

(3) If the ship leaves the harbour immediately upon commencement of fumigation activities, a fumigation supervisor shall be on board until such time as

1. the fumigated load has been unloaded, or
2. the requirements specified in paragraph 2 have been fulfilled.

(4) The provisions of No. 5.4 (2) shall apply as well.

(5) The gas-tightness of all fumigated areas shall be verified at least every eight hours while the ship is in transit. The results of this verification shall be recorded in the ship's log.

(6) A minimum of 24 hours prior to the arrival of any fumigated ship, the competent harbour officials shall be notified of the time of fumigation and the areas and transport containers that are, have been or are to be under fumigation.

#### 5.8 Additional regulations pertaining to specific fumigants

##### 5.8.1 Bromomethane

(1) If any area that is to be fumigated with bromomethane is to be entered for purposes of opening bottle valves, holders of the relevant competence certificate shall be deployed in sufficient numbers to ensure that such area can be evacuated no more than ten minutes after the valve on the first bottle is opened.

(2) Any enclosed area whose bromomethane concentration exceeds 2 grams per cubic meter shall be off limits. Any enclosed area whose bromomethane concentration exceeds 0.4 grams per cubic meter shall be occupied for a maximum of ten minutes by persons equipped with respiratory protection.

(3) Fumigation in greenhouses or out of doors shall be realized solely under gas-tight covers or tarps. Warning signs pursuant to No. 5.6 (2) shall be posted at any location at which such fumigation is realized.

##### 5.8.2 Hydrogen cyanide

(1) A maximum of 30 grams per cubic meter (2.7 volume percent in air) of hydrogen cyanide shall be used for the fumigation of enclosed spaces. The use of additional fumigant shall be admissible after two hours have elapsed.

(2) The holder of a competence certificate shall not use more than 100 kilograms of fumigant over the course of a single work day.

##### 5.8.3 Hydrogen phosphide

(1) Application of Nos. 5.3.2 and 5.3.3 shall be excluded for hydrogen phosphide use out of doors.

(2) Preparations that produce hydrogen phosphide shall contain an additive that prevents the spontaneous combustion of hydrogen phosphide.

(3) Hydrogen phosphide fumigation shall be realized using a fumigant mixture that does not have the capacity to produce an explosive mixture of gas and air.

##### 5.8.4 Formaldehyde

The fumigation supervisor shall deem a fumigated area and the objects therein safe for use only insofar as an appropriate proof procedure has verified that the concentration of formaldehyde in the area does not exceed 0.1 milliliter per cubic meter.

**Appendix to Section 8.1.2****Extract of the German regulation TRGS 512 “Fumigations” (edition January 2007)  
concerning the storage of dangerous goods**

*unauthorized translation!*

**6 Keeping and storing of fumigants**

- (1) Very toxic and toxic fumigants must be kept or stored in such way, that they do not endanger human health and the environment. Suitable and reasonable precautions must be taken to prevent, as far as possible, wrongful or mistaken use. They must be kept under lock and key or stored in such a way that only experts according to number 4.3 have access to them (see also TRGS 514).
- (2) Hazardous substances must not be transferred to or stored in containers which as a result of their shapes or labelling may confuse the content with foods. They may only be kept away from pharmaceuticals, foods or feedstuffs.

**Extract of the German regulation TRGS 514**

*unauthorized translation!*

The Technical Rule for Hazardous Substances (TRGS) 514 (edition September 2008) contains special safety measurements for the storage of very toxic and toxic substances in packages and portable containers.

Contents:

1. Scope
2. Definition of terms and explanations
3. Safety measures
4. Operating regulations
5. Personal protective equipment
6. Protective hygiene measures
7. Rescue equipment and First aid

With regard to the scope of the rules on handling contained in the Hazardous Substances Ordinance (GefStoffV) as well as the generally applicable definitions of terms, attention is drawn to §§ 2 and 3 of the Hazardous Substances Ordinance.

