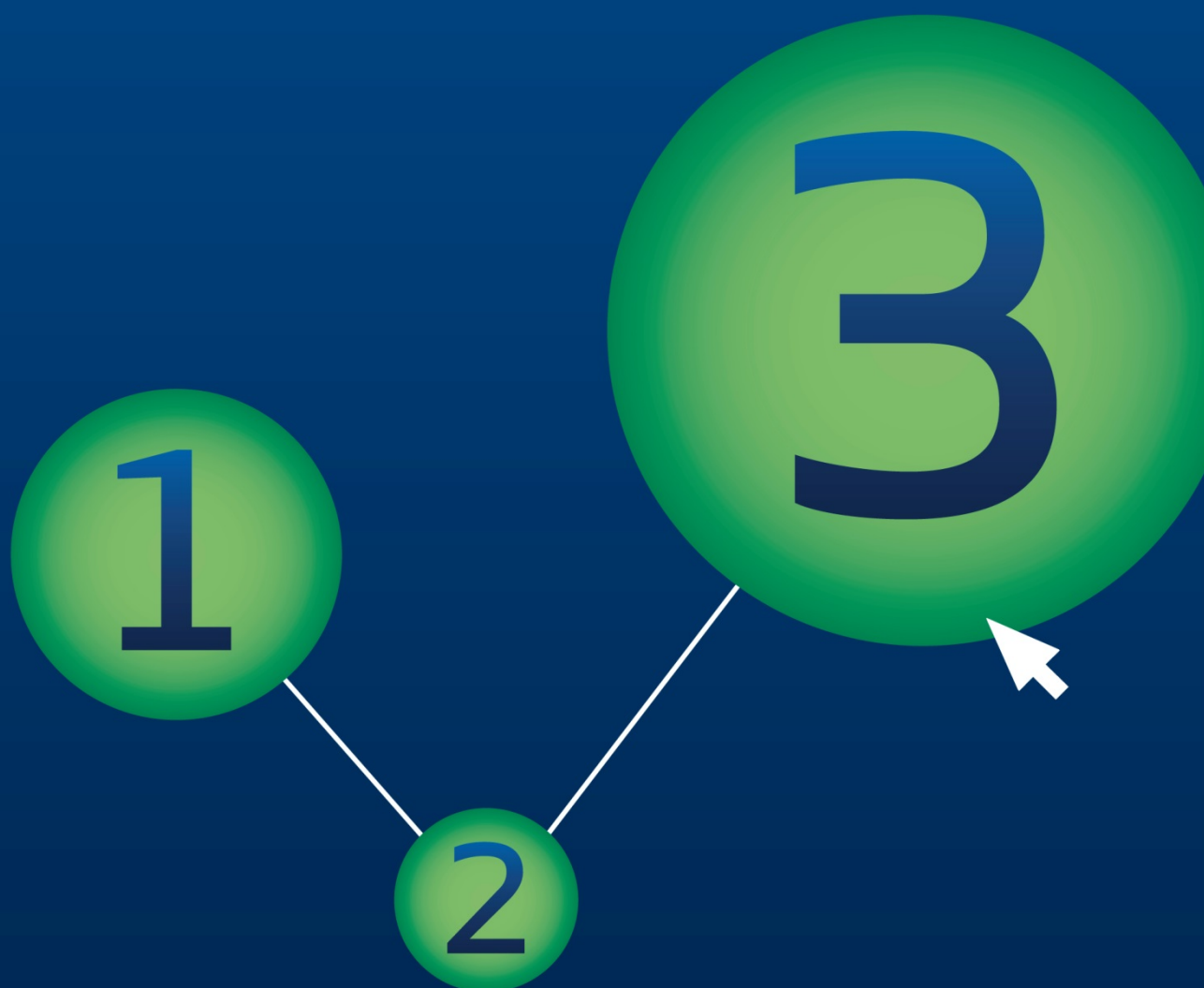


Biocides Submission Manual

Process of dissemination
Part 4: IUCLID sections 9-13



Version	Changes	Date
Version 1.0	First version	November 2015
Version 2.0	IUCLID 6 screenshots, completely revised	July 2016

BSM Process of dissemination, Part 4: IUCLID sections 9-13

Reference: ECHA-16-B-29-EN

Publ. date: July 2016

Language: EN

© European Chemicals Agency, 2016

Cover page © European Chemicals Agency

Legal notice:

Neither the European Chemicals Agency nor any person acting on behalf of the European Chemicals Agency is responsible for the use which might be made of the following information. A wealth of additional information on the European Chemicals Agency is available on the Internet. It can be accessed through the ECHA website (<http://echa.europa.eu>).

Reproduction is authorised provided the source is fully acknowledged in the form "Source: European Chemicals Agency, <http://echa.europa.eu/>", and provided written notification is given to the ECHA Communications Unit (publications@echa.europa.eu).

If you have questions or comments in relation to this document please send them (quote the reference and issue date) using the information request form. The information request form is accessible from the Contact ECHA page at: <http://echa.europa.eu/contact>

European Chemicals Agency

Mailing address: P.O. Box 400, 00121 Helsinki, Finland

Visiting address: Annankatu 10, Helsinki, Finland

Table of Contents

Introduction	5
Objective	5
Biocides Submission Manuals	5
Article 67 of the Biocidal Products Regulation (BPR)	6
The Dissemination portal	6
Web address	6
Searching in the portal	7
Intellectual property rights	7
The Dissemination process	7
When is the information published?	7
Filtering	8
Publication	8
Dissemination preview for IUCLID	8
Confidentiality	9
Process of dissemination, Part 3: IUCLID sections 9-13	10
Legend	11
Note to the legend	11
9. Ecotoxicological studies (also in Biocidal Product)	13
9.1 Toxicity to aquatic organisms	17
9.1.1 Short-term toxicity testing on fish (also in Biocidal Product under 9.2.1.1)	17
9.1.2 Short-term toxicity to aquatic invertebrates (also in Biocidal Product under 9.2.1.2)	23
9.1.3 Growth inhibition study on algae (also in Biocidal Product under 9.2.1.3)	29
9.1.5 Inhibition of microbial activity (also in Biocidal Product under 9.2.1.5)	35
9.1.6 Further toxicity studies on aquatic organisms (also in Biocidal Product under 9.2.1.6)	41
9.1.6.1 Long-term toxicity testing on fish (also in Biocidal Product under 9.2.1.6.1)	42
9.1.6.2 Long-term toxicity testing on invertebrates (also in Biocidal Product under 9.2.1.6.2)	48
9.1.7 Bioaccumulation in appropriate aquatic species (also in Biocidal Product under 9.2.1.7)	54
9.1.8 Effects on any other specific, non-target organisms (flora and fauna) believed to be at risk (also in Biocidal Product under 9.2.1.8)	60
9.1.9 Studies on sediment dwelling organisms (also in Biocidal Product under 9.2.1.9)	66
9.1.10 Effects on aquatic macrophytes (also in Biocidal Product under 9.2.1.10)	72
9.2 Terrestrial toxicity, initial tests	78
9.2.1 Effects on soil microorganisms (also in Biocidal Product under 9.2.2.1)	78
9.2.2 Effects on earthworms or other soil-dwelling non-target invertebrates (also in Biocidal Product under 9.2.2.2)	83
9.2.3 Acute toxicity to plants (also in Biocidal Product under 9.2.2.3)	89
9.4 Effects on birds (also in Biocidal Product under 9.2.4)	95
9.5 Effects on arthropods (also in Biocidal Product under 9.2.5)	101
9.6 Bioconcentration terrestrial (also in Biocidal Product under 9.2.6)	107
9.8 Effects on other non-target, non-aquatic organisms (also in Biocidal Product under 9.2.8)	112

9.10 Identification of endocrine activity (<i>also in Biocidal Product under 9.2. Further Ecotoxicological studies</i>)	118
10. Environmental fate and behaviour (<i>also in Biocidal Product</i>)	122
10.1 Fate and behaviour in water and sediment.....	123
10.1.1 Degradation, initial studies	123
10.1.1.1 Abiotic	123
10.1.1.1.a Abiotic: Hydrolysis as a function of pH and identification of breakdown products	123
10.1.1.1.b Abiotic: Phototransformation in water, including identification of transformation products ...	129
10.1.1.2 Biotic → Ready / Inherent biodegradability.....	135
10.1.2 Adsorption / desorption.....	140
10.1.3 Rate and route of degradation including identification of metabolites and degradation products	146
10.1.5 Field studies.....	152
10.2 Fate and behaviour in soil.....	156
10.2.1 Laboratory study on rate and route of degradation including identification of the processes involved.....	156
10.2.3 Soil accumulation studies (<i>also in Biocidal Product under 10.1 Foreseeable routes of entry into the environment on the basis of the use envisaged</i>)	162
10.3 Fate and behaviour in air.....	166
10.3.1 Phototransformation in air	166
10.3 <i>Biocidal Product – Leaching behaviour</i>	171
10.4 Additional information on environmental fate and behaviour (<i>also in Biocidal Product under 10.2 Further studies on fate and behaviour in the environment</i>).....	175
10.4 <i>Biocidal Product – Testing for distribution and dissipation</i>	179
10.6 Monitoring: Identification of all degradation products (>10%) in the studies on degradation in soil, water and sediments	183
11. Measures to protect humans, animals and the environment.....	187
11. <i>Biocidal Product – Measures to protect humans, animals and the environment</i>	189
12. Classification & Labelling.....	191
12.1 GHS.....	191
12.1 <i>Biocidal Product – GHS</i>	195
12.2 DSD – DPD	199
12.2 <i>Biocidal Product – DSD-DPD</i>	201
12.3 <i>Biocidal Product – Packaging (12.7. in Annex III of BPR)</i>	203
13. Summary and evaluation (<i>also in Biocidal Product</i>)	204
13.1 PBT assessment	205

Introduction

Objective

This manual gives guidance on the online access provided by ECHA to information on active substances approved, and biocidal products authorised, under the Biocidal Products Regulation (BPR) (where the data is in IUCLID). It is aimed at industry, and in particular, at managers and technical experts in companies who are responsible for making sure that comprehensive information is entered in the dossiers.

Biocides Submission Manuals

This manual is part of the Biocides Submission Manual (BSM) series concerning technical guides, application instructions and process manuals and also includes:

Technical guides:

Using IUCLID, which describes how to prepare a general IUCLID dossier, giving you details on the different functionalities in IUCLID, as well as explaining the different sections contained within a dossier.

Using R4BP 3, which describes how to create user accounts in R4BP 3 through ECHA Accounts and gives a detailed description of the various functionalities of the system.

Using the SPC Editor, which describes how to prepare a summary of the product characteristics (SPC) required for certain application types.

Application instructions:

Application instruction manuals give guidance on how to submit applications concerning various processes concerning active substance approvals and biocidal product authorisations.

- **Active substances**
- **National authorisations**
- **Simplified authorisations**
- **Union authorisations**
- **Technical equivalence and chemical similarity**

Process manuals:

Invoicing in R4BP 3, which describes the general information related to invoices and credit notes issued by ECHA following the submission of an application.

Confidentiality requests for biocide applications, which describes how to make confidentiality claims in IUCLID and which dossier information can be claimed confidential.



All of the Biocides Submission Manuals, including the technical guides, application instructions and related processes can be found from the [ECHA website](#).

Article 67 of the Biocidal Products Regulation (BPR)

Article 67 of the Biocidal Products Regulation (BPR) stipulates that the European Chemicals Agency (ECHA) shall publish certain information it holds on approved biocidal active substances and authorised biocidal products free of charge over the internet. This information is published on the ECHA website, in the section 'Information on Chemicals'.

However, in certain cases information can be withheld, if the applicant submitting the information also indicates they wish to keep the information confidential, and submits a justification as to why publishing the information would be potentially harmful to the commercial interests of the applicant or any other party concerned. Relevant authority will then assess such justifications in accordance with Article 67(3) and (4), and where the justification is accepted, the information concerned will not be published. Claiming information confidential may be subject to a fee.

Where urgent action is essential to protect human health, safety or the environment, such as emergency situations, ECHA may disclose additional information, in accordance with Article 66(2). This will not happen in an automated way, and is therefore not covered in this manual.

This manual provides information about the online access to information on active substances approved, and biocidal products authorised, under the BPR.¹ The document will help readers to understand:

- what are the steps in the dissemination process; and
- which information will be made publicly available on the ECHA website.

The manual presents what ECHA directly publishes from the IUCLID dossiers and it does not cover information from other documents such as SPCs², assessment reports, etc.

Because of technical progress with IUCLID and the Dissemination portal, this manual will be regularly updated. We recommend that you check our website regularly to make sure that you have the most recent version of the document.

In addition to this manual, a specific dissemination preview will be made available for applicants in the near future. This tool will enable applicants to verify – when preparing a dossier in IUCLID – which information will be published on the ECHA website, as explained further below (see [Dissemination preview for IUCLID](#)).

The Dissemination portal

Web address

Information on chemicals can be accessed through the ECHA website; detailed information on approved active substances or authorised biocidal products can be accessed through the ECHA website following the below paths:

Active substances: ECHA Website > Information on Chemicals > Biocidal Active Substances:
<http://echa.europa.eu/information-on-chemicals/biocidal-active-substances>

¹ The dissemination obligations under Article 67 of the BPR also apply to information on active substances and biocidal products where the approval/authorisation application was submitted under the Biocidal Products Directive (BPD) (Directive 98/8/EC). The process for doing so is different to that described in this manual due to the use of IUCLID for the BPR dossiers.

² The information contained in the SPC associated with the authorisation of a biocidal product will be published on the ECHA website. It is the responsibility of the authorisation holder and Member State competent authority to make sure that this document does not contain confidential information.

Biocidal products: ECHA Website > Information on Chemicals > Biocidal Products:

<http://echa.europa.eu/information-on-chemicals/biocidal-products>

Searching in the portal

Before you can begin to search for active substances and/or biocidal products in the Dissemination portal, you have to acknowledge that you have taken note of the Legal Notice explaining the nature of the information, the right to use the information, and possible limits to the right to use the information.

You can search for an active substance in the portal by its name, EC number, CAS number, type, biocide ID and approval information.

A biocidal product can be searched by its name, asset number, product-type and name, EC number or CAS number, approval ID of its active substance as well as by authorisation data.

A list of all the active substances or biocidal products in the database can be obtained by searching without entering any search criteria.

Search results can be reordered by clicking on the individual column headers.

ECHA has also developed InfoCards and Brief Profiles for substances where details on the substance classification, uses and exposure and scientific properties are summarised and aggregated. Whilst currently they are primarily based on the data submitted in REACH registrations, they also include data from other sources, including the C&L Inventory, other REACH regulatory processes, and data from the BPR and PIC regulation.

Intellectual property rights

You should use the information available in the Dissemination portal with care. Reproduction and use or further distribution of the information is subject to copyright laws and may require the permission of the owner of that information.

Data protection periods apply to data submitted for the purposes of the BPD or of the BPR. Remember that in line with Article 59(1) of the BPR, the competent authorities may not use that data for the benefit of a subsequent applicant unless it has a letter of access or the data protection periods have expired.

