable
Processed commodities	Not provided and not required
Residue pattern in processed commodities similar to residue pattern in raw commodities?	Not applicable
Plant residue definition for monitoring	Not proposed and not required
Plant residue definition for risk assessment	Not proposed and not required
Conversion factor (monitoring to risk assessment)	Not applicable

Metabolism in livestock (Annex IIA, point 6.2 and 6.7, Annex IIIA, point 8.1 and 8.6)

Animals covered	Not provided and not required
Animal residue definition for monitoring	Not applicable
Animal residue definition for risk assessment	Not applicable
Conversion factor (monitoring to risk assessment)	Not applicable
Metabolism in rat and ruminant similar (yes/no)	Not applicable
Fat soluble residue: (yes/no)	Not applicable

Residues in succeeding crops (Annex IIA, point 6.6, Annex IIIA, point 8.5)

Not applicable. The residue arising from the use of the plant protection product is indistinguishable from naturally occurring residues present in any treated crop.

Stability of residues (Annex IIA, point 6 introduction, Annex IIIA, point 8 Introduction)

Not provided and not required



Residues from livestock feeding studies (Annex IIA, point 6.4, Annex IIIA, point 8.3)

Expected intakes by livestock ≥ 0.1 mg/kg diet (dry weight basis) (yes/no - If yes, specify the level)

Potential for accumulation (yes/no):

Metabolism studies indicate potential level of residues ≥ 0.01 mg/kg in edible tissues (yes/no)

Muscle Liver Kidney Fat Milk Eggs

Ruminant:	Poultry:	Pig:			
Conditions of requ	irement of feeding s	tudies			
Not relevant.	Not relevant.	Not relevant.			
Not relevant.	Not relevant.	Not relevant.			
Not relevant.	Not relevant.	Not relevant.			
poultry studies con	Feeding studies (Specify the feeding rate in cattle and poultry studies considered as relevant) Residue levels in matrices: Mean (max) mg/kg				
Not relevant.	Not relevant.	Not relevant.			
Not relevant.	Not relevant.	Not relevant.			
Not relevant.	Not relevant.	Not relevant.			
Not relevant.	Not relevant.	Not relevant.			
Not relevant.					
	Not relevant.				

Summary of residues data according to the representative uses on raw agricultural commodities and feedingstuffs (Annex IIA, point 6.3, Annex IIIA, point 8.2)

Crop	Northern Southern Region, field or glasshouse,	Trials results relevant to the representative uses (a)	Recommendation/com ments	MRL estimated from trials according to representativ e use	HR (c)	STMR (b)
Residue	trials for pota	ssium hydrogen carbonat	e not provided and not re	quired.		_

⁽a) Numbers of trials in which particular residue levels were reported e.g.~3x < 0.01,~0.01,~6x~0.02,~0.04,~0.08,~2x~0.15,~0.17

⁽b) Supervised Trials Median Residue *i.e.* the median residue level estimated on the basis of supervised trials relating to the representative use

⁽c) Highest residue



Consumer risk assessment (Annex IIA, point 6.9, Annex IIIA, point 8.8)⁷

ADI	Not proposed and not required.
TMDI (% ADI) according to WHO European diet	Not required
TMDI (% ADI) according to national (to be specified) diets	Not required.
IEDI (WHO European Diet) (% ADI)	Not required.
NEDI (specify diet) (% ADI)	Not required.
Factors included in IEDI and NEDI	Not required.
ARfD	Not proposed and not required.
IESTI (% ARfD)	Not required.
NESTI (% ARfD) according to national (to be specified) large portion consumption data	Not required.
Factors included in IESTI and NESTI	Not required.

⁷ To be done on the basis of WHO guidelines and recommendations with the deviations within the EU so far accepted (especially diets).

Processing factors (Annex IIA, point 6.5, Annex IIIA, point 8.4)

Crop/ process/ processed product	Numberof studies	Processing factors		Amount	
		Transfer factor ⁸	Yield factor ⁸	transferred (%) (Optional)	
Not provided and not required.					