Link to public version of TRGS 514:

[http://www.baua.de/de/Themen-von-A-Z/Gefahrstoffe/TRGS/TRGS-514.html?\\_\\_nnn=true&\\_\\_nnn=true](http://www.baua.de/de/Themen-von-A-Z/Gefahrstoffe/TRGS/TRGS-514.html?__nnn=true&__nnn=true)

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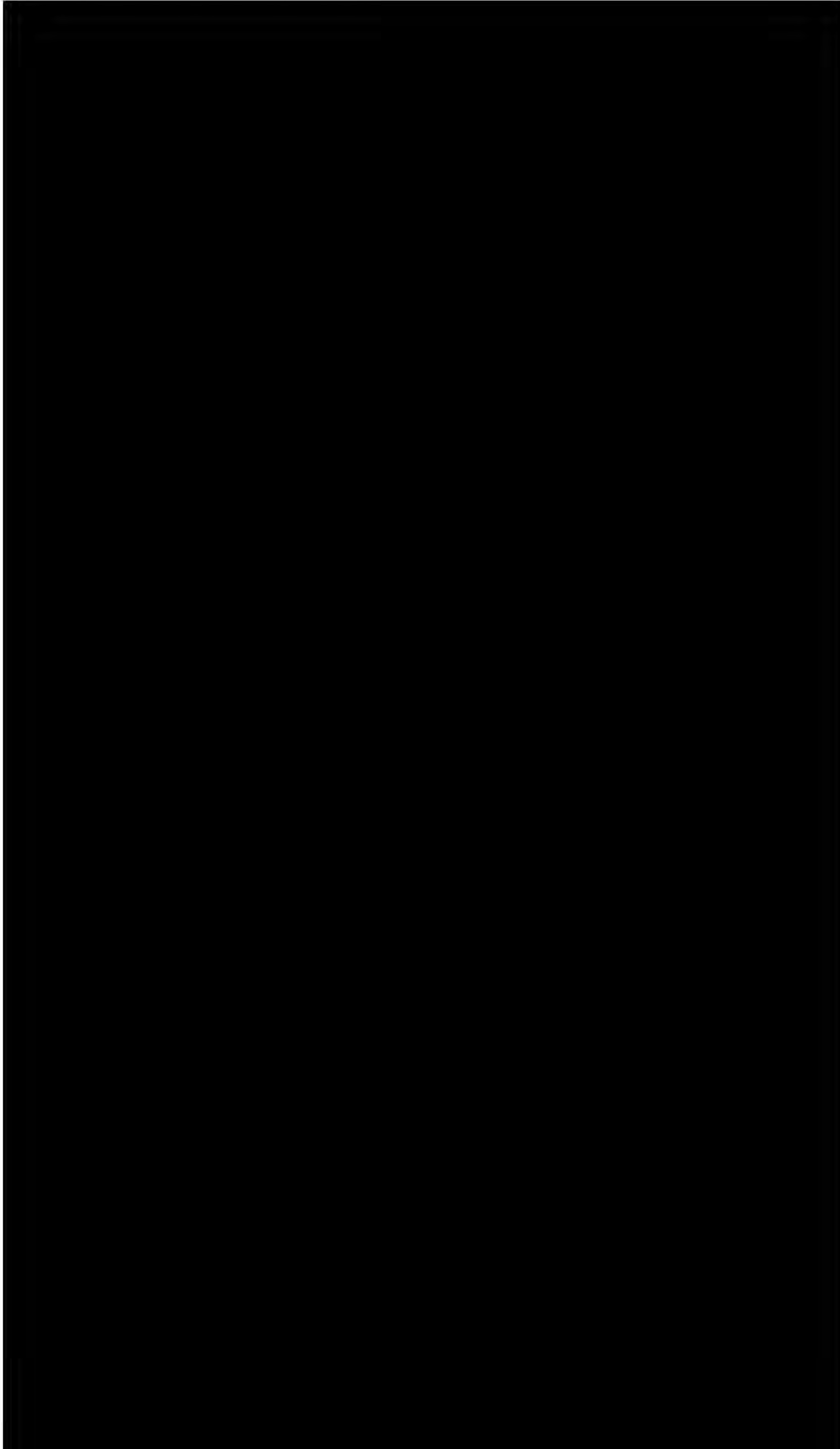
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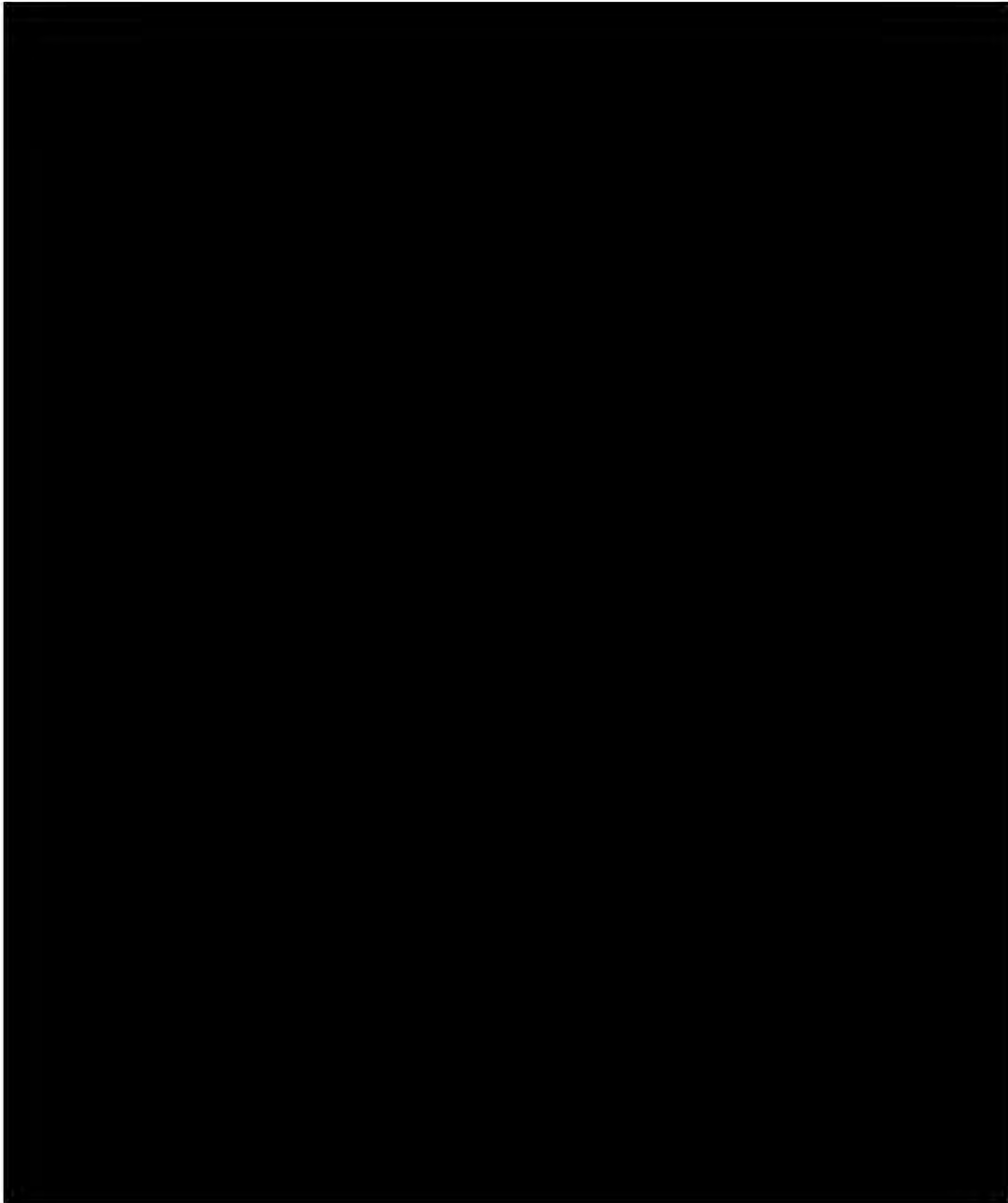
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**APPENDIX 3.2 TO CHAPTER 8:**