The data protection periods are set in Article 60 of the BPR ("*Protection of Data held by competent authorities or the Agency*") and Article 95(5) of the BPR ("*Transitional measures concerning access to the active substance dossier*").

The Dissemination process

When is the information published?

For active substances, the process of disseminating information from an application dossier starts as soon as the European Commission has adopted an Implementing Regulation providing that the active substance is approved.

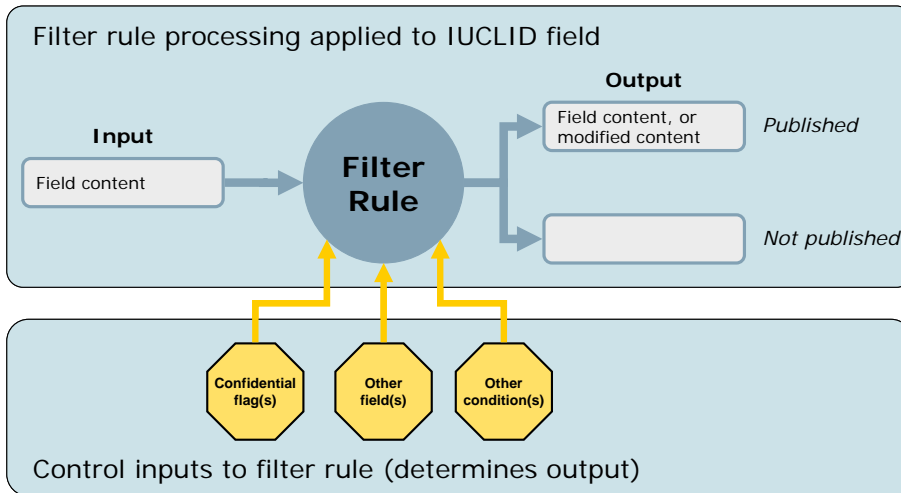
For biocidal products, the dissemination process starts from the date a biocidal product is authorised.

At this point, the associated dossier – which by now only contains confidentiality requests upheld by the relevant authority – is prepared for dissemination, as described below:

Filtering

The most important step in the dissemination process of the IUCLID information is the filtering step in which information not meant to be published is removed from the dossier, along with information flagged or claimed to be confidential (Figure 1).

Figure 1: Filter rules



The filtering of IUCLID dossiers is performed by an IT tool which has been programmed with Filter rules. Filter Rules are based on the Article 67 of the BPR and are applied to each field in the IUCLID dossier determining whether the field content should be published or not. After the filtering step is completed, the filtered dossier will contain only the information from the fields that is to be published.

Dossier filtering (the removal of information not to be published) is an automated process and is independent of which text you provide in a certain field. Please review your dossier before you submit it, to make sure the correct content is provided. If you provide confidential information in a field which is set to be published (e.g. the guidance on safe use), **the information will become visible on the internet.**

In this manual (consisting of four separate parts), you can see the filter rule that applies for all IUCLID 6 fields from the *BPR Active substance information* and *BPR Biocidal product authorisation* templates. The filter rules are explained in the [legend](#).

The other BPR templates for the substances (substances of concern and non-active substances) and representative biocidal products are currently not covered in this manual. Further information on how this information will be published will be made available in due time.

Publication

ECHA disseminates information at substance/product level. All documents and information from a IUCLID dossier will be linked to other relevant substance data identifiers. After the filtering step, the dossier is processed to create a set of html web pages. A batch of these web pages is regularly published on the ECHA website. As part of the same step, the relevant data and metadata are published to allow the search results to be searched and filtered.

Dissemination preview for IUCLID

ECHA is developing a IUCLID plugin to enable applicants to simulate which information from their IUCLID dossier will be removed during filtering, and which information will be made publicly available over the internet. The Dissemination preview will allow applicants to use it while they are preparing their BPR dossier in IUCLID. The purpose of the tool is to help

applicants to prepare dossiers that can be published without revealing confidential business information. More information on the tool will be available on the ECHA website in due time.

Confidentiality

The IUCLID template allows applicants to set confidentiality request flags on information covered by Article 67(3) and (4) of the BPR. For information that an applicant wishes to keep confidential, a confidentiality request must be justified and the request will only be implemented if it is upheld by the relevant authority. For guidance on confidentiality claims, please see the Biocides Submission Manual on the Process of confidentiality requests for biocide applications.³

³ http://echa.europa.eu/documents/10162/14938692/bsm_10_confidentiality_requests_en.pdf

Process of dissemination, Part 3: IUCLID sections 9-13







In this manual, you can find the filter rule – illustrated with screenshots – for every field in sections 9, 10, 11, 12 and 13 of the IUCLID 6 dossier. The filter rule automatically determines if the content of the field is published on the ECHA website or not.

The legend on the next page briefly explains the different filter rules.

Unless otherwise stated, the rules of the section refer to the active substance template.

To allow for easy navigation, the sections in this manual use the same numbering as the sections in IUCLID in the appropriate BPR view. For information on the other sections of the IUCLID dossier, please consult the relevant part of the manual (the manual is divided into four parts, each dealing with separate groups of IUCLID sections).

Legend

	<p>Is always automatically published.</p> <ul style="list-style-type: none"> This rule concerns information listed in Article 67(1) and (2).
	<p>Is not automatically published.</p> <ul style="list-style-type: none"> This rule concerns information normally deemed to undermine the protection of the commercial interests of the persons concerned, or information which does not relate to the hazard and safe use of the substance.
	<p>Is automatically published unless confidentiality has been claimed on this section.</p> <ul style="list-style-type: none"> This rule concerns information listed in Article 67(3) and (4).
	<p>Fields that are always automatically published in the active substance dossier can be claimed confidential in the biocidal product dossier. Confidentiality flags and indication of endpoint addressed will be published.</p> <ul style="list-style-type: none"> This rule concerns information listed in Article 67(4).
	<p>The bibliographic references author, title, and source are published according to the following rule, with the most important criteria listed first:</p> <ul style="list-style-type: none"> are not published if the endpoint record is claimed confidential, unless the reference type is publication, review article or handbook; are not published if the reference type is study report or company data; are not published if at least one of following is provided: testing lab, report number, owner company or study number.
	<p>This information will be published from other source (e.g. SPC, R4BP) and not from the IUCLID dossier.</p>

Any confidentiality claim must be justified and the claim will only be implemented if it is upheld by the relevant authority. Justifications for claiming the confidentiality are not published.

Note to the legend

Field with a link

Fields containing a link to a record or information do not carry a filter rule as information they contain is published according to the individual filter rules set within the given record or information to which they refer.

Test material and identity of transformation products – conditioned filter rule

The test material and the identity of transformation products will be published unless:

- the reference substance describing the material itself is flagged confidential, or
- the endpoint study record is flagged confidential.

Justification for type of information – conditioned filter rule

Justification for type of information will be published unless:

- the reference substances linked to the endpoint study record have been flagged confidential, or
- the endpoint study record is flagged confidential.

For read-across, the information is not published if the study record in the related information is flagged confidential, or the test material reference substance in the related information is flagged confidential.

(Robust) study summary data

Fields referring to (robust) study summary data will only be published if the endpoint study record is not requested confidential.

9. Ecotoxicological studies (also in Biocidal Product)

NB: Biocidal Product **A** → **C**

Administrative data ^

Hazard for aquatic organisms ^

Freshwater ^

Hazard assessment conclusion

A ...

A **A** ...

Assessment factor

A

Extrapolation method

A ...

PNEC freshwater (intermittent releases)

A **A** ...

Explanations on the hazard conclusion

Normal Agency FB 8 A B I U

N

Marine water ^

Hazard assessment conclusion

A ...

A **A** ...

Assessment factor

A

Extrapolation method

A ...

PNEC marine water (intermittent releases)

A **A** ...

Explanations on the hazard conclusion

Normal Agency FB 8 A B I U

N

STP ^

Hazard assessment conclusion

A ...

A **A** ...

Assessment factor

A

Extrapolation method

A ...

Explanations on the hazard conclusion

Normal Agency FB 8 A B I U

N

Sediment (freshwater) ^

Hazard assessment conclusion

A ...

A ...

Assessment factor

A

Extrapolation method

A ...

Explanations on the hazard conclusion

Normal Agency FB 8 A B I U

N

Sediment (marine water) ^

Hazard assessment conclusion

A ...

A **A** ...

Assessment factor

A

Extrapolation method

A ...

Explanations on the hazard conclusion

Normal Agency FB 8 A B I U

N

Hazard for air ^


Air ^

Hazard assessment conclusion

...

...

Explanations on the hazard conclusion



Hazard for terrestrial organisms ^

Soil ^

Hazard assessment conclusion

...


...

Assessment factor

Extrapolation method

...

Explanations on the hazard conclusion



Hazard for predators ^

Secondary poisoning ^


Hazard assessment conclusion

...

...

Assessment factor

Explanations on the hazard conclusion



Additional information ^

Normal Agency FB 8 A B I U [list icons] A [undo] [redo]

C

Conclusion on classification ^

Normal Agency FB 8 A B I U [list icons] A [undo] [redo]

A

Administrative data ^

Endpoint
A ... **Remarks** N ...

Type of information
A ... **Other** A ... **Remarks** N ...

Adequacy of study
A ...
 Robust study summary N
 Used for classification N
 Used for SDS N

Study period
C ...

Reliability
A ... **Other** A ...

Rationale for reliability incl. deficiencies
C ... **Other** C ... **Remarks** N ...

Data waiving
C ...

Justification for data waiving
C

Justification for type of information
C

Attached justification

Attached justification	Reason / purpose
N	N

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
A	A	N

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	A	A	B	N	N	A	N	A	N

+ Add... Delete Move up Move down Go to link target

Data access
N ... **Other** N ... **Remarks** N ...

Data protection claimed
N ... **Remarks** N ...

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

+ Add... Edit... X Delete ↑ Move up ↓ Move down

Principles of method if other than guideline

A | X

A ...

GLP compliance

C ... Remarks **N** ...

Test material ^

Test material information

Specific details on test material used for the study

A | X

C ...

Sampling and analysis ^

Analytical monitoring

C ... Remarks **N** ...

Details on sampling

A | X

C ...

Details on analytical methods

A | X

N ...

Test solutions ^

Vehicle

C ... Remarks **N** ...

Details on test solutions

A | X

C ...

Test organisms ^

Test organisms (species)

A ... Other **A** ...

Details on test organisms

A | X

C ...

Study design ^

Test type

A ... Other **A** ...

Water media type

A ... Other **A** ... Remarks **N** ...

Attached full study report

N

+ Add... Open Properties X Delete Move up Move down

Illustration (picture/graph)

N

Zoom Load Delete

Applicant's summary and conclusion ^

Validity criteria fulfilled

A ... Remarks N ...

Conclusions

C ...

Executive summary

Normal Aharoni 8 A B I U ...

C

9.1.2 Short-term toxicity to aquatic invertebrates (also in Biocidal Product under 9.2.1.2)

NB: Biocidal Product A → C

Administrative data ^

A

Link to relevant study record(s) ^

Study name / type

+ Add...
✕ Delete
↑ Move up
↓ Move down
➤ Go to link target

Description of key information ^

A

Key value for chemical safety assessment ^

EC50/LC50 for freshwater invertebrates

A

A ...

EC50/LC50 for marine water invertebrates

A

A ...

Additional information ^

C

Administrative data ^

Endpoint A ... Remarks N ...

Type of information A ... Other A ... Remarks N ...

Adequacy of study A ...

Robust study summary N

Used for classification N

Used for SDS N

Study period C ...

Reliability A ... Other A ...

Rationale for reliability incl. deficiencies C ... Other C ... Remarks N ...

Data waiving C ...

Justification for data waiving C

Justification for type of information C

Attached justification

Attached justification	Reason / purpose
N	N

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
A	A	N

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	A	A	B	N	N	A	N	A	N

+ Add... Delete Move up Move down Go to link target

Data access N ... Other N ... Remarks N ...

Data protection claimed N ... Remarks N ...

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

+ Add... Edit... Delete Move up Move down

Principles of method if other than guideline

A | X

A

GLP compliance

C ... Remarks N

Test material ^

Test material information

Specific details on test material used for the study

A | X

C

Sampling and analysis ^

Analytical monitoring

C ... Remarks N

Details on sampling

A | X

C

Details on analytical methods

A | X

N

Test solutions ^

Vehicle

C ... Remarks N

Details on test solutions

A | X

C

Test organisms ^

Test organisms (species)

A ... Other A

Details on test organisms

A | X

C

Study design ^	
Test type	<input type="text" value="A"/> ... Other <input type="text" value="A"/> ...
Water media type	<input type="text" value="A"/> ... Other <input type="text" value="A"/> ... Remarks <input type="text" value="N"/> ...
Limit test	<input type="text" value="A"/> ...
Total exposure duration	<input type="text" value="A"/> <input type="text" value="A"/> ...
Remarks on exposure duration	<input type="text" value="A"/> ...
Post exposure observation period	<input type="text" value="C"/> ...
Test conditions ^	
Hardness	<input type="text" value="C"/> ...
Test temperature	<input type="text" value="C"/> ...
pH	<input type="text" value="C"/> ...
Dissolved oxygen	<input type="text" value="C"/> ...
Salinity	<input type="text" value="C"/> ...
Conductivity	<input type="text" value="C"/> ...
Nominal and measured concentrations	<input type="text" value="C"/> ...
Details on test conditions	<input type="text" value="A"/> <input type="text" value="X"/> ...
Reference substance (positive control)	<input type="text" value="A"/> ... Remarks <input type="text" value="N"/> ...

Any other information on materials and methods incl. tables ^

Normal Aharoni 8 A B I U

N

Results and discussion ^

Effect concentrations

Key result	Duration	Dose descriptor	Effect conc.	Nominal / measured	Conc. based on	Basis for effect	Remarks on result
A	A	A	A	A	A	A	A

+ Add... Edit... Delete Move up Move down

Details on results

A X

A

Results with reference substance (positive control)

A X

C

Reported statistics and error estimates

C

Any other information on results incl. tables ^

Normal Aharoni 8 A B I U


A

Overall remarks, attachments ^

Overall remarks

Normal Aharoni 8 A B I U

N

Attached background material	
Attached document	Remarks
N	N
+ Add... Edit... Delete Move up Move down	
Attached full study report	
N	
+ Add... Open Properties Delete Move up Move down	
Illustration (picture/graph)	
N	
Zoom + Load Delete	
Applicant's summary and conclusion ^	
Validity criteria fulfilled	
A	Remarks N
Conclusions	
C	
Executive summary	
	
C	

9.1.3 Growth inhibition study on algae (also in Biocidal Product under 9.2.1.3)

NB: Biocidal Product **A** → **C**

Administrative data ^

A

Link to relevant study record(s) ^

Study name / type

+ Add... X Delete ↑ Move up ↓ Move down > Go to link target

Description of key information ^

Normal Aharoni 8 A^T | B | I | U | 1/2 = A⁹ ↻

A

Key value for chemical safety assessment ^

EC50/LC50 for freshwater algae **A** **A** ...

EC50/LC50 for marine water algae **A** **A** ...

EC10, LC10 or NOEC for freshwater algae **A** **A** ...