Proposed MRLs (Annex IIA, point 6.7, Annex IIIA, point 8.6)

No MRL proposed

Not required. The residue arising from the use of the plant protection product is indistinguishable from naturally occurring residues present in any treated crop

When the MRL is proposed at the LOQ, this should be annotated by an asterisk after the figure.



Chapter 2.5: Fate and Behaviour in the Environment

Route of degradation (aerobic) in soil

(Annex IIA, point 7.1.1.1.1)

Potassium hydrogen carbonate completely dissociates to potassium and bicarbonate ions in the presence of water.

Route of degradation in soil - Supplemental studies

(Annex IIA, point 7.1.1.1.2)

None available.

Rate of degradation in soil

(Annex IIA, point 7.1.1.2, Annex IIIA, point 9.1.1)

Not applicable: Potassium hydrogen carbonate completely dissociates to potassium and bicarbonate ions in the presence of water.

Soil adsorption/desorption

(Annex IIA, point 7.1.2) It should be noted that no data was presented for this section nor was any data requested.

Potassium is strongly bound in soil and a rapid equilibrium is observed between soluble and exchangeable forms.

Mobility in soil

Application rate

(Annex IIA, point 7.1.3, Annex IIIA, point 9.1.2) It should be noted that no data was presented for this section nor was any data requested.

Potassium has a very low mobility in soil due to cation binding to negatively charged soil components.

PEC (soil) (Annex IIIA, point 9.1.3)
Method of calculation

Calculations were based on a lumped application of 40.8 kg a.s./ha corresponding to the maximum number of recommended doses and the highest rate of application in a season.

Potassium hydrogen carbonate spontaneously dissociates to potassium and bicarbonate ions in moist soils. Consequently initial PECs were calculated for the potassium and bicarbonate ions.

% plant interception: 50

Crops: Apples & vines.

Number of applications: 1 lumped application of the active substance. That is, the active substance is applied eight times per season with no loss of residues.

Application rate: 40.8 kg a.s./ha per season [8 x 5.10 kg a.s/ha]

PEC_S	Single application	Single	Multiple	Multiple application
	Actual	application	application actual	actual
mg /kg soil		Time weighted		



Initial

	average	K^{+}	HCO ₃	KHCO ₃
Only the PEC soil to assessment is included points[EPCO Manual September 2005].	in the list of end		16.6	27.2

Route and rate of degradation in water

(Annex IIA, point 7.2.1)

Not applicable: Potassium hydrogen carbonate completely dissociates to potassium and bicarbonate ions in the presence of water. Bicarbonate is produced from various natural sources, particularly carbonate based rocks and respiration of aquatic plants during the hours of darkness. Typical levels found in natural surface waters adjacent to agricultural land are between 100-500 mg/L.

Potassium is an essential nutrient for aquatic plants and micro-organisms and has a well known cycle via the food chain.

PEC (surface water)

(Annex IIIA, point 9.2.3) Method of calculation

Application rate

Main route of entry

Given the nature of the active substance it was not considered appropriate to use the FOCUS model to determine the PEC of potassium hydrogen carbonate in surface waters. Instead, an estimation based on spray drift with no degradation between applications was used.

40.8 kg a.s ha[8 x 5.10 kg a.s./ha]

Spray drift [100 %]

PEC _{SW}	Single app Actu		Sing applic	_	Multiple application actual		ltiple apple weighted	
mg as/L	K ⁺	HCO ₃	Time we	Ŭ		K ⁺	HCO ₃	KHCO ₃
Initial (100% spray drift)	-	-	-		-	5.30	8.30	13.6
Apple crop [3 m buffer zone]	1	-	-		-	1.55	2.42	3.97
Vine crop [3 m buffer zone]	-	-	-		-	0.42	0.66	1.088



PEC (groundwater)

Method of calculation and type of study (e.g. modelling, monitoring, lysimeter) Application rate PEC_{GW}
Maximum concentration
Average annual concentration

Not calculated. The dissolution products of Potassium hydrogen carbonate are naturally occurring in the environment. For example, bicarbonate is produced from various natural sources, particularly carbonate-based rocks:

 $CaCO_{3(S)} \leftrightarrows Ca^{2+}_{(aq)} + CO_3^{2-}_{(aq)}$ $CO_3^{2-}_{(aq)} + H_2O_{(1)} \leftrightarrows HCO_3^{-}_{(aq)} + OH^{-}_{(aq)}$ and respiration of aquatic plants during the hours of darkness.