**Realistic worst case scenario**



**Worst case scenario**





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## Section A9

## Classification and labelling, packaging

## Annex Point IIB IX

Official  
use only**1 PACKAGING**

Not relevant.

**2 LABELLING****2.1 CLASSIFICATION AND LABELLING OF  
ALP PREMIX AND PHOSPHINE**

Please see included documents Figure 2\_1\_01, 2\_1\_02 and 2\_1\_03.

**2.2 UP-TO-DATE LABEL ACCORDING TO DIRECTIVE  
1999/45/EC**

Not relevant

**2.3 ADDITIONAL LABELLING REQUIREMENTS  
ACCORDING TO BPD ARTICLE 20 (3)**

Not relevant.

<b>Evaluation by Competent Authorities</b>	
	Use separate "evaluation boxes" to provide transparency as to the comments and views submitted
	<b>EVALUATION BY RAPPORTEUR MEMBER STATE</b>
<b>Date</b>	<i>Give date of action</i>
<b>Materials and Methods</b>	<i>State if the applicants version is acceptable or indicate relevant discrepancies referring to the (sub) heading numbers and to applicant's summary and conclusion.</i>
<b>Results and discussion</b>	<i>Adopt applicant's version or include revised version. If necessary, discuss relevant deviations from applicant's view referring to the (sub)heading numbers</i>
<b>Conclusion</b>	<b>Other conclusions:</b> <i>(Adopt applicant's version or include revised version)</i>
<b>Reliability</b>	<i>Based on the assessment of materials and methods include appropriate reliability indicator</i>
<b>Acceptability</b>	acceptable / not acceptable <i>(give reasons if necessary, e.g. if a study is considered acceptable despite a poor reliability indicator. Discuss the relevance of deficiencies and indicate if repeat is necessary.)</i>
<b>Remarks</b>	
	<b>COMMENTS FROM ...</b>
<b>Date</b>	<i>Give date of comments submitted</i>
<b>Materials and Methods</b>	<i>Discuss additional relevant discrepancies referring to the (sub)heading numbers and to applicant's summary and conclusion. Discuss if deviating from view of rapporteur member state</i>

**Section A9****Classification and labelling, packaging****Annex Point IIB IX**

<b>Results and discussion</b>	<i>Discuss if deviating from view of rapporteur member state</i>
<b>Conclusion</b>	<i>Discuss if deviating from view of rapporteur member state</i>
<b>Reliability</b>	<i>Discuss if deviating from view of rapporteur member state</i>
<b>Acceptability</b>	<i>Discuss if deviating from view of rapporteur member state</i>
<b>Remarks</b>	

**Figure 2\_1\_01:** Classification and Labelling: Aluminium phosphide (according to Directive 67/548/EEC and 1999/45/EC)

<p><b>Justified proposals for the classification and labelling of:</b></p> <p style="text-align: center;"><b>AIP-Premix</b></p> <p>according to Directive 67/548/EEC and 1999/45/EEC</p> <p><b>The product contains the following dangerous substances in the concentrations stated:</b></p> <p><b>85 % aluminiumphosphide:</b></p> <p><b>F, T+, N, R 15/29-21-26/28-32-50</b>          CAS No.: 20859-73-8          Acute dermal LD<sub>50</sub> (rat): 900 mg/kg<sup>(1)</sup>          Acute oral LD<sub>50</sub> (rat) = 8,7 mg/kg<sup>(2,3)</sup>          LC<sub>50</sub> inhalative (rat, 4hrs.): hydrogen phosphide 11 ppm = 0.015 mg/l<sup>(1)</sup>          Aqueous toxicity:          LC<sub>50</sub> (rainbow trout, 96 hrs.) = 9,7 x 10<sup>3</sup> ppm<sup>(4)</sup>          EC<sub>50</sub> (daphnia magna, 24 hrs.) = 0,2 mg/l<sup>(5)</sup></p>	<p>N, R 50: Classification of the preparation as per Annex III part B, table 2 of Directive 1999/45/EEC based on the classification of Aluminium phosphide as set down in Directive 67/548/EEC: results in N, R 50, as the concentration of Aluminium phosphide is <math>\geq 25</math> %.</p> <p><b>Labelling of the preparation:</b></p> <p>Danger symbols: F, T+, N          Indications of danger: flammable, very toxic, dangerous for the environment          Risk phrases: R 15/29-21-26/28-32-50          Safety phrases: S 1/2-3/9/14-30-36/37-45-61</p>
<p><b>The preparation is classified as follows:</b></p> <ul style="list-style-type: none"> <li>- as per Annex VI, No. 2.2.4 as well as No. 3.2.8 of Directive 67/548/EEC: <b>F, R 15/29</b></li> <li>- as per Annex VI, No. 3.2.3 of Directive 67/548/EEC: <b>R 21</b></li> <li>- as per article 3 (1) as well as Annex II of Directive 1999/45/EEC and Annex VI, No. 3.2.1 of Directive 67/548/EEC: <b>T+, R 26/28</b></li> <li>- as per Annex VI, No. 3.2.8 of Directive 67/548/EEC: <b>R 32</b></li> <li>- as per Annex III, part B, table 2 of Directive 1999/45/EEC: <b>N, R 50</b></li> </ul>	<p>1) WHO Environmental Health Criteria 73: Phosphine and Selected Metal Phosphides: a) page 18, b) page 72, c) page 75          2) International Bio-Research Inc., D-Hannover: Acute oral toxicity of Aluminium phosphide in rats (1.1.1977)          3) Weitz, R.S. &amp; Brown, R.M. (1975): Acute and subacute inhalation toxicities of phosphine, phenylphosphine and triphenylphosphine. Am. Ind. Assoc. J., 36: 452-458.          4) Laboratory for Pharmacology and Toxicology, D-Hamburg: Prüfung der akuten Toxizität von Aluminiumphosphid an Regenbogenforellen (24.11.1984)          5) Okolima, D-Burgwedel: Daphnientoxizität mit Aluminiumphosphid, 1986          MSDS of Ammoniumcarbamid</p>
<p><b>Explanation:</b></p> <p>T+, R 28: Classification of the preparation as per Annex I No. 1.1 of Directive 1999/45/EEC based on the classification of Aluminium phosphide as set down in Directive 67/548/EEC: results in T+, R 28, as the concentration of Aluminium phosphide is <math>\geq 7</math> %.</p>	