EC10, LC10 or NOEC for marine water algae **A** **A** ...

Additional information ^

Normal Aharoni 8 A^T | B | I | U | 1/2 = A⁹ ↻

C

Administrative data ^

Endpoint
A ... Remarks N ...

Type of information
A ... Other A ... Remarks N ...

Adequacy of study
A ...
 Robust study summary N
 Used for classification N
 Used for SDS N

Study period
C ...

Reliability
A ... Other A ...

Rationale for reliability incl. deficiencies
C ... Other C ... Remarks N ...

Data waiving
C ...

Justification for data waiving
C

Justification for type of information
C

Attached justification

Attached justification	Reason / purpose
N	N

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
A	A	N

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	A	A	B	N	N	A	N	A	N

+ Add... Delete Move up Move down Go to link target

Data access
N ... Other N ... Remarks N ...

Data protection claimed
N ... Remarks N ...

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Principles of method if other than guideline

A ↕ ✕ ▼

A

...

GLP compliance

C
...
Remarks
N
...

Test material ^

Test material information

> 🔗 ✕

Specific details on test material used for the study

A ↕ ✕ ▼

C

...

Sampling and analysis ^

Analytical monitoring

C
...
Remarks
N
...

Details on sampling

A ↕ ✕ ▼

C

...

Details on analytical methods

A ↕ ✕ ▼

N

...

Test solutions ^

Vehicle

C
...
Remarks
N
...

Details on test solutions

A ↕ ✕ ▼

C

...

Test organisms ^

Test organisms (species)

A
...
Other
A
...

Details on test organisms

A ↕ ✕ ▼

C

...

Study design ^	
Test type	<input type="text" value="A"/> ... Other <input type="text" value="A"/> ...
Water media type	<input type="text" value="A"/> ... Other <input type="text" value="A"/> ... Remarks <input type="text" value="N"/> ...
Limit test	<input type="text" value="A"/> ...
Total exposure duration	<input type="text" value="A"/> <input type="text" value="A"/> ...
Remarks on exposure duration	<input type="text" value="A"/> ...
Post exposure observation period	<input type="text" value="C"/> ...
Test conditions ^	
Hardness	<input type="text" value="C"/> ...
Test temperature	<input type="text" value="C"/> ...
pH	<input type="text" value="C"/> ...
Dissolved oxygen	<input type="text" value="C"/> ...
Salinity	<input type="text" value="C"/> ...
Conductivity	<input type="text" value="C"/> ...
Nominal and measured concentrations	<input type="text" value="C"/> ...
Details on test conditions	<input type="text" value="C"/> ...
Reference substance (positive control)	<input type="text" value="A"/> ... Remarks <input type="text" value="N"/> ...

9.1.5 Inhibition of microbial activity (also in Biocidal Product under 9.2.1.5)

NB: Biocidal Product **A** → **C**

Administrative data ^

A

Link to relevant study record(s) ^

Study name / type

+ Add... X Delete ↑ Move up ↓ Move down > Go to link target

Description of key information ^

Normal Aharoni 8 A[▼] | **B** | *I* | U | [List icons] | [Table icons] | [Link icon] | [Undo] | [Redo]

A

Key value for chemical safety assessment ^

EC50 or LC50 for microorganisms

A **A** ...

EC10, LC10 or NOEC for microorganisms

A **A** ...

Additional information ^

Normal Aharoni 8 A[▼] | **B** | *I* | U | [List icons] | [Table icons] | [Link icon] | [Undo] | [Redo]

C

Administrative data ^

Endpoint
 ... Remarks ...

Type of information
 ... Other ... Remarks ...

Adequacy of study
 ...
 Robust study summary
 Used for classification
 Used for SDS

Study period
 ...

Reliability
 ... Other ...

Rationale for reliability incl. deficiencies
 ... Other ... Remarks ...

Data waiving
 ...

Justification for data waiving

Justification for type of information

Attached justification

Attached justification	Reason / purpose
<input type="text" value="N"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
<input type="text" value="B"/>	<input type="text" value="B"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="B"/>	<input type="text" value="N"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Delete Move up Move down Go to link target

Data access
 ... Other ... Remarks ...

Data protection claimed
 ... Remarks ...

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Principles of method if other than guideline

A
✕

A
⋮

GLP compliance

C
Remarks N

Test material ^

Test material information

> ∞ ✕

Specific details on test material used for the study

A
✕

C
⋮

Sampling and analysis ^

Analytical monitoring

C
Remarks N

Details on sampling

A
✕

C
⋮

Details on analytical methods

A
✕

N
⋮

Test solutions ^

Vehicle

C
Remarks N

Details on test solutions

A
✕

C
⋮

Test organisms ^

Test organisms (species)

A
Other N

Details on inoculum

A
✕

C
⋮

Study design	
Test type	<input type="text" value="A"/> ... <input type="text" value="Other"/> <input type="text" value="A"/> ...
Water media type	<input type="text" value="A"/> ... <input type="text" value="Other"/> <input type="text" value="A"/> ... <input type="text" value="Remarks"/> <input type="text" value="N"/> ...
Limit test	<input type="text" value="A"/> ...
Total exposure duration	<input type="text" value="A"/> <input type="text" value="A"/> ...
Remarks on exposure duration	<input type="text" value="A"/> ...
Post exposure observation period	<input type="text" value="C"/> ...
Test conditions	
Hardness	<input type="text" value="C"/> ...
Test temperature	<input type="text" value="C"/> ...
pH	<input type="text" value="C"/> ...
Dissolved oxygen	<input type="text" value="C"/> ...
Salinity	<input type="text" value="C"/> ...
Conductivity	<input type="text" value="C"/> ...
Nominal and measured concentrations	<input type="text" value="C"/> ...
Details on test conditions	<input type="text" value="A"/> <input type="text" value="X"/>
	<input type="text" value="C"/> ...
Reference substance (positive control)	<input type="text" value="A"/> ... <input type="text" value="Remarks"/> <input type="text" value="N"/> ...

Any other information on materials and methods incl. tables ^

Normal Aharoni 8 A B I U | | | | 1 2 = | A

N

Results and discussion ^

Effect concentrations

Key result	Duration	Dose descriptor	Effect conc.	Nominal / measured	Conc. based on	Basis for effect	Remarks on result
A	A	A	A	A	A	A	A

+ Add...
Edit...
Delete
Move up
Move down

Details on results

A X

A

Results with reference substance (positive control)

A X

C

Reported statistics and error estimates

C

Any other information on results incl. tables ^

Normal Aharoni 8 A B I U | | | | 1 2 = | A

A

Overall remarks, attachments ^

Overall remarks

Normal Aharoni 8 A B I U | | | | 1 2 = | A

N

Attached background material

Attached document	Remarks
N	N

+ Add... Edit... Delete Move up Move down

Attached full study report

N

+ Add... Open Properties Delete Move up Move down

Illustration (picture/graph)

N

Zoom Load Delete

Applicant's summary and conclusion ^

Validity criteria fulfilled

A	Remarks	N
---	---------	---

Conclusions

C


Executive summary

Normal Aharoni 8 A B I U [text alignment icons] [bullet list icon] [numbered list icon] [undo icon] [redo icon]


C

9.1.6 Further toxicity studies on aquatic organisms (also in Biocidal Product under 9.2.1.6)


Administrative data ^




Description of key information ^




Normal Aharoni 8 A B I U



Additional information ^



Normal Aharoni 8 A B I U



9.1.6.1 Long-term toxicity testing on fish (*also in Biocidal Product under 9.2.1.6.1*)

NB: Biocidal Product **A** → **C**

Administrative data ^

A

Link to relevant study record(s) ^

Study name / type

+ Add...
✕ Delete
↑ Move up
↓ Move down
➤ Go to link target

Description of key information ^

A

Key value for chemical safety assessment ^

EC10, LC10 or NOEC for freshwater fish

A

A ...

EC10, LC10 or NOECEC10, LC10 or NOEC for marine water fish

A

A ...

Additional information ^

C

Administrative data ^

Endpoint
 ... Remarks

Type of information
 ... Other ... Remarks

Adequacy of study
 ...
 Robust study summary
 Used for classification
 Used for SDS

Study period

Reliability
 ... Other

Rationale for reliability incl. deficiencies
 ... Other ... Remarks

Data waiving

Justification for data waiving

Justification for type of information

Attached justification

Attached justification	Reason / purpose
<input type="text" value="N"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
<input type="text" value="B"/>	<input type="text" value="B"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="B"/>	<input type="text" value="N"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Delete Move up Move down Go to link target

Data access
 ... Other ... Remarks

Data protection claimed
 ... Remarks

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

Principles of method if other than guideline

A | X

A

GLP compliance

C ... Remarks N

Test material ^

Test material information

Specific details on test material used for the study

A | X

C

Sampling and analysis ^

Analytical monitoring

A ... Remarks N

Details on sampling

A | X

C

Details on analytical methods

A | X

N

Test solutions ^

Vehicle

C ... Remarks N

Details on test solutions

A | X

C

Test organisms ^

Test organisms (species)

A ... Other A

Details on test organisms

A | X

C

Study design ^

Test type
A ... Other A ...

Water media type
A ... Other A ... Remarks N ...

Limit test
A ...

Total exposure duration
A A ...

Remarks on exposure duration
A ...

Post exposure observation period
C ...

Test conditions ^

Hardness
C ...

Test temperature
C ...

pH
C ...

Dissolved oxygen
C ...

Salinity
C ...

Conductivity
C ...

Nominal and measured concentrations
C ...

Details on test conditions
A X v
C ...

Reference substance (positive control)
A ... Remarks N ...

Any other information on materials and methods incl. tables ^

Normal Aharoni 8 A B I U ≡ ≡ ≡ 1/2 ≡ ≡ ≡ A ≡ ≡

N

Attached full study report

N

+ Add... Open Properties Delete Move up Move down

Illustration (picture/graph)

N

Zoom Load Delete

Attached full study report

N

+ Add... Open Properties Delete Move up Move down

Illustration (picture/graph)

N

Zoom Load Delete

Applicant's summary and conclusion ^

Validity criteria fulfilled

A ... Remarks N ...

Conclusions



C ...

Executive summary


Normal Aharoni 8 A B I U ...

C

9.1.6.2 Long-term toxicity testing on invertebrates (also in Biocidal Product under 9.2.1.6.2)

NB: Biocidal Product  → 

Administrative data ^




Link to relevant study record(s) ^

Study name / type


+ Add...
✕ Delete
↑ Move up
↓ Move down
➤ Go to link target


Description of key information ^




Key value for chemical safety assessment ^


EC10, LC10 or NOEC for freshwater invertebrates




 ...

EC10, LC10 or NOEC for marine water invertebrates



 ...

Additional information ^



Administrative data ^

🗑️
A

Endpoint A Remarks N

Type of information A Other A Remarks N

Adequacy of study A

Robust study summary N

Used for classification N

Used for SDS N

Study period C

Reliability A Other A

Rationale for reliability incl. deficiencies C Other C Remarks N

Data waiving C

Justification for data waiving C

Justification for type of information C

Attached justification	Reason / purpose
N	N

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Reason / purpose	Related information	Remarks
A	A	N

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down
➤ Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	A	A	B	N	N	A	N	A	N

+ Add...
✕ Delete
↑ Move up
↓ Move down
➤ Go to link target

Data access N Other N Remarks N

Data protection claimed N Remarks N

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

Principles of method if other than guideline

A, X

A

GLP compliance

C ... Remarks N

Test material ^

Test material information

Specific details on test material used for the study

A, X

C

Sampling and analysis ^

Analytical monitoring

C ... Remarks N

Details on sampling

A, X

C

Details on analytical methods

A, X

N

Test solutions ^

Vehicle

C ... Remarks N

Details on test solutions

A, X

C

Test organisms ^

Test organisms (species)

A ... Other A

Details on test organisms

A, X

C

Study design ^	
Test type	A ... Other A ...
Water media type	A ... Other A ... Remarks N ...
Limit test	A ...
Total exposure duration	A A ...
Remarks on exposure duration	A ...
Post exposure observation period	C ...
Test conditions ^	
Hardness	C ...
Test temperature	C ...
pH	C ...
Dissolved oxygen	C ...
Salinity	C ...
Conductivity	C ...
Nominal and measured concentrations	C ...
Details on test conditions	A ₂ X ...
Reference substance (positive control)	A ... Remarks N ...

Attached background material

Attached document	Remarks
N	N

+ Add... Edit... Delete Move up Move down

Attached full study report

N

+ Add... Open Properties Delete Move up Move down

Illustration (picture/graph)

N

Zoom Load Delete

Applicant's summary and conclusion ^

Validity criteria fulfilled

A ... Remarks **N** ...

Conclusions

C ...

Executive summary

Normal Aharoni 8 A B I U ...

C

9.1.7 Bioaccumulation in appropriate aquatic species *(also in Biocidal Product under 9.2.1.7)*

NB: Biocidal Product **A** → **C**

Administrative data ^

A

Link to relevant study record(s) ^

Study name / type

+ Add...
✕ Delete
↑ Move up
↓ Move down
➤ Go to link target

Description of key information ^

Normal
Aharoni
8
A
B
I
U
≡
≡
≡
1/2
≡
A
↶

A

Key value for chemical safety assessment ^

BCF (aquatic species)

A
A
...

BMF in fish (dimensionless)

A

Additional information ^

Normal
Aharoni
8
A
B
I
U
≡
≡
≡
1/2
≡
A
↶

C

Administrative data ^

Endpoint
A ... Remarks N ...

Type of information
A ... Other A ... Remarks N ...

Adequacy of study
A ...
 Robust study summary N
 Used for classification N
 Used for SDS N

Study period
C ...

Reliability
A ... Other A ...

Rationale for reliability incl. deficiencies
C ... Other C ... Remarks N ...

Data waiving
C ...

Justification for data waiving
C

Justification for type of information
C

Attached justification

Attached justification	Reason / purpose
N	N

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
A	A	N

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	A	A	B	N	N	A	N	A	N

+ Add... Delete Move up Move down Go to link target

Data access
N ... Other N ... Remarks N ...

Data protection claimed
N ... Remarks N ...

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

Principles of method if other than guideline

A | X

A

GLP compliance

C ... Remarks N

Test material ^

Test material information

Specific details on test material used for the study

A | X

C

Radiolabelling

A ... Remarks N

Sampling and analysis ^

Details on sampling

A | X

C

Details on analytical methods

A | X

N

Test solutions ^

Vehicle

C ... Remarks N

Details on preparation of test solutions, spiked fish food or sediment

A | X

C

Test organisms ^

Test organisms (species)

A ... Other A

Details on test organisms

A | X

C

Study design ^

Route of exposure
 ... Other ... Remarks ...

Justification for method
 ... Other ... Remarks ...

Test type
 ... Other ...

Water / sediment media type
 ... Other ... Remarks ...

Total exposure / uptake duration
 ...

Total depuration duration
 ...

Test conditions ^

Hardness
 ...

Test temperature
 ...

pH
 ...

Dissolved oxygen
 ...

TOC
 ...

Salinity
 ...

Conductivity
 ...

Details on test conditions
 | ...

Nominal and measured concentrations
 ...