Potassium is an essential nutrient for aquatic plants and micro-organisms and has a well known cycle via the food chain.

Fate and behaviour in air

(Annex IIA, point 7.2.2, Annex III, point 9.3) It should be noted that no data was presented for this section nor was any data requested.

Direct photolysis in air

Quantum yield of direct phototransformation at > 290 nm

Photochemical oxidative degradation in air Volatilisation

PEC (air)

Method of calculation

Not applicable. Potassium hydrogen carbonate is not volatile and does not degrade in air.

Not applicable

Not applicable

Not applicable

Not applicable

PEC_A

Maximum concentration

Not applicable

Definition of the Residue

(Annex IIA, point 7.3) Relevant to the environment

Not applicable, Potassium hydrogen carbonate is naturally present in the environment.

Monitoring data, if available

(Annex IIA, point 7.4) Soil

(indicate location and type of study)

Surface water

(indicate location and type of study)

Ground water

(indicate location and type of study)

Air

(indicate location and type of study)

Not applicable, potassium and bicarbonate ions are naturally present in the environment.

Potassium and bicarbonate ions are naturally present in sediments in surface water.

Potassium and bicarbonate are naturally present in groundwater.

Not applicable: Potassium hydrogen carbonate is not volatile

List of studies submitted.

Waivers were requested by the notifier for potassium hydrogen carbonate for all environmental fate studies. This was accepted by the RMS as potassium hydrogen carbonate is a natural component of the environment. Inputs from the use of 'Armicarb 85SP' are expected to be negligible compared with natural background levels. A data gap has been identified for studies or peer reviewed scientific literature to support the naturally occurring levels of potassium proposed by the notifier.



Chapter 2.6 Effects on Non-target Species

Effects on terrestrial vertebrates (Annex IIA, point 8.1, Annex IIIA, points 10.1 and 10.3)

Species	Test substance	Time scale	End point (mg/kg bw/day)	End point (mg/kg feed)
Birds ‡				
Standard tests not available.	Not required.			
Chicken	NaHCO ₃ *	Acute	>8,075*	10 000
Chicken	NaHCO ₃ *	Short-term	>8,075*	10 000
Chicken	NaHCO ₃ *	Long-term	>8,075*	10 000
Mammals				
Rat	KHCO ₃	Acute	2064	-
Long-term and additional his	gher tier studies			
Not available				

^{*:} endpoint is based on summaries of long-term feeding studies where chickens were fed with high concentration of sodium bicarbonate in the standard feed and no adverse effects were observed. 1.04 kg feed /kg bw/day was assumed to be consumed by the chickens. The value is expressed as HCO₃, therefore refers only to bicarbonate.

Toxicity/exposure ratios for terrestrial vertebrates (Annex IIIA, points 10.1 and 10.3)

Crop and application rate: Apples & grapevines, 8 x 5100 g a.s./ha

Indicator species	Time scale	ETE	TER	Annex VI Trigger
Tier 1 (Birds)				
representative insect eating bird	Acute	276	>29	10
representative insect eating bird	Short-term	154	>52	10
representative insect eating bird	Long-term	154	>52	5
Higher tier refinement (Birds)				
Not required				
Tier 1 (Mammals)				
Rat	Acute	964	2.1	10
Long-term and higher tier refine	ement (Mammals)			
Not available – data gap	·			·

Toxicity data for aquatic species (most sensitive species of each group) (Annex IIA, point 8.2, Annex IIIA, point 10.2)

Group	Test substance	Time-scale	End point	Toxicity ¹
		(Test type)		(mg/L)
Laboratory tests ‡				
Fish				
Rainbow Trout	KHCO ₃	96 hr (flow-through)	Mortality,LC ₅₀	>1200 (nom)
Bluegill Sunfish	KHCO ₃	96 hr (flow-through)	Mortality,LC ₅₀	>1200 (nom)
Aquatic invertebrate				
Daphnia magna	KHCO ₃	48 hr (flow-through)	Mortality, EC ₅₀	>860 (_{nom})
Sediment dwelling organisms				
Not available. Not required				
Algae				
Not available - data gap				



Group	Test substance	Time-scale	End point	Toxicity ¹
		(Test type)		(mg/L)
Higher plant				
Not available. Not required				
Microcosm or mesocosm tests	}			
Not available. Not required				

 $^{^{1}}$ indicate whether based on nominal ($_{nom}$) or mean measured concentrations ($_{mm}$). In the case of preparations indicate whether end points are presented as units of preparation or a.s.