**Figure 2\_1\_02:** Classification and Labelling: Aluminium phosphide  
(according to Regulation (EC) No. 1272/2008)

Signal Word:  
Danger

Hazard statement:  
H260, H300, H311, H400, EUH029, EUH032

Precautinary statements:  
P233, P234, P273, P301+P310, P321, P335, P370+P378, P402+P404, P405, P501

Laudenbach, 26.06.2011

Justified proposals for the classification and labelling of:

ALP-Premix

according to Regulation (EC) No 1272/2008

The product contains the following dangerous substances in the concentrations stated:

85 % aluminiumphosphide:

GHS02, GHS06, GHS09  
H260, H300, H311, H400, EUH029, EUH032  
CAS No.: 20859-73-8  
Acute dermal LD<sub>50</sub> (rat): 900 mg/kg<sup>1)</sup>  
Acute oral LD<sub>50</sub> (rat) = 8,7 mg/kg<sup>1,2)</sup>  
LC<sub>50</sub> inhalative (rat, 4hrs.):  
hydrogen phosphide 11 ppm = 0,015 mg/l<sup>3)</sup>  
Aqueous toxicity:  
LC<sub>50</sub> (rainbow trout, 96 hrs.) = 0,7 x 10<sup>-3</sup> ppm<sup>4)</sup>  
EC<sub>50</sub> (daphnia magna, 24 hrs.) = 0,2 mg/l<sup>5)</sup>

The preparation is classified as follows:

<b>Current entry in Annex VI, CLP Regulation</b>	Water-react. 1 H260 EUH029 EUH032 Acute Tox. 2+ H300 Aquatic Acute 1 H400 M = 100
<b>Current proposal for consideration by RAC</b>	Acute Tox. 3 H311 Acute Tox. 2 H300
<b>Resulting harmonised classification (future entry in Annex VI, CLP Regulation)</b>	Water-react. 1 H260 EUH029 EUH032 Acute Tox. 2 H300 Acute Tox. 3 H311 Aquatic Acute 1 H400 M = 100

GHS02, GHS06, GHS09

**Figure 2\_1\_03:** Classification and Labelling: Phosphine  
(according to Directive 67/548/EEC and 1999/45/EC)

**Justified proposals for the classification and labelling of:**

**Phosphine**

**according to Directive 67/548/EEC and 1999/45/EEC**

**The product contains the following dangerous substances in the concentrations stated:**

Phosphine : F+, T+, N, R-12-17-26-34-50  
CAS No.: 7803-51-2

**Labelling of the preparation:**

Danger symbols: F+, T+, N  
Indications of danger: extremely flammable, very toxic, dangerous for the environment  
Risk phrases: R 12-17-26-34-50  
Safety phrases: S 1/2-28-36/37-45-61-63

**according to Regulation (EC) No 1272/2008**

**The product contains the following dangerous substances in the concentrations stated:**

Phosphine: GHS02, GHS04, GHS06, GHS05, GHS09  
H220, H330, H314, H400  
CAS No.: 7803-51-2

**Labelling of the preparation:**

Pictograms: GHS02, GHS04, GHS06, GHS05, GHS09  
Signal Word: Danger  
Hazard statement: H220, H330, H314, H400

Laudenbach, 12.07.2011



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Excel- Tabelle Produktzulassung Gesamtübersicht (Produktteil)