Reference substance (positive control)
 ... Remarks ...

Details on estimation of bioconcentration
 | ...

Any other information on materials and methods incl. tables ^

Normal Aharoni 8 A B I U

N

Results and discussion ^

Lipid content

Lipid content	Time point	Remarks on result
A	A	A

+ Add... Edit... Delete Move up Move down

Bioaccumulation factor

Key result	Conc. / dose	Temp.	pH	Type	Value	Basis	Time of plateau	Calculation b...	Remarks on r...
A	<input type="checkbox"/>	A	A	A	A	A	A	A	A

+ Add... Edit... Delete Move up Move down

Depuration

Key result	Elimination	Parameter	Depuration time (DT)	Remarks on result
A	<input type="checkbox"/>	A	A	A

Rate constants

Key result	Rate constant	Value	Remarks on result
A	<input type="checkbox"/>	A	A

+ Add... Edit... Delete Move up Move down

Details on kinetic parameters

A₀ | X **A** ...

Metabolites

C ...

Results with reference substance (positive control)

C ...



Details on results

A₀ | X **A** ...



Reported statistics

C ...

9.1.8 Effects on any other specific, non-target organisms (flora and fauna) believed to be at risk (*also in Biocidal Product under 9.2.1.8*)

NB: Biocidal Product  → 

Administrative data ^


 


Link to relevant study record(s) ^

Study name / type


+ Add...
✕ Delete
↑ Move up
↓ Move down
> Go to link target


Description of key information ^





Additional information ^





Administrative data ^

Endpoint
 ... Remarks

Type of information
 ... Other ... Remarks

Adequacy of study
 ...
 Robust study summary
 Used for classification
 Used for SDS

Study period

Reliability
 ... Other

Rationale for reliability incl. deficiencies
 ... Other ... Remarks

Data waiving

Justification for data waiving

Justification for type of information

Attached justification

Attached justification	Reason / purpose
<input type="text" value="N"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
<input type="text" value="B"/>	<input type="text" value="B"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="B"/>	<input type="text" value="N"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Delete Move up Move down Go to link target

Data access
 ... Other ... Remarks

Data protection claimed
 ... Remarks

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

Principles of method if other than guideline

A | X

A

GLP compliance

C ... Remarks N

Test material ^

Test material information

Specific details on test material used for the study

A | X

C

Sampling and analysis ^

Analytical monitoring

C ... Remarks N

Details on sampling

A | X

C

Details on analytical methods

A | X

N

Test solutions ^

Vehicle

C ... Remarks N

Details on test solutions

A | X

C

Test organisms ^

Aquatic vertebrate type (other than fish)

A ... Other A

Test organisms (species)

A ... Other A

Details on test organisms

A | X

C

Study design ^

Test type
 ... Other ...

Water media type
 ... Other ... Remarks ...

Limit test
 ...

Total exposure duration
 ...

Remarks on exposure duration
 ...

Post exposure observation period
 ...

Test conditions ^

Hardness
 ...

Test temperature
 ...

pH
 ...

Dissolved oxygen
 ...

Salinity
 ...

Conductivity
 ...

Nominal and measured concentrations
 ...

Details on test conditions
 | ...

Reference substance (positive control)
 ... Remarks ...

Overall remarks, attachments ^

Overall remarks

Normal Aharoni 8 A B I U | | | 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 57 58 59 60 61 62 63 64 65 66 67 68 69 70 71 72 73 74 75 76 77 78 79 80 81 82 83 84 85 86 87 88 89 90 91 92 93 94 95 96 97 98 99 100

N

Attached background material

Attached document	Remarks
N	N

+ Add... Edit... X Delete ↑ Move up ↓ Move down

Attached full study report

N

+ Add... Open Properties X Delete ↑ Move up ↓ Move down

Illustration (picture/graph)

N

Zoom Load X Delete

Applicant's summary and conclusion ^

Validity criteria fulfilled

A	Remarks	N
---	---------	---

Conclusions



C

Executive summary


Normal Aharoni 8 A B I U | | | 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 57 58 59 60 61 62 63 64 65 66 67 68 69 70 71 72 73 74 75 76 77 78 79 80 81 82 83 84 85 86 87 88 89 90 91 92 93 94 95 96 97 98 99 100

C

9.1.9 Studies on sediment dwelling organisms (also in Biocidal Product under 9.2.1.9)

NB: Biocidal Product  → 

Administrative data ^




Link to relevant study record(s) ^

Study name / type


+ Add...
✕ Delete
↑ Move up
↓ Move down
> Go to link target


Description of key information ^




Key value for chemical safety assessment ^


EC50 or LC50 for freshwater sediment







EC50 or LC50 for marine water sediment







EC10, LC10 or NOEC for freshwater sediment






EC10, LC10 or NOEC for marine water sediment





Additional information ^



Administrative data ^

Endpoint
A ... Remarks N ...

Type of information
A ... Other A ... Remarks N ...

Adequacy of study
A ...
 Robust study summary N
 Used for classification N
 Used for SDS N

Study period
C ...

Reliability
A ... Other A ...

Rationale for reliability incl. deficiencies
C ... Other C ... Remarks N ...

Data waiving
C ...

Justification for data waiving
C

Justification for type of information
C

Attached justification

Attached justification	Reason / purpose
N	N

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
A	A	N

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	A	A	B	N	N	A	N	A	N

+ Add... Delete Move up Move down Go to link target

Data access
N ... Other N ... Remarks N ...

Data protection claimed
N ... Remarks N ...

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

Principles of method if other than guideline

A | X

A

GLP compliance

C ... Remarks N

Test material ^

Test material information

Specific details on test material used for the study

A | X

C

Sampling and analysis ^

Analytical monitoring

C ... Remarks N

Details on sampling

A | X

C

Details on analytical methods

A | X

N

Test substrate ^

Vehicle

C ... Remarks N

Details on sediment and application

A | X

C

Test organisms ^

Test organisms (species)

A ... Other A

Details on test organisms

A | X

C

Study design ^

Study type
 ... Other ... Remarks ...

Test type
 ... Other ...

Water media type
 ... Other ... Remarks ...

Type of sediment
 ... Remarks ...

Limit test
 ...

Exposure duration

Duration	Exposure phase	Remarks
<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="A"/>

Post exposure observation period
 ...

Test conditions ^

Hardness
 ...

Test temperature
 ...

pH
 ...

Dissolved oxygen
 ...

Salinity
 ...

Ammonia
 ...

Conductivity
 ...

Nominal and measured concentrations
 ...

Details on test conditions
 ...

Reference substance (positive control)
 ... Remarks ...

Attached background material

Attached document	Remarks
N	N

+ Add... Edit... Delete Move up Move down

Attached full study report

N

+ Add... Open Properties Delete Move up Move down

Illustration (picture/graph)

N

Zoom Load Delete

Applicant's summary and conclusion ^

Validity criteria fulfilled

A	...	Remarks	N	...
----------	-----	---------	----------	-----

Conclusions

C ...

Executive summary

Normal Aharoni 8 A B I U | | | 1 2 3 4 5 6 7 8 9 10 11 12 | A |

C

9.1.10 Effects on aquatic macrophytes (also in Biocidal Product under 9.2.1.10)

NB: Biocidal Product A → C

Administrative data ^

A

Link to relevant study record(s) ^

Study name / type

+ Add...
✕ Delete
↑ Move up
↓ Move down
➤ Go to link target

Description of key information ^

A

Key value for chemical safety assessment ^

EC50/LC50 for freshwater plants

A

A ...

EC50/LC50 for marine water plants

A

A ...

EC10, LC10 or NOEC for freshwater plants

A

A ...

EC10, LC10 or NOEC for marine water plants

A

A ...

Additional information ^

C

Administrative data ^

Endpoint
 ... Remarks

Type of information
 ... Other ... Remarks

Adequacy of study
 ...
 Robust study summary
 Used for classification
 Used for SDS

Study period

Reliability
 ... Other

Rationale for reliability incl. deficiencies
 ... Other ... Remarks

Data waiving

Justification for data waiving

Justification for type of information

Attached justification

Attached justification	Reason / purpose
<input type="text" value="N"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
<input type="text" value="B"/>	<input type="text" value="B"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="B"/>	<input type="text" value="N"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Delete Move up Move down Go to link target

Data access
 ... Other ... Remarks

Data protection claimed
 ... Remarks

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

Principles of method if other than guideline

A | X

A

GLP compliance

C ... Remarks N

Test material ^

Test material information

Specific details on test material used for the study

A | X

C

Sampling and analysis ^

Analytical monitoring

C ... Remarks N

Details on sampling

A | X

C

Details on analytical methods

A | X

N

Test solutions ^

Vehicle

C ... Remarks N

Details on test solutions

A | X

C

Test organisms ^

Test organisms (species)

A ... Other A

Details on test organisms

A | X

C

Results and discussion ^

Effect concentrations

Key result	Duration	Dose descriptor	Effect conc.	Nominal / measured	Conc. based on	Basis for effect	Remarks on result
A <input type="checkbox"/>	A	A	A	A	A	A	A

+ Add... Edit... Delete Move up Move down

Details on results

A X v

A ...

Results with reference substance (positive control)

A X v

C ...

Reported statistics and error estimates

C ...

Any other information on results incl. tables ^

A

Overall remarks, attachments ^

Overall remarks

[Rich text editor toolbar with icons for copy, paste, undo, redo, bold, italic, underline, list, link, unlink, etc.]

N

Attached background material

Attached document	Remarks
N	N

+ Add... Edit... X Delete ↑ Move up ↓ Move down

Attached full study report

N

+ Add... Open Properties X Delete ↑ Move up ↓ Move down

Illustration (picture/graph)

N

Zoom Load X Delete

Applicant's summary and conclusion ^

Validity criteria fulfilled

A	Remarks	N
---	---------	---

Conclusions

C

Executive summary

[Rich text editor toolbar with icons for copy, paste, undo, redo, bold, italic, underline, list, link, unlink, etc.]

C

Administrative data ^

Endpoint
 ... Remarks

Type of information
 ... Other ... Remarks

Adequacy of study
 ...
 Robust study summary
 Used for classification
 Used for SDS

Study period

Reliability
 ... Other

Rationale for reliability incl. deficiencies
 ... Other ... Remarks

Data waiving

Justification for data waiving

Justification for type of information

Attached justification

Attached justification	Reason / purpose
<input type="text" value="N"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
<input type="text" value="B"/>	<input type="text" value="B"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="B"/>	<input type="text" value="N"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Delete Move up Move down Go to link target

Data access
 ... Other ... Remarks

Data protection claimed
 ... Remarks

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

Principles of method if other than guideline

A, X

A

GLP compliance

C ... Remarks N

Test material ^

Test material information

Specific details on test material used for the study

A, X

C

Sampling and analysis ^

Analytical monitoring

C ... Remarks N

Details on sampling

A, X

C

Details on analytical methods

A, X

N

Test substrate ^

Vehicle

C ... Remarks N

Details on preparation and application of test substrate

A, X

C

Test organisms ^

Test organisms (inoculum)

A ... Other A

Study design ^

Total exposure duration

A A ...

Remarks

A

Any other information on results incl. tables ^

Normal Aharoni 8 A **B** *I* U ≡ ≡ ≡ 1/2 ≡ A ↺

A

Overall remarks, attachments ^

Overall remarks

Normal Aharoni 8 A **B** *I* U ≡ ≡ ≡ 1/2 ≡ A ↺

N

Attached background material

Attached document	Remarks
N	N

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Attached full study report

N

+ Add...
📄 Open
✎ Properties
✕ Delete
↑ Move up
↓ Move down

Illustration (picture/graph)

N

🔍 Zoom
+ Load
✕ Delete

Applicant's summary and conclusion ^

Validity criteria fulfilled

A

...

N

Remarks

N

...

Conclusions

C

Executive summary

Normal Aharoni 8 A **B** *I* U ≡ ≡ ≡ 1/2 ≡ A ↺

C

Administrative data ^

Endpoint
A ... Remarks N ...

Type of information
A ... Other A ... Remarks N ...

Adequacy of study
A ...
 Robust study summary N
 Used for classification N
 Used for SDS N

Study period
C ...

Reliability
A ... Other A ...

Rationale for reliability incl. deficiencies
C ... Other C ... Remarks N ...

Data waiving
C ...

Justification for data waiving
C

Justification for type of information
C

Attached justification

Attached justification	Reason / purpose
N	N

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
A	A	N

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	A	A	B	N	N	A	N	A	N

+ Add... Delete Move up Move down Go to link target

Data access
N ... Other N ... Remarks N ...

Data protection claimed
N ... Remarks N ...

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Principles of method if other than guideline

A ✕

A

GLP compliance

C ... Remarks N

Test material ^

Test material information

> 🔗 ✕

Specific details on test material used for the study

A ✕

C

Sampling and analysis ^

Analytical monitoring

C ... Remarks N

Details on sampling

A ✕

C

Details on analytical methods

A ✕

N

Test substrate ^

Vehicle

C ... Remarks N

Details on preparation and application of test substrate

A ✕

C

Test organisms ^

Test organisms (species)

A ... Other A

Animal group

A ... Other A

Details on test organisms

A ✕

C

Study design ^

Study type
 ... Other Remarks ...

Substrate type
 ... Other Remarks ...

Limit test
 ...

Total exposure duration
 ...

Remarks
 ...

Post exposure observation period
 ...

Test conditions ^

Test temperature
 ...

pH
 ...

Moisture
 ...

Details on test conditions
 | ...

Nominal and measured concentrations
 ...

Reference substance (positive control)
 ... Remarks ...

Any other information on materials and methods incl. tables ^

B *I* U |

Applicant's summary and conclusion ^

Validity criteria fulfilled

A ... Remarks **N** ...

Conclusions

C ...

Executive summary

Normal Aharoni 8 A B I U ...

C

Administrative data ^

Endpoint
 ... Remarks

Type of information
 ... Other ... Remarks

Adequacy of study
 ...
 Robust study summary
 Used for classification
 Used for SDS

Study period

Reliability
 ... Other

Rationale for reliability incl. deficiencies
 ... Other ... Remarks

Data waiving

Justification for data waiving

Justification for type of information

Attached justification

Attached justification	Reason / purpose
<input type="text" value="N"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
<input type="text" value="B"/>	<input type="text" value="B"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="B"/>	<input type="text" value="N"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Delete Move up Move down Go to link target

Data access
 ... Other ... Remarks

Data protection claimed
 ... Remarks

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

+ Add... Edit... Delete Move up Move down

Principles of method if other than guideline

A X

A

GLP compliance

C ... Remarks N

Test material ^

Test material information

Specific details on test material used for the study

A X

C

Sampling and analysis ^

Analytical monitoring

C ... Remarks N

Details on sampling

A X

C

Details on analytical methods

A X

N

Test substrate ^

Vehicle

C ... Remarks N

Details on preparation and application of test substrate

A X

C

Test organisms ^

Species

A ... Other A

Plant group

A ... Other A

Details on test organisms

A X

A

Study design ^

Test type
 ... Other ...

Study type
 ... Other ... Remarks ...

Substrate type
 ... Other ... Remarks ...

Limit test
 ...

Total exposure duration
 ...