Toxicity/exposure ratios for the most sensitive aquatic organisms (Annex IIIA, point 10.2)

Worst-case PEC_{SW} of KHCO₃ following a lumped dose of 8 applications at the maximum recommended dose rate of Arimcarb 85SP: PEC_{SW} for overspray is 13.6 mg/L; Apples (3m from edge of field) is 3.97 mg/L; Vines (3m from edge of field) is 1.088 mg/L.

Organism	Toxicity end point (mg/L)	TER _A overspray	TER _A 3m from apples	TER _A 3m from vines	Annex VI Trigger
Fish	LC ₅₀ 1200	> 88	> 302	> 1103	100
Aquatic invertebrates	EC ₅₀ 860	> 63	> 217	> 790	100

Bioconcentration
No data. Not required

Effects on honeybees (Annex IIA, point 8.3.1, Annex IIIA, point 10.4)

Test substance	Acute oral toxicity	Acute contact toxicity
	(LD ₅₀ µg a.s./bee)	(LD ₅₀ μg a.s./bee)
KHCO ₃	No data. Not required.	>24
Field or semi-field tests		
No data. Not required.		

Hazard quotients for honey bees (Annex IIIA, point 10.4)

Test	Exposure	Dose g a.s./ha	LD ₅₀ μg/bee	Hazard quotient	Annex VI
substance	Route			Q_{HC}	Trigger
KHCO ₃	Contact	5100	>24	<212	50

Effects on other arthropod species (Annex IIA, point 8.3.2, Annex IIIA, point 10.5)

No data - data gap

Effects on earthworms, other soil macro-organisms and soil micro-organisms (Annex IIA points 8.4 and 8.5. Annex IIIA, points, 10.6 and 10.7)

Earthworms
No data - data gap
Other soil macro-organisms
No data - data gap



Soil micro-organisms		
No data - data gap		

Effects on non target plants (Annex IIA, point 8.6, Annex IIIA, point 10.8)

Preliminary screening data

No data - data gap

Effects on biological methods for sewage treatment (Annex IIA 8.7)

No data. Not required

Ecotoxicologically relevant compounds (consider parent and all relevant metabolites requiring further assessment from the fate section)

Potassium hydrogen carbonate

Classification and proposed labelling with regard to ecotoxicological data (Annex IIA, point 10 and Annex IIIA, point 12.3)

RMS proposal

Active substance No classification proposed



ABBREVIATIONS

1/n slope of Freundlich isotherm

 λ wavelength

ε decadic molar extinction coefficient

°C degree Celsius (centigrade)

μg microgram

μm micrometer (micron)
 a.s. active substance
 AChE acetylcholinesterase
 ADE actual dermal exposure
 ADI acceptable daily intake
 AF assessment factor

AOEL acceptable operator exposure level

AP alkaline phosphatase
AR applied radioactivity
ARfD acute reference dose

AST aspartate aminotransferase (SGOT)

AV avoidance factor
BCF bioconcentration factor
BUN blood urea nitrogen

bw body weight

CAS Chemical Abstracts Service

CFU colony forming units
ChE cholinesterase
CI confidence interval

CIPAC Collaborative International Pesticides Analytical Council Limited

CL confidence limits

cm centimetre

d day

DAA days after application
DAR draft assessment report
DAT days after treatment

DM dry matter

 DT_{50} period required for 50 percent disappearance (define method of estimation) DT_{90} period required for 90 percent disappearance (define method of estimation)

dw dry weight

EbC₅₀ effective concentration (biomass)

EC₅₀ effective concentration ECHA European Chemical Agency EEC European Economic Community