BAuA, 22.04.2008

BPD Annex point	TNsG Dossier Prep and Study Eval. Chapter No.	TNsG on Data requirements for biocidal product types	Author(s)	Year	Title	Company	Source (when different from company)	Report No	Report Date	(Un)Published	Study evaluated in CAR (Y/N)	Data protection (Y/N), Confidential ( C)	Data owner/ Sponsor (give name, address)	Letter of Access (Y/N)	GLP (Y/N)	Link
		<b>Applicant</b>														
IIB I	1.1	Name and address, etc.														
	1.2	Manufacturer/formulator of biocidal product and the active substance(s)														
		<b>Identify</b>														
IIB II.2.1	2.1	Trade name or proposed trade name,														
	2.1.1	Trade name														
	2.1.2	Manufacturer's														
IIB II.2.2	2.2	Detailed quantitative and qualitative information														
	2.2.1	Trade name														
	2.2.2	IUPAC name														
	2.2.3	CAS No.														
	2.2.4	EC No.:														
	2.2.5	Other														
	2.2.6	Molecular														
	2.2.7	Structural														
	2.2.8	Classification according														
IIB II.2.3	2.3	Physical state and														
	2.3.1	Physical														
	2.3.2	Nature														
		Exposure data														
		Physical, chemical and technical properties														
IIB III.3.1	3.1	Appearance														



Excel- Tabelle Produktzulassung Gesamtübersicht (Produktteil)

BAuA, 22.04.2008

BPD Annex point	TNSG Dossier Prep and Study Eval. Chapter No.	TNSG on Data requirements for biocidal product types	Author(s)	Year	Title	Company	Source (when different from company)	Report No	Report Date	(Un)Published	Study evaluated in CAR (Y/N)	Data protection (Y/N), Confidential (C)	Data owner/ Sponsor (give name, address)	Letter of Access (Y/N)	GLP (Y/N)	Link
	3.1.1	Physical state														
	3.1.2	Colour														
	3.1.3	Odour														
IIB III.3.2	3.2	Explosive properties														
IIB III.3.3	3.3	Oxidising properties														
IIB III.3.4	3.4_01	Flash-point and other indications of flammability or spontaneous ignition														
IIB III.3.4	3.4_02	Flash-point and other indications of flammability or spontaneous ignition														
IIB III.3.5	3.5	Acidity/alkalinity and if														
IIB III.3.6	3.6_01	Relative density														

BPD Annex point	TNSG Dossier Prep and Study Eval. Chapter No.	TNSG on Data requirements for biocidal product types	Author(s)	Year	Title	Company	Source (when different from company)	Report No	Report Date	(Un)Published	Study evaluated in CAR (Y/N)	Data protection (Y/N), Confidential (C)	Data owner/ Sponsor (give name, address)	Letter of Access (Y/N)	GLP (Y/N)	Link
IIB III.3.6	3.6_02	Relative density														
IIB III.3.7	3.7_01	Storage stability - stability and shelf-life														
IIB III.3.7	3.7_02	Storage stability - stability and shelf-life														
IIB III.3.8	3.8	Technical characteristics of the biocidal														
IIB III.3.8	3.8	Technical characteristics of the biocidal														
IIB III.3.9	3.9	Physical and chemical compatibility with other														
	3.10	Surface tension and viscosity														
	3.10.1	Surface tension														
	3.10.2	Viscosity														
	3.11	Particle size distribution														

BPD Annex point	TNSG Dossier Prep and Study Eval. Chapter No.	TNSG on Data requirements for biocidal product types	Author(s)	Year	Title	Company	Source (when different from company)	Report No	Report Date	(Un)Published	Study evaluated in CAR (Y/N)	Data protection (Y/N), Confidential (C)	Data owner/ Sponsor (give name, address)	Letter of Access (Y/N)	GLP (Y/N)	Link
		Methods of identification and analysis														
IIB IV.4.1	4.1	Analytical method for determining the concentrations of the active substance(s) in the biocidal product														
IIB IV.4.2	4.2	Analytical methods including recovery rates and the limits of determination for toxicologically and ecotoxicologically														
		Intended uses and efficacy														
IIB V.5.1	5.1	Product type and field of use envisaged														
	5.1.1	Product type														
	5.1.2	Overall use pattern														
IIB V.5.2	5.2	Method of application including description of system used														

Excel- Tabelle Produktzulassung Gesamtübersicht (Produktteil)

BAuA, 22.04.2008

BPD Annex point	TNsG Dossier Prep and Study Eval. Chapter No.	TNsG on Data requirements for biocidal product types	Author(s)	Year	Title	Company	Source (when different from company)	Report No	Report Date	(Un)Published	Study evaluated in CAR (Y/N)	Data protection (Y/N), Confidential (C)	Data owner/ Sponsor (give name, address)	Letter of Access (Y/N)	GLP (Y/N)	Link
IIB V.5.3	5.3	Application rate and if appropriate, the final concentration of the biocidal product and active substance in the system in which the preparation is to be used, e.g. cooling water, surface water, water used for heating														
IIB V.5.4	5.4	Number and timing of applications, and where relevant, any particular information relating to geographical variations, climatic variations, or necessary waiting periods to protect man and animals														