Remarks
 ...

Post exposure observation period
 ...

Justification for exposure duration
 ...

Test conditions ^

Test temperature
 ...

pH
 ...

Moisture
 ...

Details on test conditions
 ...

Nominal and measured concentrations
 ...

Reference substance (positive control)
 ... Remarks ...

Any other information on materials and methods incl. tables ^

Normal Aharoni 8 A B I U 1/2 1/3 1/4 1/5 1/6 1/7 1/8 1/9 1/10 1/11 1/12 1/13 1/14 1/15 1/16 1/17 1/18 1/19 1/20 1/21 1/22 1/23 1/24 1/25 1/26 1/27 1/28 1/29 1/30 1/31 1/32 1/33 1/34 1/35 1/36 1/37 1/38 1/39 1/40 1/41 1/42 1/43 1/44 1/45 1/46 1/47 1/48 1/49 1/50 1/51 1/52 1/53 1/54 1/55 1/56 1/57 1/58 1/59 1/60 1/61 1/62 1/63 1/64 1/65 1/66 1/67 1/68 1/69 1/70 1/71 1/72 1/73 1/74 1/75 1/76 1/77 1/78 1/79 1/80 1/81 1/82 1/83 1/84 1/85 1/86 1/87 1/88 1/89 1/90 1/91 1/92 1/93 1/94 1/95 1/96 1/97 1/98 1/99 1/100

Applicant's summary and conclusion ^

Validity criteria fulfilled

A ... Remarks **N** ...

Conclusions

C ...

Executive summary

Normal Aharoni 8 A B I U ...

C

9.4 Effects on birds (also in Biocidal Product under 9.2.4)

NB: Biocidal Product **A** → **C**

Administrative data ^

A

Link to relevant study record(s) ^

Study name / type

+ Add... X Delete ↑ Move up ↓ Move down > Go to link target

Description of key information ^

Normal Aharoni 8 A[†] B I U [List icons] [Undo/Redo] [Copy/Paste]

A

Key value for chemical safety assessment ^

Short-term EC50 or LC50 for birds

A **A** ...

Long-term EC10, LC10 or NOEC for birds

A **A** ...

Additional information ^

Normal Aharoni 8 A[†] B I U [List icons] [Undo/Redo] [Copy/Paste]

C

Administrative data ^

Endpoint
A ... **Remarks** N ...

Type of information
A ... **Other** A ... **Remarks** N ...

Adequacy of study
A ...
 Robust study summary N
 Used for classification N
 Used for SDS N

Study period
C ...

Reliability
A ... **Other** A ...

Rationale for reliability incl. deficiencies
C ... **Other** C ... **Remarks** N ...

Data waiving
C ...

Justification for data waiving
C

Justification for type of information
C

Attached justification

Attached justification	Reason / purpose
N	N

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
A	A	N

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	A	A	B	N	N	A	N	A	N

+ Add... Delete Move up Move down Go to link target

Data access
N ... **Other** N ... **Remarks** N ...

Data protection claimed
N ... **Remarks** N ...

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

Principles of method if other than guideline

A, X

A

GLP compliance

C ... Remarks N

Test material ^

Test material information

Specific details on test material used for the study

A, X

C

Dose method

A ... Other A ... Remarks N

Analytical monitoring

C ... Remarks N

Vehicle

C ... Remarks N

Details on preparation and analysis of diet

A, X

C

Test organisms ^

Test organisms (species)

A ... Other A

Details on test organisms

A, X

C

Study design ^

Limit test

A

Total exposure duration (if not bolus)

A A

Remarks

A

Post exposure observation period

C

No. of animals per sex per dose and/or stage

C

Control animals

▼ C

Nominal and measured doses / concentrations

C ...

Details on test conditions

A, X

C ...

Examinations ^

Details on examinations and observations

A, X

C ...

Details on reproductive parameters

A, X

C ...

Reference substance (positive control)

A ... Remarks N ...

Any other information on materials and methods incl. tables ^

Normal Aharoni 8 A B I U ...

N

Results and discussion ^

Effect levels

Key result	Duration (if not bolus)	Dose descriptor	Effect level	Conc. / dose based ...	Basis for effect	Remarks on result
A <input type="checkbox"/>	A	A	A	A	A	A

+ Add... Edit... X Delete ↑ Move up ↓ Move down

Repellency factors (if applicable)

A ...

Mortality and sub-lethal effects

A, X

A ...

Effects on reproduction

A, X

A ...

Results with reference substance (positive control)

A, X

C ...

Further details on results

C

Reported statistics and error estimates

C

Any other information on results incl. tables ^

Normal Aharoni 8 A B I U

A

Overall remarks, attachments ^

Overall remarks

Normal Aharoni 8 A B I U

N

Attached background material

Attached document	Remarks
N	N

+ Add... Edit... X Delete ↑ Move up ↓ Move down

Attached full study report

N

+ Add... Open Properties X Delete ↑ Move up ↓ Move down

Illustration (picture/graph)

N

Q Zoom + Load X Delete

Applicant's summary and conclusion ^

Validity criteria fulfilled

A ... Remarks N ...

Conclusions

C ...

Executive summary

Normal Aharoni 8 A B I U ...

C

Administrative data ^

Endpoint
 ... Remarks

Type of information
 ... Other ... Remarks

Adequacy of study
 ...
 Robust study summary
 Used for classification
 Used for SDS

Study period

Reliability
 ... Other

Rationale for reliability incl. deficiencies
 ... Other ... Remarks

Data waiving

Justification for data waiving

Justification for type of information

Attached justification

Attached justification	Reason / purpose
<input type="text" value="N"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
<input type="text" value="B"/>	<input type="text" value="B"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="B"/>	<input type="text" value="N"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Delete Move up Move down Go to link target

Data access
 ... Other ... Remarks

Data protection claimed
 ... Remarks

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

Principles of method if other than guideline

A | X

A

GLP compliance

C ... Remarks N

Application method

A ... Other A

Test material ^

Test material information

Specific details on test material used for the study

A | X

C

Sampling and analysis ^

Analytical monitoring

C ... Remarks N

Details on sampling

A | X

C

Details on analytical methods

A | X

N

Test substrate ^

Vehicle

C ... Remarks N

Details on preparation and application of test substrate

A | X

C

Test organisms ^

Test organisms (species)

A ... Other A

Animal group

A ... Other A

Details on test organisms

A | X

C

Study design ^

Study type

Limit test
 ...

Total exposure duration
 ...

Remarks
 ...

Post exposure observation period
 ...

Test conditions ^

Test temperature
 ...

pH (if soil or dung study)
 ...

Humidity
 ...

Photoperiod and lighting
 ...

Details on test conditions
 | v
 ...

Nominal and measured concentrations
 ...

Reference substance (positive control)

Any other information on materials and methods incl. tables ^



Rich text editor toolbar with icons for bold, italic, underline, list, link, etc.


Results and discussion ^

Effect concentrations



Key result	Duration	Dose descriptor	Effect conc.	Nominal / measured	Conc. based on	Basis for effect	Remarks on result
<input type="checkbox"/> <input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="A"/>


Details on results

A  


A 


Results with reference substance (positive control)


A  

C 


Reported statistics and error estimates

C 


Any other information on results incl. tables 



A

Overall remarks, attachments 

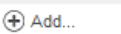
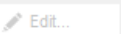

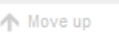
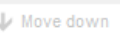
Overall remarks



N

Attached background material

Attached document	Remarks
N	N

Attached full study report

N

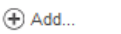





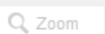


     

Illustration (picture/graph)

N

Applicant's summary and conclusion ^

Validity criteria fulfilled

A ... Remarks **N** ...

Conclusions

C ...

Executive summary

Normal Aharoni 8 A B I U ...

C

Administrative data ^

Endpoint
A ... **Remarks** N ...

Type of information
A ... **Other** A ... **Remarks** N ...

Adequacy of study
A ...
 Robust study summary N
 Used for classification N
 Used for SDS N

Study period
C ...

Reliability
A ... **Other** A ...

Rationale for reliability incl. deficiencies
C ... **Other** C ... **Remarks** N ...

Data waiving
C ...

Justification for data waiving
C

Justification for type of information
C

Attached justification

Attached justification	Reason / purpose
N	N

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
A	A	N

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	A	A	B	N	N	A	N	A	N

+ Add... Delete Move up Move down Go to link target

Data access
N ... **Other** N ... **Remarks** N ...

Data protection claimed
N ... **Remarks** N ...

Overall remarks, attachments ^

Overall remarks

Normal Aharoni 8 A B I U | | | 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 57 58 59 60 61 62 63 64 65 66 67 68 69 70 71 72 73 74 75 76 77 78 79 80 81 82 83 84 85 86 87 88 89 90 91 92 93 94 95 96 97 98 99 100

N

Attached background material

Attached document	Remarks
N	N

+ Add... Edit... X Delete ↑ Move up ↓ Move down

Attached full study report

N

+ Add... Open Properties X Delete ↑ Move up ↓ Move down

Illustration (picture/graph)

N

Zoom Load X Delete

Applicant's summary and conclusion ^

Validity criteria fulfilled

A	Remarks	N
---	---------	---

Conclusions

C

Executive summary

Normal Aharoni 8 A B I U | | | 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 57 58 59 60 61 62 63 64 65 66 67 68 69 70 71 72 73 74 75 76 77 78 79 80 81 82 83 84 85 86 87 88 89 90 91 92 93 94 95 96 97 98 99 100

C

9.8 Effects on other non-target, non-aquatic organisms (*also in Biocidal Product under 9.2.8*)

NB: Biocidal Product A → C

Administrative data ^

A

Link to relevant study record(s) ^

Study name / type

+ Add...
✕ Delete
↑ Move up
↓ Move down
> Go to link target

Description of key information ^

A

Key value for chemical safety assessment ^

Short-term EC50 or LC50 for mammals

A

A ...

Long-term EC10, LC10 or NOEC for mammals

A

A ...

Additional information ^

C

Administrative data ^

Endpoint
 ... Remarks

Type of information
 ... Other ... Remarks

Adequacy of study
 ...
 Robust study summary
 Used for classification
 Used for SDS

Study period

Reliability
 ... Other

Rationale for reliability incl. deficiencies
 ... Other ... Remarks

Data waiving

Justification for data waiving

Justification for type of information

Attached justification

Attached justification	Reason / purpose
<input type="text" value="N"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
<input type="text" value="B"/>	<input type="text" value="B"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="B"/>	<input type="text" value="N"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Delete Move up Move down Go to link target

Data access
 ... Other ... Remarks

Data protection claimed
 ... Remarks

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

Principles of method if other than guideline

A | X

A

GLP compliance

C ... Remarks N

Test material ^

Test material information

Specific details on test material used for the study

A | X

C

Sampling and analysis ^

Analytical monitoring

C ... Remarks N

Details on sampling

A | X

C

Details on analytical methods

A | X

N

Test substrate ^

Vehicle

C ... Remarks N

Details on preparation and application of test substrate

A | X

C

Test organisms ^

Test organisms (species)

A ... Other A

Details on test organisms

A | X

C

Study design ^

Study type
A ... Other A ... Remarks N ...

Limit test
A ...

Total exposure duration
A A ...

Remarks
A ...

Post exposure observation period
C ...

Test conditions ^

Test temperature
C ...

Humidity
C ...

Photoperiod and lighting
C ...

Details on test conditions
A, X
C ...

Nominal and measured concentrations
C ...

Reference substance (positive control)
A ... Remarks N ...

Any other information on materials and methods incl. tables ^

Normal Aharoni 8 A B I U | | | 1/2 = | A |

N

Illustration (picture/graph)

N

Zoom Load Delete

Applicant's summary and conclusion ^

Validity criteria fulfilled

A Remarks **N**

Conclusions

C

Executive summary

Normal Aharoni 8 A B I U | | | 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 57 58 59 60 61 62 63 64 65 66 67 68 69 70 71 72 73 74 75 76 77 78 79 80 81 82 83 84 85 86 87 88 89 90 91 92 93 94 95 96 97 98 99 100

C

Administrative data ^

Endpoint
 ... Remarks

Type of information
 ... Other ... Remarks

Adequacy of study
 ...
 Robust study summary
 Used for classification
 Used for SDS

Study period

Reliability
 ... Other

Rationale for reliability incl. deficiencies
 ... Other ... Remarks

Data waiving

Justification for data waiving

Justification for type of information

Attached justification

Attached justification	Reason / purpose
<input type="text" value="N"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
<input type="text" value="B"/>	<input type="text" value="B"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="B"/>	<input type="text" value="N"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Delete Move up Move down Go to link target

Data access
 ... Other ... Remarks

Data protection claimed
 ... Remarks



10.1 Fate and behaviour in water and sediment

10.1.1 Degradation, initial studies

10.1.1.1 Abiotic






10.1.1.1.a Abiotic: Hydrolysis as a function of pH and identification of breakdown products

Administrative data ^

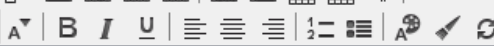
 


Link to relevant study record(s) ^

Study name / type

 Add...  Delete  Move up  Move down  Go to link target



Description of key information ^

Normal Aharoni 8 





Key value for chemical safety assessment ^


Half-life for hydrolysis


  ...

at the temperature of

  ...

Additional information ^

Normal Aharoni 8 



Administrative data ^

Endpoint
A ... **Remarks** N ...

Type of information
A ... **Other** A ... **Remarks** N ...

Adequacy of study
A ...
 Robust study summary N
 Used for classification N
 Used for SDS N

Study period
C ...

Reliability
A ... **Other** A ...

Rationale for reliability incl. deficiencies
C ... **Other** C ... **Remarks** N ...

Data waiving
C ...

Justification for data waiving
C

Justification for type of information
C

Attached justification

Attached justification	Reason / purpose
N	N

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
A	A	N

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	A	A	B	N	N	A	N	A	N

+ Add... Delete Move up Move down Go to link target

Data access
N ... **Other** N ... **Remarks** N ...

Data protection claimed
N ... **Remarks** N ...

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

Principles of method if other than guideline

A | X

A

GLP compliance

C ... Remarks N

Test material ^

Test material information

Specific details on test material used for the study

A | X

C

Radiolabelling

A ... Remarks N

Study design ^

Analytical monitoring

C ... Remarks N

Details on sampling

A | X

C

Details on analytical methods

A | X

N

Buffers

A | X

C

Estimation method (if used)

C

Details on test conditions

A | X

C

Other kinetic parameters

A

Details on results

A

Results with reference substance

C

Any other information on results incl. tables ^

Normal Aharoni 8 A B I U

A

Overall remarks, attachments ^

Overall remarks

Normal Aharoni 8 A B I U

N

Attached background material

Attached document	Remarks
N	N

+ Add... Edit... Delete Move up Move down

Attached full study report

N

+ Add... Open Properties Delete Move up Move down

Illustration (picture/graph)

N

Zoom Load Delete

Applicant's summary and conclusion ^

Validity criteria fulfilled

A ... Remarks **N** ...

Conclusions

C ...

Executive summary

Normal Aharoni 8 A B I U ...

C

Administrative data ^

Endpoint
A ... **Remarks** N ...