EINECS European Inventory of Existing Commercial Chemical Substances

ELINCS European List of New Chemical Substances

EMDI estimated maximum daily intake ER_{50} emergence rate/effective rate, median ErC_{50} effective concentration (growth rate)

EU European Union

EUROPOEM European Predictive Operator Exposure Model

f(twa) time weighted average factor

FAO Food and Agriculture Organisation of the United Nations

FIR Food intake rate

FOB functional observation battery

FOCUS Forum for the Co-ordination of Pesticide Fate Models and their Use

g gram

GAP good agricultural practice GC gas chromatography

GCPF Global Crop Protection Federation (formerly known as GIFAP)

GGT gamma glutamyl transferase

GM geometric mean growth stage GS glutathion **GSH** hour(s) h ha hectare Hb haemoglobin haematocrit Hct hectolitre hL.

HPLC high pressure liquid chromatography

or high performance liquid chromatography

HPLC-MS high pressure liquid chromatography – mass spectrometry

HQ hazard quotient

IEDIinternational estimated daily intakeIESTIinternational estimated short-term intakeISOInternational Organisation for StandardisationIUPACInternational Union of Pure and Applied Chemistry

JMPR Joint Meeting on the FAO Panel of Experts on Pesticide Residues in Food and

the Environment and the WHO Expert Group on Pesticide Residues (Joint

Meeting on Pesticide Residues)

K_{doc} organic carbon linear adsorption coefficient

kg kilogram

K_{Foc} Freundlich organic carbon adsorption coefficient

L litre

LC liquid chromatography LC_{50} lethal concentration, median

LC-MS liquid chromatography-mass spectrometry

LC-MS-MS liquid chromatography with tandem mass spectrometry

LOAEL lethal dose, median; dosis letalis media LOAEL lowest observable adverse effect level

LOD limit of detection

LOQ limit of quantification (determination)

m metre

M/L mixing and loading
MAF multiple application factor
MCH mean corpuscular haemoglobin

MCHC mean corpuscular haemoglobin concentration

MCV mean corpuscular volume

mg milligram
mL millilitre
mm millimetre
mN milli-newton

MRL maximum residue limit or level

MS mass spectrometry

MSDS material safety data sheet MTD maximum tolerated dose

MWHC maximum water holding capacity
NESTI national estimated short-term intake

ng nanogram

NOAEC no observed adverse effect concentration

NOAEL no observed adverse effect level



NOEC no observed effect concentration

NOEL no observed effect level OM organic matter content

Pa pascal

PD proportion of different food types
PEC predicted environmental concentration
PECair predicted environmental concentration in air

PEC_{gw} predicted environmental concentration in ground water PEC_{sed} predicted environmental concentration in sediment PEC_{soil} predicted environmental concentration in soil

PEC_{sw} predicted environmental concentration in surface water

pH pH-value

PHED pesticide handler's exposure data

PHI pre-harvest interval

PIE potential inhalation exposure

pK_a negative logarithm (to the base 10) of the dissociation constant

 P_{ow} partition coefficient between n-octanol and water

PPE personal protective equipment

ppm parts per million (10⁻⁶) ppp plant protection product

PT proportion of diet obtained in the treated area

PTT partial thromboplastin time

QSAR quantitative structure-activity relationship

r² coefficient of determination RPE respiratory protective equipment

RUD residue per unit dose
SC suspension concentrate
SD standard deviation
SFO single first-order

SSD species sensitivity distribution STMR supervised trials median residue $t_{1/2}$ half-life (define method of estimation)

TER toxicity exposure ratio

TER_A toxicity exposure ratio for acute exposure

TER_{LT} toxicity exposure ratio following chronic exposure TER_{ST} toxicity exposure ratio following repeated exposure

TK technical concentrate TLV threshold limit value

TMDI theoretical maximum daily intake

TRR total radioactive residue

TSH thyroid stimulating hormone (thyrotropin)

TWA time weighted average UDS unscheduled DNA synthesis

UV ultraviolet
W/S water/sediment
w/v weight per volume
w/w weight per weight
WBC white blood cell

WG water dispersible granule WHO World Health Organisation

wk week yr year