BPD Annex point	TNSG Dossier Prep and Study Eval. Chapter No.	TNSG on Data requirements for biocidal product types	Author(s)	Year	Title	Company	Source (when different from company)	Report No	Report Date	(Un)Published	Study evaluated in CAR (Y/N)	Data protection (Y/N), Confidential (C)	Data owner/ Sponsor (give name, address)	Letter of Access (Y/N)	GLP (Y/N)	Link
IIB V.5.5	5.5	Function, e.g. fungicide, rodenticide, insecticide, bactericide														
IIB V.5.6	5.6	Pest organism(s) to be														
	5.6.1	Pest organism(s) to be controlled														
	5.6.2	Products, objects or organisms to be protected														
IIB V.5.7	5.7	Effects on target organisms														
IIB V.5.8	5.8_01	Mode of action (including time delay)														
IIB V.5.8	5.8_02	Mode of action (including time delay)														
IIB V.5.8	5.8_03	Mode of action (including time delay)														
IIB V.5.8	5.8_04	Mode of action (including time delay)														
IIB V.5.9	5.9	User: industrial, professional, general public (non-professional)														

Excel- Tabelle Produktzulassung Gesamtübersicht (Produktteil)

BAuA, 22.04.2008

BPD Annex point	TNsG Dossier Prep and Study Eval. Chapter No.	TNsG on Data requirements for biocidal product types	Author(s)	Year	Title	Company	Source (when different from company)	Report No	Report Date	(Un)Published	Study evaluated in CAR (Y/N)	Data protection (Y/N), Confidential ( C)	Data owner/ Sponsor (give name, address)	Letter of Access (Y/N)	GLP (Y/N)	Link
IIB V.5.10	5.10	The proposed label claims for the product and efficacy data to support these claims, including any available standard protocols used, laboratory tests, or field trials, where appropriate														
	5.10.1	Proposed label claims for the product														
	5.10.2_01	Efficacy data														
	5.10.2_02	Efficacy data														
	5.10.2_03	Efficacy data														

BPD Annex point	TNSG Dossier Prep and Study Eval. Chapter No.	TNSG on Data requirements for biocidal product types	Author(s)	Year	Title	Company	Source (when different from company)	Report No	Report Date	(Un)Published	Study evaluated in CAR (Y/N)	Data protection (Y/N), Confidential (C)	Data owner/ Sponsor (give name, address)	Letter of Access (Y/N)	GLP (Y/N)	Link
IIB V.5.11	5.11	Any other known limitations on efficacy including resistance														
	5.11.1	Use-related restriction														
	5.11.2	Prevention of the development resistance														
	5.11.3	Concomitant use with other (biocidal) products														
		Toxicologic														
IIB VI.6.1	6.1	Acute toxicity														
IIB VI.6.1.1	6.1.1	Oral														
IIB VI.6.1.2	6.1.2	Dermal														
IIB VI.6.1.3	6.1.3	Inhalation														
IIB VI.6.1.4	6.1.4	For biocidal products that are intended to be														
IIB VI.6.2	6.2	Skin and eye irritation														

BPD Annex point	TNSG Dossier Prep and Study Eval. Chapter No.	TNSG on Data requirements for biocidal product types	Author(s)	Year	Title	Company	Source (when different from company)	Report No	Report Date	(Un)Published	Study evaluated in CAR (Y/N)	Data protection (Y/N), Confidential (C)	Data owner/ Sponsor (give name, address)	Letter of Access (Y/N)	GLP (Y/N)	Link
IIB VI.6.2	6.2.1	Skin irritation														
IIB VI.6.2	6.2.2	Eye irritation														
IIB VI.6.3	6.3	Skin sensitisation														
IIB VI.6.4	6.4	Information on dermal absorption														
IIB VI.6.5	6.5	Available toxicological data relating to toxicologically relevant non-active substances (i.e. substances of concern)														
IIB VI.6.6	6.6	Information related to the exposure of the biocidal product														
IIIB XI	6.7	Further human health-related studies														
IIIB XI.1	6.7.1	Food and feedingstuffs studies														