Type of information
A ... **Other** A ... **Remarks** N ...

Adequacy of study
A ...
 Robust study summary N
 Used for classification N
 Used for SDS N

Study period
C ...

Reliability
A ... **Other** A ...

Rationale for reliability incl. deficiencies
C ... **Other** C ... **Remarks** N ...

Data waiving
C ...

Justification for data waiving
C

Justification for type of information
C

Attached justification

Attached justification	Reason / purpose
N	N

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
A	A	N

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	A	A	B	N	N	A	N	A	N

+ Add... Delete Move up Move down Go to link target

Data access
N ... **Other** N ... **Remarks** N ...

Data protection claimed
N ... **Remarks** N ...

Materials and methods ^

Study type
 ... Other ... Remarks ...

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="A"/>

+ Add... Edit... Delete Move up Move down

Principles of method if other than guideline
 ...

GLP compliance
 ... Remarks ...

Test material ^

Test material information
 > 🔗 ✕

Specific details on test material used for the study
 ...

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="A"/>

+ Add... Edit... Delete Move up Move down

Principles of method if other than guideline
 ...

GLP compliance
 ... Remarks ...

Study design ^

Radiolabelling
 ... Remarks ...

Analytical method

Details on sampling
 ...

Details on analytical methods
 ...

Buffers

A₂ | X

C
...

Light source

A ... Other A ... Remarks N ...

Light spectrum: wavelength in nm

A A A A

Relative light intensity

A A A A

Details on light source

A₂ | X

C
...

Sensitiser (for indirect photolysis)

Type of sensitiser	Details on sensitiser	Concentration of sensitiser
A	A	A

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Details on test conditions

A₂ | X

C
...

Duration of test at given test condition

Duration	Temp.	Initial conc. measured
C	C	C

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Reference substance

C ... Remarks N ...

Dark controls

C ... Remarks N ...

Computational methods

A₂ | X

C
...

Any other information on materials and methods incl. tables ^

N

Results and discussion ^

Preliminary study

C ...

Test performance

C ...

Spectrum of substance

Parameter	Value	Remarks
A	A	A

+ Add... Edit... Delete Move up Move down

% Degradation

Key result	% Degr.	St. dev.	Sampling time	Test condition	Remarks on result
A <input type="checkbox"/>	A	A	A	A	A

+ Add... Edit... Delete Move up Move down

Quantum yield (for direct photolysis)

A

Rate constant (for indirect photolysis)

A A A A ... Other A ...

Dissipation half-life of parent compound

Key result	DT50	Test condition	Remarks on result
A <input type="checkbox"/>	A	A	A

+ Add... Edit... Delete Move up Move down

Predicted environmental photolytic half-life

C ...

Transformation products

C ... Remarks C ...

Identity of transformation products

No.	Reference substance
C	C

+ Add... Edit... Delete Move up Move down Go to link target

Details on results

A X



A ...

Results with reference substance

C ...






10.1.1.2 Biotic → Ready / Inherent biodegradability

Administrative data ^


 


Link to relevant study record(s) ^

Study name / type

 Add...  Delete  Move up  Move down  Go to link target


Description of key information ^







Key value for chemical safety assessment ^

Biodegradation in water

 ...

Additional information ^





Administrative data ^

Endpoint
A ... **Remarks** N ...

Type of information
A ... **Other** A ... **Remarks** N ...

Adequacy of study
A ...
 Robust study summary N
 Used for classification N
 Used for SDS N

Study period
C ...

Reliability
A ... **Other** A ...

Rationale for reliability incl. deficiencies
C ... **Other** C ... **Remarks** N ...

Data waiving
C ...

Justification for data waiving
C

Justification for type of information
C

Attached justification

Attached justification	Reason / purpose
N	N

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
A	A	N

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	A	A	B	N	N	A	N	A	N

+ Add... Delete Move up Move down Go to link target

Data access
N ... **Other** N ... **Remarks** N ...

Data protection claimed
N ... **Remarks** N ...

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

Principles of method if other than guideline

A, X

A

GLP compliance

C ... Remarks N

Test material ^

Test material information

Specific details on test material used for the study

A, X

C

Study design ^

Oxygen conditions

A ... Other A ... Remarks N

Inoculum or test system

A ... Other A ... Remarks N

Details on inoculum

A, X

C

Duration of test (contact time)

A A A A A

Initial test substance concentration

Initial conc.	Based on
A	A

Parameter followed for biodegradation estimation

Parameter followed for biodegradation estimation

C

Details on analytical methods
A | X

N ...

Details on study design
A | X

C ...

Reference substance

Reference substance

C

+ Add... Edit... Delete Move up Move down

Any other information on materials and methods incl. tables ^

Normal Aharoni 8 A B I U

N

Results and discussion ^

Preliminary study

C ...

Test performance

C ...

% Degradation

Key result	Parameter	Value	St. dev.	Sampling time	Remarks on result
A <input type="checkbox"/>	A	A	A	A	A

+ Add... Edit... Delete Move up Move down

Details on results

A ...

BOD5 / COD results ^

BOD5 / COD

Key result	Parameter	Value	Remarks on result
A <input type="checkbox"/>	A	A	A

+ Add... Edit... Delete Move up Move down

Results with reference substance

C ...

Any other information on results incl. tables ^

Normal Aharoni 8 A B I U

A

Overall remarks, attachments ^

Overall remarks

Normal Aharoni 8 A B I U

N

Attached background material

Attached document	Remarks
N	N

+ Add... Edit... Delete Move up Move down

Attached full study report

N

+ Add... Open Properties Delete Move up Move down

Illustration (picture/graph)

N

Zoom Load Delete

Applicant's summary and conclusion ^

Validity criteria fulfilled

A Remarks **N**

Interpretation of results

C Other **C** Remarks **N**

Conclusions

C


Executive summary

Normal Aharoni 8 A B I U

C


10.1.2 Adsorption / desorption

Administrative data ^




Link to relevant study record(s) ^


Study name / type




Description of key information ^






Key value for chemical safety assessment ^



Koc at 20 °C 

Log Koc at 20 °C 


Other adsorption coefficients

Type  ... Other 

Value in L/kg 

at the temperature of  

Additional information ^



Administrative data ^

Endpoint
A ... Remarks N ...

Type of information
A ... Other A ... Remarks N ...

Adequacy of study
A ...
 Robust study summary N
 Used for classification N
 Used for SDS N

Study period
C ...

Reliability
A ... Other A ...

Rationale for reliability incl. deficiencies
C ... Other C ... Remarks N ...

Data waiving
C ...

Justification for data waiving
C

Justification for type of information
C

Attached justification

Attached justification	Reason / purpose
N	N

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
A	A	N

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	A	A	B	N	N	A	N	A	N

+ Add... Delete Move up Move down Go to link target

Data access
N ... Other N ... Remarks N ...

Data protection claimed
N ... Remarks N ...

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

⊕ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Principles of method if other than guideline

A ✕

A

GLP compliance

C ... Remarks N

Type of method

A ... Other A ... Remarks N

Media

A ... Other A

Test material ^

Test material information

> ∞

Specific details on test material used for the study

A ✕

C

Radiolabelling

A ... Remarks N

Study design ^

Test temperature

C

HPLC method ^

Details on study design: HPLC method

A ✕

C

Batch equilibrium or other method ^

Analytical monitoring

C ... Remarks N

Details on sampling

A ✕

C

Details on analytical methods

A ✕

N

Matrix properties

Matrix no.	Matrix type	% Clay	% Silt	% Sand	% Org. carbon	pH	CEC	Bulk density
C	C	C	C	C	C	C	C	C

Details on matrix

A | X

C

Details on test conditions

A | X

C

Duration of adsorption equilibration

Sample No.	Duration	Initial conc. measured	pH	Temp.	Remarks
A	A	A	A	A	A

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Duration of desorption equilibration

Sample no.	Duration	Conc. of adsorbed test...	pH	Temp.	Remarks
A	A	A	A	A	A

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Computational methods

A | X

C

Any other information on materials and methods incl. tables ^

N

Results and discussion ^

Adsorption coefficient

Key result	Sample No.	Type	Value	pH	Temp.	Matrix	% Org. carbon	Remarks on
A <input type="checkbox"/>	A	A	A	A	A	A	A	A

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Partition coefficients

Key result	Sample No.	Phase system	Type	Value	Temp.	pH	Matrix	% Org. carbon	Remarks on
A <input type="checkbox"/>	A	A	A	A	A	A	A	A	A

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Results: HPLC method ^

Details on results (HPLC method)

A | X

A

Results: Batch equilibrium or other method ^

Adsorption and desorption constants

C

Recovery of test material

C

Concentration of test substance at end of adsorption equilibration period

C

Concentration of test substance at end of desorption equilibration period

C

Mass balance (%) at end of adsorption phase

Sample no.	Duration	% Adsorption	Remarks on result
C	C	C	C

+ Add... | Edit... | Delete | Move up | Move down

Mass balance (%) at end of desorption phase

Sample no.	Duration	% Desorption	Remarks on result
C	C	C	C

+ Add... | Edit... | Delete | Move up | Move down

Transformation products

C ... Remarks C

Identity of transformation products

No.	Reference substance
C	C

+ Add... | Edit... | Delete | Move up | Move down | Go to link targ

Details on results (Batch equilibrium method)

A | X

A

Statistics

C

Any other information on results incl. tables ^

A

Overall remarks, attachments ^

Overall remarks

Normal Aharoni 8 A B I U | | | 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 57 58 59 60 61 62 63 64 65 66 67 68 69 70 71 72 73 74 75 76 77 78 79 80 81 82 83 84 85 86 87 88 89 90 91 92 93 94 95 96 97 98 99 100

N

Attached background material

Attached document	Remarks
N	N

+ Add... Edit... X Delete ↑ Move up ↓ Move down

Attached full study report

N

+ Add... Open Properties X Delete ↑ Move up ↓ Move down

Illustration (picture/graph)

N

Zoom + Load X Delete

Applicant's summary and conclusion ^

Validity criteria fulfilled

A ... Remarks N

Conclusions

C

Executive summary

Normal Aharoni 8 A B I U | | | 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 57 58 59 60 61 62 63 64 65 66 67 68 69 70 71 72 73 74 75 76 77 78 79 80 81 82 83 84 85 86 87 88 89 90 91 92 93 94 95 96 97 98 99 100

C

Administrative data ^

Endpoint
A ... Remarks N ...

Type of information
A ... Other A ... Remarks N ...

Adequacy of study
A ...
 Robust study summary N
 Used for classification N
 Used for SDS N

Study period
C ...

Reliability
A ... Other A ...

Rationale for reliability incl. deficiencies
C ... Other C ... Remarks N ...

Data waiving
C ...

Justification for data waiving
C

Justification for type of information
C

Attached justification

Attached justification	Reason / purpose
N	N

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
A	A	N

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	A	A	B	N	N	A	N	A	N

+ Add... Delete Move up Move down Go to link target

Data access
N ... Other N ... Remarks N ...

Data protection claimed
N ... Remarks N ...

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

Principles of method if other than guideline

A, X

A

GLP compliance

C ... Remarks N

Test material ^

Test material information

Specific details on test material used for the study

A, X

C

Radiolabelling

A ... Remarks N

Study design ^

Oxygen conditions

A ... Other A ... Remarks N

Inoculum or test system

A ... Other A ... Remarks N

Details on source and properties of surface water

A, X

C

Details on source and properties of sediment

A, X

C

Details on inoculum

A, X

C

Duration of test (contact time)

A A A A A

Initial test substance concentration

Initial conc. A	Based on A
--	---

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Parameter followed for biodegradation estimation

▼ C

Details on analytical methods

A₁ | ✕
▼

N
...

Details on study design

A₁ | ✕
▼

C
...

Reference substance

Reference substance

C

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Any other information on materials and methods incl. tables ^

N

Results and discussion ^

Test performance

C
...

Mean total recovery

Compartment	% Recovery	St. dev.	Remarks on result
A	A	A	A

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

% Degradation

Key result	% Degr.	St. dev.	Parameter	Sampling time	Remarks on result
A	A	A	A	A	A

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Overall remarks, attachments ^

Overall remarks

Normal Aharoni 8 A B I U | | | 1 2 | A ↻

N

Attached background material

Attached document	Remarks
N	N

+ Add... Edit... X Delete ↑ Move up ↓ Move down

Attached full study report

N

+ Add... Open Properties X Delete ↑ Move up ↓ Move down

Illustration (picture/graph)

N

Zoom + Load X Delete

Applicant's summary and conclusion ^

Validity criteria fulfilled

A ... Remarks N

Conclusions

C



Executive summary

Normal Aharoni 8 A B I U | | | 1 2 | A ↻

C






10.1.5 Field studies

Administrative data ^


 


Link to relevant study record(s) ^

Study name / type


 Add...  Delete  Move up  Move down  Go to link target


Description of key information ^





Additional information ^





Administrative data ^

Endpoint
A ... Remarks N ...

Type of information
A ... Other A ... Remarks N ...

Adequacy of study
A ...
 Robust study summary N
 Used for classification N
 Used for SDS N

Study period
C ...

Reliability
A ... Other A ...

Rationale for reliability incl. deficiencies
C ... Other C ... Remarks N ...

Data waiving
C ...

Justification for data waiving
C

Justification for type of information
C

Attached justification

Attached justification	Reason / purpose
N	N

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
A	A	N

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	A	A	B	N	N	A	N	A	N

+ Add... Delete Move up Move down Go to link target



Data access
N ... Other N ... Remarks N ...

Data protection claimed
N ... Remarks N ...

10.2 Fate and behaviour in soil






10.2.1 Laboratory study on rate and route of degradation including identification of the processes involved

Administrative data ^






















 

Link to relevant study record(s) ^

Study name / type

 Add...  Delete  Move up  Move down  Go to link target

Description of key information ^

Normal Aharoni 8 A B I U                     

Administrative data ^

Endpoint
A ... Remarks N ...

Type of information
A ... Other A ... Remarks N ...

Adequacy of study
A ...
 Robust study summary N
 Used for classification N
 Used for SDS N

Study period
C ...

Reliability
A ... Other A ...

Rationale for reliability incl. deficiencies
C ... Other C ... Remarks N ...

Data waiving
C ...

Justification for data waiving
C

Justification for type of information
C

Attached justification

Attached justification	Reason / purpose
N	N

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
A	A	N

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	A	A	B	N	N	A	N	A	N

+ Add... Delete Move up Move down Go to link target

Data access
N ... Other N ... Remarks N ...

Data protection claimed
N ... Remarks N ...

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

Principles of method if other than guideline

A, X

A

GLP compliance

C ... Remarks N

Test type

A ... Other A

Test material ^

Test material information

Specific details on test material used for the study

A, X

C

Radiolabelling

A ... Remarks N

Study design ^

Oxygen conditions

A ... Other A ... Remarks N

Soil classification

A ... Other A ... Remarks N

Year

A

Soil properties

Soil no.	Soil type	% Clay	% Silt	% Sand	% Org. C	pH	CEC	Bulk density (g...
A	A	A	A	A	A	A	A	A

Details on soil characteristics

A, X

C

Duration of test (contact time)

Soil No.	Duration
A	A

Initial test substance concentration

Soil No.	Initial conc.	Based on
A	A	A

Details on transformation products

C
...

Evaporation of parent compound

C
...
Remarks
C
...

Volatile metabolites

C
...
Remarks
C
...

Residues

C
...
Remarks
C
...

Details on results

A
...

Results with reference substance

C
...

Any other information on results incl. tables ^

A

Overall remarks, attachments ^

Overall remarks

N

Attached background material

Attached document	Remarks
N	N

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Attached full study report

N

+ Add...
📄 Open
🔗 Properties
✕ Delete
↑ Move up
↓ Move down

Illustration (picture/graph)

N

🔍 Zoom
+ Load
✕ Delete

Applicant's summary and conclusion ^

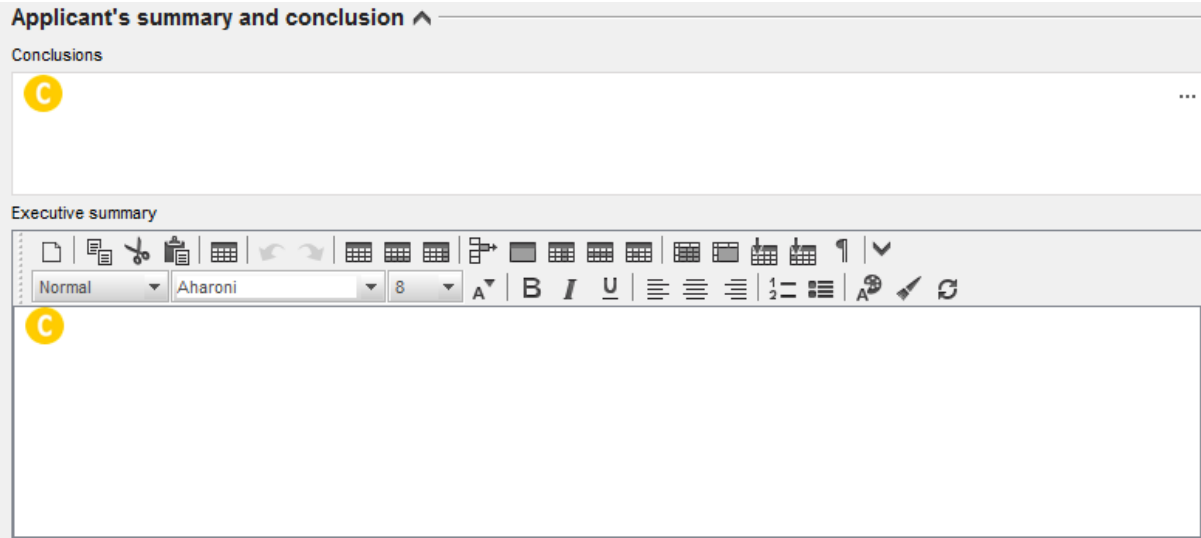
Conclusions

C

Executive summary

Normal Aharoni 8 A B I U



C

The image shows a screenshot of the IUCLID software interface. At the top, there is a header bar with the text "Applicant's summary and conclusion" followed by a small upward-pointing arrow. Below this, there are two main text input areas. The first is labeled "Conclusions" and contains a yellow circle with the letter "C" in the top-left corner and a three-dot menu icon in the top-right corner. The second is labeled "Executive summary" and also contains a yellow circle with the letter "C" in the top-left corner. Between these two text areas is a rich text editor toolbar. The toolbar includes various icons for text editing (copy, paste, undo, redo), alignment (left, center, right, justified), and formatting (bold, italic, underline, list, link, unlink, indent, outdent). Below the toolbar, the "Executive summary" text area is visible, with the yellow circle "C" in the top-left corner.

10.2.3 Soil accumulation studies (also in Biocidal Product under 10.1 Foreseeable routes of entry into the environment on the basis of the use envisaged)

NB: Biocidal Product  → 

Administrative data ^


 


Link to relevant study record(s) ^

Study name / type


+ Add...
✕ Delete
↑ Move up
↓ Move down
➤ Go to link target


Description of key information ^





Additional information ^





Administrative data ^

Endpoint
 ... Remarks

Type of information
 ... Other ... Remarks

Adequacy of study
 ...
 Robust study summary
 Used for classification
 Used for SDS

Study period

Reliability
 ... Other

Rationale for reliability incl. deficiencies
 ... Other ... Remarks

Data waiving

Justification for data waiving

Justification for type of information

Attached justification

Attached justification	Reason / purpose
<input type="text" value="N"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
<input type="text" value="B"/>	<input type="text" value="B"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="B"/>	<input type="text" value="N"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Delete Move up Move down Go to link target

Data access
 ... Other ... Remarks

Data protection claimed
 ... Remarks

Materials and methods ^

Model
 ... Other ...

Calculation programme
 ...

Release year

Media
 ... Other ...

Test material ^

Test material information


Specific details on test material used for the study
 ...

Study design ^

Test substance input data
 ...

Environmental properties
 ...

Any other information on materials and methods incl. tables ^



Results and discussion ^

Percent distribution in media ^

Air (%)

Water (%)

Soil (%)

Sediment (%)


Susp. sediment (%)

Biota (%)

Aerosol (%)

Other distribution results
 ...

Any other information on results incl. tables ^



Administrative data ^

Endpoint
 ... Remarks

Type of information
 ... Other ... Remarks

Adequacy of study
 ...
 Robust study summary
 Used for classification
 Used for SDS

Study period

Reliability
 ... Other

Rationale for reliability incl. deficiencies
 ... Other ... Remarks

Data waiving

Justification for data waiving

Justification for type of information

Attached justification

Attached justification	Reason / purpose
<input type="text" value="N"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
<input type="text" value="B"/>	<input type="text" value="B"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="B"/>	<input type="text" value="N"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Delete Move up Move down Go to link target

Data access
 ... Other ... Remarks

Data protection claimed
 ... Remarks

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Principles of method if other than guideline

A, ✕

A

⋮

GLP compliance

C
Remarks N

Test material ^

Test material information

> ∞ ✕

Specific details on test material used for the study

A, ✕

C

⋮

Study design ^

Estimation method (if used)

A, ✕

C

⋮

Light source

A
Other A Remarks N

Light spectrum: wavelength in nm

A A A A

Relative light intensity

A A A A

Details on light source

A, ✕

C

⋮

Details on test conditions

C

⋮

Duration of test at given test condition

Duration	Temp.	Initial conc. measured
C	C	C

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Reference substance

C
Remarks N

Any other information on results incl. tables ^

Normal Aharoni 8 A B I U ...

A

Overall remarks, attachments ^

Overall remarks

Normal Aharoni 8 A B I U ...

N

Attached background material

Attached document	Remarks
N	N

+ Add... Edit... X Delete ↑ Move up ↓ Move down

Attached full study report

N

+ Add... Open Properties X Delete ↑ Move up ↓ Move down

Illustration (picture/graph)

N

Zoom Load X Delete

Applicant's summary and conclusion ^

Validity criteria fulfilled

A	Remarks	N
---	---------	---

Conclusions

C

Executive summary

Normal Aharoni 8 A B I U ...

C

10.3 Biocidal Product – Leaching behaviour

Administrative data ^

Endpoint
 ... Remarks

Type of information
 ... Other ... Remarks

Adequacy of study
 ...
 Robust study summary
 Used for classification
 Used for SDS

Study period

Reliability
 ... Other

Rationale for reliability incl. deficiencies
 ... Other ... Remarks

Data waiving

Justification for data waiving

Justification for type of information
 |

Attached justification

Attached justification	Reason / purpose
<input type="text" value="N"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
<input type="text" value="C"/>	<input type="text" value="C"/>	<input type="text" value="C"/>

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
<input type="text" value="B"/>	<input type="text" value="B"/>	<input type="text" value="C"/>	<input type="text" value="C"/>	<input type="text" value="B"/>	<input type="text" value="N"/>	<input type="text" value="N"/>	<input type="text" value="C"/>	<input type="text" value="N"/>	<input type="text" value="C"/>	<input type="text" value="N"/>

+ Add... Delete Move up Move down Go to link target

Data access
 ... Other ... Remarks

Data protection claimed
 ... Remarks

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
C	C	C	C

+ Add... Edit... Delete Move up Move down

Principles of method if other than guideline

A | X

C

GLP compliance

C Remarks N

Study type

C Other C Remarks N

Company study no. ⚠

C

Report date

C

Test water

C Remarks N

Test material ^

Test material information

Specific details on test material used for the study

A | X

C

Wood preservative ^

Supplier of the preservative (name)

C

Type of formulation

C Other C

Preservative (specific and unique name or code)

C Other C

Active Ingredient(s) (trade or common name)

C Other C

Composition (in % m/m)

C

Coformulants

C

Retention or loading specified for tested wood

C Remarks N

10.4 Additional information on environmental fate and behaviour *(also in Biocidal Product under 10.2 Further studies on fate and behaviour in the environment)*

NB: Biocidal Product **A** → **C**

Administrative data ^

A

Description of key information ^

Normal Aharoni 8 A B I U

A

Additional information ^

Normal Aharoni 8 A B I U

C

Administrative data ^

Endpoint
A ... Remarks N ...

Type of information
A ... Other A ... Remarks N ...

Adequacy of study
A ...
 Robust study summary N
 Used for classification N
 Used for SDS N

Study period
C ...

Reliability
A ... Other A ...

Rationale for reliability incl. deficiencies
C ... Other C ... Remarks N ...

Data waiving
C ...

Justification for data waiving
C

Justification for type of information
C

Attached justification

Attached justification	Reason / purpose
N	N

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
A	A	N

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	A	A	B	N	N	A	N	A	N

+ Add... Delete Move up Move down Go to link target

Data access
N ... Other N ... Remarks N ...

Data protection claimed
N ... Remarks N ...

Materials and methods

Test guideline

Qualifier	Guideline	Version / remarks	Deviations

Add...
 Edit...
 Delete
 Move up
 Move down

Principles of method if other than guideline

GLP compliance

... Remarks

Type of study / information

Test material

Test material information

Specific details on test material used for the study

Any other information on materials and methods incl. tables

Results and discussion

Any other information on results incl. tables

Overall remarks, attachments ^

Overall remarks

Normal Aharoni 8
A B I U
≡ ≡ ≡
1/2 ≡
A ↻

N

Attached background material

Attached document	Remarks
N	N

+ Add...
 ✎ Edit...
 ✕ Delete
 ↑ Move up
 ↓ Move down

Attached full study report

N

+ Add...
 📄 Open
 ✎ Properties
 ✕ Delete
 ↑ Move up
 ↓ Move down

Illustration (picture/graph)

N

🔍 Zoom
 + Load
 ✕ Delete

Applicant's summary and conclusion ^

Conclusions

C
...


Executive summary

Normal Aharoni 8
A B I U
≡ ≡ ≡
1/2 ≡
A ↻

C






10.4 Biocidal Product – Testing for distribution and dissipation

Administrative data ^





Link to relevant study record(s) ^

Study name / type


 Add...  Delete  Move up  Move down  Go to link target


Description of key information ^





Additional information ^





Administrative data ^

Endpoint
 ... Remarks **N** ...

Type of information
 ... Other **C** ... Remarks **N** ...

Adequacy of study
 ...
 Robust study summary **N**
 Used for classification **N**
 Used for SDS **N**

Study period
 ...

Reliability
 ... Other **C** ...

Rationale for reliability incl. deficiencies
 ... Other **C** ... Remarks **N** ...

Data waiving
 ...

Justification for data waiving

Justification for type of information
 ...

Attached justification

Attached justification	Reason / purpose
N	N

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
C	C	C

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
B	B	C	C	B	N	N	C	N	C	N

+ Add... Delete Move up Move down Go to link target

Data access
 ... Other **N** ... Remarks **N** ...

Data protection claimed
 ... Remarks **N** ...

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
C	C	C	C

+ Add... Edit... Delete Move up Move down

Principles of method if other than guideline

A X

C

GLP compliance

C ... Remarks N

Type of study

C ... Other C

Media

C ... Other C

Test material ^

Test material information

Specific details on test material used for the study

A X

C

Any other information on materials and methods incl. tables ^

Normal Aharoni 8 A B I U

N

Results and discussion ^

Any other information on results incl. tables ^

Normal Aharoni 8 A B I U

C

Overall remarks, attachments ^

Overall remarks

Normal Aharoni 8 A B I U [List icons]

N

Attached background material

Attached document	Remarks
N	N

+ Add...
Edit...
X Delete
↑ Move up
↓ Move down

Attached full study report

N

+ Add...
Open
Properties
X Delete
↑ Move up
↓ Move down

Illustration (picture/graph)

N

Zoom
+ Load
X Delete

Applicant's summary and conclusion ^

Conclusions

C

Executive summary

Normal Aharoni 8 A B I U [List icons]

C

Administrative data ^

Endpoint
 ... Remarks

Type of information
 ... Other ... Remarks

Adequacy of study
 ...
 Robust study summary
 Used for classification
 Used for SDS

Study period

Reliability
 ... Other

Rationale for reliability incl. deficiencies
 ... Other ... Remarks

Data waiving

Justification for data waiving

Justification for type of information

Attached justification

Attached justification	Reason / purpose
<input type="text" value="N"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down

Cross-reference

Reason / purpose	Related information	Remarks
<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Edit... Delete Move up Move down Go to link target

Data source ^

Reference

Title	Author	Reference t...	Year	Bibliographi...	Testing labo...	Report no.	Company o...	Company st...	Report date	Remarks
<input type="text" value="B"/>	<input type="text" value="B"/>	<input type="text" value="A"/>	<input type="text" value="A"/>	<input type="text" value="B"/>	<input type="text" value="N"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>	<input type="text" value="A"/>	<input type="text" value="N"/>

+ Add... Delete Move up Move down Go to link target

Data access
 ... Other ... Remarks

Data protection claimed
 ... Remarks

Materials and methods ^

Test guideline

Qualifier	Guideline	Version / remarks	Deviations
A	A	A	A

+ Add... Edit... Delete Move up Move down

Principles of method if other than guideline

A X

A

GLP compliance

C ... Remarks N

Type of measurement

A ... Other A ... Remarks N

Media

A ... Other A

Test material ^

Test material information

Specific details on test material used for the study

A X

C

Study design ^

Details on sampling

A X

C

Details on analytical methods

A X

N

Any other information on materials and methods incl. tables ^

Normal Aharoni 8 A B I U ...