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IIIB XI.1.1	6.7.1.1	<i>If residues of the biocidal product remain on feedingstuffs for a significant period of time, then feeding and metabolism studies in livestock shall be required to permit evaluation of residues in food of animal origin</i>														
IIIB XI.1.2	6.7.1.2	<i>Effects of industrial processing and/or domestic preparation on the nature and magnitude of residues of the biocidal product</i>														
IIIB XI.2	6.7.2	<i>Other test(s) related to the exposure to humans</i>														
		<i>Ecotoxicology data for the biocidal product</i>														
IIIB VII.7.1	7.1	<i>Foreseeable routes of entry into</i>														
IIIB VII.7.2	7.2	<i>Information on the ecotoxicology of the</i>														

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IIB VII.7.3	7.3	Available ecotoxicological information														
IIIB XII.1	7.4	Where relevant all the information required in accordance with paragraph A7.1 and A7.2														
IIIB XII.2	7.5	Testing for distribution and dissipation in the following: (a) soil (b) water (c) air														
	(A7.5.3.1.1)	Acute oral toxicity														
IIIB XIII.1	7.6	Effects on birds														
IIIB XIII.1	7.6.1	Acute oral toxicity to birds, if not already done in accordance with Annex IIB, section VII														
IIIB XIII.2	7.7	Effects on aquatic organisms														
IIIB XIII.2.1	7.7.1	In case of application on, in, or near to surface waters														
IIIB XIII.2.1.1	7.7.1.1	Particular studies with fish and other aquatic organisms														

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IIIB XIII.2.1.2	7.7.1.2	<i>Residue data in fish concerning the active substance and including toxicologically relevant metabolites</i>														
IIIB XIII.2.1.3	7.7.1.3	<i>The studies referred to in Annex IIIA, section XIII parts 2.1, 2.2 and 2.3 may be required for relevant component of the biocidal product</i>														
IIIB XIII.2.2	7.7.2	<i>If the biocidal product is to be sprayed near to surface waters then an overspray study may be required to assess risks to aquatic organisms under field conditions</i>														
	7.8	<i>Effects on other non-target organisms</i>														
IIIB XIII.3.1	7.8.1	<i>Toxicity to terrestrial vertebrates other than birds</i>														
IIIB XIII.3.2	7.8.2	<i>Toxicity to honeybees</i>														

Excel- Tabelle Produktzulassung Gesamtübersicht (Produktteil)

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IIIB XIII 3.3	7.8.3	<i>Effects on other beneficial arthropods other than bees</i>														
IIIB XIII 3.4	7.8.4	<i>Effects on earthworms and other soil non-target macro-organisms, believed to be at risk</i>														
IIIB XIII 3.5	7.8.5	<i>Effects on soil non-target micro-organisms</i>														
IIIB XIII 3.6	7.8.6	<i>Effects on any other specific, non-target organisms (flora and fauna) believed to be at risk</i>														
IIIB XIII.3.7	7.8.7	<i>If the biocidal product is in the form of bait or granules.</i>														
IIIB XIII.3.7 1	7.8.7.1	<i>Supervised trials to assess risks to non-target organisms under field conditions</i>														

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IIB XIII.3.7.2	7.8.7.2	Studies on acceptance by ingestion of the biocidal product by any non-target organisms thought to be at risk														
		Measures to be adopted														
IIB VIII.8.1	8.1	Recommended methods and precautions concerning handling, use, storage, transport or fire														
IIB VIII.8.2	8.2	Specific treatment in case of an accident, e.g. first-aid measures, antidotes, medical treatment if available; emergency measures to protect the environment														
IIB VIII.8.3	8.3_01	Procedures, if any, for cleaning application equipment														

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	8.3_01	Procedures, if any, for cleaning application equipment														
	8.3_01	Procedures, if any, for cleaning application equipment														
IIB VIII.8.4	8.4	Identity of relevant combustion products in cases of fire														
IIB VIII.8.5	8.5	Procedures for waste management of the biocidal product and its packaging for industry, professional users and the general public (non-professional users)														
IIB VIII.8.6	8.6	Possibility of destruction or decontamination														
IIB VIII.8.7	8.7	Observations on undesirable or unintended side-effects, e.g. on beneficial and other non-target organisms														

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IIB VIII.8.8	8.8	Specify any repellents or poison control measures included in the preparation that are present to prevent action against non-target														
		Classification, packaging and labelling														
IIB IX	9.	Classification & labelling														
IIB IX		Packaging and labelling														
		Safety data														
		SDS (b.p.)														
		SDS (b.p.) (substances of concern)														
		Confidential data														
		Reference lists														
		Reference list, by section number														
		Reference list, by author (in alphabetical order)														
		Profile and results of														