N

Results and discussion ^

Concentration

Key result	Country	Location	Substance or metabolite	Conc.	Remarks on result
A	A	A	A	A	A

+ Add... Edit... Delete Move up Move down

Details on results

A

Any other information on results incl. tables ^

Normal | Aharoni | 8 | A | B | I | U | | | | 1/2 = | A | ✎ | ↺

A

Overall remarks, attachments ^

Overall remarks

Normal | Aharoni | 8 | A | B | I | U | | | | 1/2 = | A | ✎ | ↺

N

Attached background material

Attached document	Remarks
N	N

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Attached full study report

N

+ Add...
📄 Open
✎ Properties
✕ Delete
↑ Move up
↓ Move down

Illustration (picture/graph)

N

🔍 Zoom
+ Load
✕ Delete

Applicant's summary and conclusion ^

Conclusions












C

Executive summary

Normal | Aharoni | 8 | A | B | I | U | | | | 1/2 = | A | ✎ | ↺




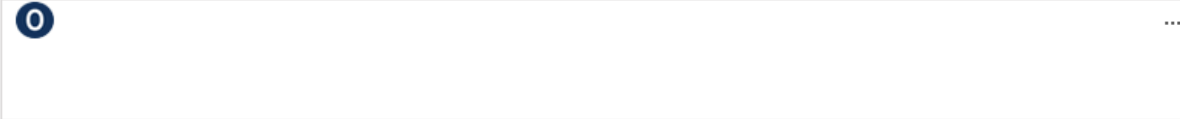





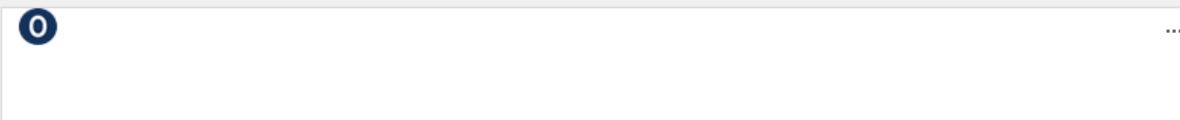

C

11. Measures to protect humans, animals and the environment

Administrative data ^	
 	
Measures to protect humans, animals and the environment ^	
Recommended methods and precautions concerning storage of active substance/biocidal product; shelf-life of biocidal product	
	...
Recommended methods and precautions concerning handling and transport	
	...
Recommended methods and precautions concerning fire; in case of fire nature of reaction products, combustion gases etc.	
	...
Particulars of likely direct or indirect adverse effects	
	...
First aid instructions, antidotes	
	...
Emergency measures to protect environment in case of accident	
	...
Control measures of repellents or poison included in the biocidal product, to prevent action against non-target organisms (relevant for biocidal products only)	
	...
Possibility of destruction or decontamination following release in or on the following: ^	
Air	
	...
Water, including drinking water	
	...

Soil	A	...
Procedures for waste management of active substance/biocidal product, and if appropriate, its packaging: ^		
Possibility of reuse or recycling	A	...
Possibility of neutralisation of effects	A	...
Conditions for controlled discharge including leachate qualities on disposal	A	...
Conditions for controlled incineration	A	...
Instructions for safe disposal of the biocidal product and its packaging for different groups of users (relevant for biocidal products only)	A	...
Procedures, if any, for cleaning application equipment (relevant for biocidal products only)	A	...

11. Biocidal Product – Measures to protect humans, animals and the environment

Administrative data ^	
 	
Measures to protect humans, animals and the environment ^	
Recommended methods and precautions concerning storage of active substance/biocidal product; shelf-life of biocidal product	
	...
Recommended methods and precautions concerning handling and transport	
	...
Recommended methods and precautions concerning fire; in case of fire nature of reaction products, combustion gases etc.	
	...
Particulars of likely direct or indirect adverse effects	
	...
First aid instructions, antidotes	
	...
Emergency measures to protect environment in case of accident	
	...
Control measures of repellents or poison included in the biocidal product, to prevent action against non-target organisms (relevant for biocidal products only)	
	...
Possibility of destruction or decontamination following release in or on the following: ^	
Air	
	...
Water, including drinking water	
	...

Soil	<input type="radio"/>	...
Procedures for waste management of active substance/biocidal product, and if appropriate, its packaging: ^		
Possibility of reuse or recycling	<input type="radio"/>	...
Possibility of neutralisation of effects	<input type="radio"/>	...
Conditions for controlled discharge including leachate qualities on disposal	<input type="radio"/>	...
Conditions for controlled incineration	<input type="radio"/>	...
Instructions for safe disposal of the biocidal product and its packaging for different groups of users (relevant for biocidal products only)	<input type="radio"/>	...
Procedures, if any, for cleaning application equipment (relevant for biocidal products only)	<input type="radio"/>	...

12. Classification & Labelling

12.1 GHS

N

General Information ^

Name ...

Not classified ...

Implementation ...

Remarks

A

Not classified A

Implementation ...

Remarks

A

Related composition ^

Related composition

A

+ Add... X Delete ↑ Move up ↓ Move down > Go to link target

Classification ^

Physical Hazards ^

	Hazard category	Hazard statement	Reason for no classification
Explosives	A ...	A ...	A ...
Flammable gases and chemically...	A ...	A ...	A ...
Aerosols	A ...	A ...	A ...
Oxidising gases	A ...	A ...	A ...
Gases under pressure	A ...	A ...	A ...
Flammable liquids	A ...	A ...	A ...
Flammable solids	A ...	A ...	A ...
Self-reactive substances and m...	A ...	A ...	A ...
Pyrophoric liquids	A ...	A ...	A ...
Pyrophoric solids	A ...	A ...	A ...

Self-heating substances and mi...	A	...	A	...	A	...
Substances and mixtures which ...	A	...	A	...	A	...
Oxidising liquids	A	...	A	...	A	...
Oxidising solids	A	...	A	...	A	...
Organic peroxides	A	...	A	...	A	...
Corrosive to metals	A	...	A	...	A	...
Desensitized explosives	A	...	A	...	A	...
Health hazards ^						
	Hazard category		Hazard statement		Reason for no classification	
Acute toxicity - oral	A	...	A	...	A	...
Acute toxicity - dermal	A	...	A	...	A	...
Acute toxicity - inhalation	A	...	A	...	A	...
Skin corrosion / irritation	A	...	A	...	A	...
Serious eye damage / eye irrit...	A	...	A	...	A	...
Respiratory sensitisation	A	...	A	...	A	...
Skin sensitisation	A	...	A	...	A	...
Aspiration hazard	A	...	A	...	A	...
Reproductive toxicity ^						
	Hazard category		Hazard statement		Reason for no classification	
Reproductive toxicity	A	...	A	...	A	...
Specific effect	A
Route of exposure	A	...	Remarks A			...
Effects on or via lactation	A	...	A	...	A	...
Germ cell mutagenicity ^						
	Hazard category		Hazard statement		Reason for no classification	
Germ cell mutagenicity	A	...	A	...	A	...
Route of exposure	A	...	Remarks A			...
Carcinogenicity ^						
	Hazard category		Hazard statement		Reason for no classification	
Carcinogenicity	A	...	A	...	A	...
Route of exposure	A	...	Remarks A			...
Specific target organ toxicity - single ^						
	Hazard category		Hazard statement		Reason for no classification	
Specific target organ toxicity...	A	...	A	...	A	...
Affected organs	A
Route of exposure	A	...	Remarks A			...
Specific target organ toxicity - repeated ^						
	Hazard category		Hazard statement		Reason for no classification	
Specific target organ toxicity...	A	...	A	...	A	...
Affected organs	A
Route of exposure	A	...	Remarks A			...

Specific concentration limits ^

Concentration range (%)

Hazard categories

Environmental hazards ^

Aquatic environment ^

	Hazard category	Hazard statement	Reason for no classification
Hazardous to the aquatic environment	A	A	A
Hazardous to the aquatic environment	A	A	A

M factor ^

M-Factor acute

M-Factor chronic

Ozone layer ^

	Hazard category	Hazard statement	Reason for no classification
Hazardous to the ozone layer	A	A	A

Additional hazard classes ^

Additional hazard classes

Additional hazard statements

Labelling ^

Signal word

Hazard pictogram ^

Code

Hazard statements ^

Hazard statement

Additional text

Precautionary statements ^

Precautionary statement
A

Additional text
A

Additional labelling requirements ^

CLP supplemental hazard statement

CLP supplemental hazard statement
A

Additional text
A

Additional labelling
A

Notes ^

A

12.1 Biocidal Product – GHS

N

General Information ^

Name

0

Not classified 0

Implementation

0
Other 0

Remarks

0

Normal
Aharoni
8

A
B
I
U

Related composition ^

Related composition

0

+ Add...
✕ Delete
↑ Move up
↓ Move down
➔ Go to link target

Classification ^

Physical Hazards ^

	Hazard category	Hazard statement	Reason for no classification
Explosives	0	0	0
Flammable gases and chemically...	0	0	0
Aerosols	0	0	0
Oxidising gases	0	0	0
Gases under pressure	0	0	0
Flammable liquids	0	0	0
Flammable solids	0	0	0
Self-reactive substances and m...	0	0	0
Pyrophoric liquids	0	0	0
Pyrophoric solids	0	0	0
Self-heating substances and mi...	0	0	0
Substances and mixtures which ...	0	0	0
Oxidising liquids	0	0	0
Oxidising solids	0	0	0
Organic peroxides	0	0	0
Corrosive to metals	0	0	0
Desensitized explosives	0	0	0

Health hazards ^

	Hazard category	Hazard statement	Reason for no classification
Acute toxicity - oral	0	0	0
Acute toxicity - dermal	0	0	0
Acute toxicity - inhalation	0	0	0
Skin corrosion / irritation	0	0	0
Serious eye damage / eye irrit...	0	0	0
Respiratory sensitisation	0	0	0
Skin sensitisation	0	0	0
Aspiration hazard	0	0	0

Reproductive toxicity ^

	Hazard category	Hazard statement	Reason for no classification
Reproductive toxicity	0	0	0
Specific effect	0		
Route of exposure	0	Remarks 0	
Effects on or via lactation	0	0	0

Germ cell mutagenicity ^

	Hazard category	Hazard statement	Reason for no classification
Germ cell mutagenicity	0	0	0
Route of exposure	0	Remarks 0	

Carcinogenicity ^

	Hazard category	Hazard statement	Reason for no classification
Carcinogenicity	0	0	0
Route of exposure	0	Remarks 0	

Specific target organ toxicity - single ^

	Hazard category	Hazard statement	Reason for no classification
Specific target organ toxicity...	0	0	0
Affected organs	0		
Route of exposure	0	Remarks 0	

Specific target organ toxicity - repeated ^

	Hazard category	Hazard statement	Reason for no classification
Specific target organ toxicity...	0	0	0
Affected organs	0		
Route of exposure	0	Remarks 0	

Specific concentration limits ^

Concentration range (%)

0 0 0 0

Hazard categories

0

Environmental hazards ^

Aquatic environment ^

	Hazard category	Hazard statement	Reason for no classification
Hazardous to the aquatic enviro...	0	0	0
Hazardous to the aquatic enviro...	0	0	0

M factor ^

M-Factor acute

0

M-Factor chronic

0

Ozone layer ^

	Hazard category	Hazard statement	Reason for no classification
Hazardous to the ozone layer	0	0	0

Additional hazard classes ^

Additional hazard classes

0

Additional hazard statements

0

Labelling ^

Signal word

0

Hazard pictogram ^

Code

0

Hazard statements ^

Hazard statement

0

Additional text

0

Precautionary statements ^

Precautionary statement

0

Additional text

0

Additional labelling requirements ^

CLP supplemental hazard statement

CLP supplemental hazard statement

Additional text

Additional labelling

Additional labelling

Notes ^

12.2 DSD – DPD

N

General information ^

Name ...

Not classified

Status ...

Index number ...

ATP inserted ...

ATP last update ...

Remarks ...

Related composition ^

Related composition

Classification ^

	Classification	Reason for no classification
Explosiveness
Oxidising properties
Flammability
Thermal stability
Acute toxicity
Acute toxicity - irreversible
Repeated dose toxicity
Irritation / corrosion
Sensitisation
Carcinogenicity
Mutagenicity - genetic toxic...
Toxicity to reproduction - fer...
Toxicity to reproduction - dev...
Toxicity to reproduction - bre...
Environment

Labelling ^

Indication of danger ^

Indication of danger

▼ A

Risk phrases ^

Risk phrases

▼ A

Safety phrases ^

Code

▼ A ...

Additional text

▼ A ...

Specific concentration limits ^

Concentration range (%)

▼ A A ▼ A A

Classification

▼ A

Notes ^

▼ A ...

12.2 Biocidal Product – DSD-DPD

N

General information ^

Name ...

Not classified

Status ... Other ...

Index number ...

ATP inserted ...

ATP last update ...

Remarks ...

Related composition ^

Related composition

+ Add...
✕ Delete
↑ Move up
↓ Move down
➤ Go to link target

Classification ^

	Classification	Reason for no classification
Explosiveness	<input type="text" value="0"/>	<input type="text" value="0"/> ...
Oxidising properties	<input type="text" value="0"/>	<input type="text" value="0"/> ...
Flammability	<input type="text" value="0"/>	<input type="text" value="0"/> ...
Thermal stability	<input type="text" value="0"/>	<input type="text" value="0"/> ...
Acute toxicity	<input type="text" value="0"/>	<input type="text" value="0"/> ...
Acute toxicity - irreversible ...	<input type="text" value="0"/>	<input type="text" value="0"/> ...
Repeated dose toxicity	<input type="text" value="0"/>	<input type="text" value="0"/> ...
Irritation / corrosion	<input type="text" value="0"/>	<input type="text" value="0"/> ...
Sensitisation	<input type="text" value="0"/> ...	<input type="text" value="0"/> ...
Carcinogenicity	<input type="text" value="0"/> ...	<input type="text" value="0"/> ...
Mutagenicity - genetic toxicit...	<input type="text" value="0"/> ...	<input type="text" value="0"/> ...
Toxicity to reproduction - fer...	<input type="text" value="0"/> ...	<input type="text" value="0"/> ...
Toxicity to reproduction - dev...	<input type="text" value="0"/> ...	<input type="text" value="0"/> ...
Toxicity to reproduction - bre...	<input type="text" value="0"/> ...	<input type="text" value="0"/> ...
Environment	<input type="text" value="0"/>	<input type="text" value="0"/> ...

Labelling ^

Indication of danger ^

Indication of danger
▼ 0

Risk phrases ^

Risk phrases
▼ 0

Safety phrases ^

Code
0 ... ▼

Additional text
0 ...

Specific concentration limits ^

Concentration range (%)
0 ▼ 0 0 ▼ 0

Classification
▼ 0

Notes ^

0 ... ▼

12.3 Biocidal Product – Packaging (12.7. in Annex III of BPR)

Administrative data ^

N

Packaging ^

For a biocidal product family, specify to which biocidal product(s) it applies:

0

+ Add...
✕ Delete
↑ Move up
↓ Move down
➤ Go to link target

Type of packaging in contact with the biocidal product (container type)

0

...
▼

Other
...

Size of packaging in contact with the biocidal product (container size)

0
0
0
0
0

...
▼

Other
0
...

Material of packaging in contact with the biocidal product (container material)

0

...
▼

Other
0
...

Compatibility of the biocidal product with the packaging materials proposed to be in contact with the biocidal product

0
...

Further description of the packaging in contact with the biocidal product

0
...

Safety features of the packaging

0
...

Description of the secondary packaging (not in contact with the biocidal product)

0
...

Packaging related attachments

Type of attachment	Attached document	Remarks
N	N	N

+ Add...
✎ Edit...
✕ Delete
↑ Move up
↓ Move down

Administrative data ^

A

Assessed Substance ^

Assessed substance
 ... ▼

Reference substance
 > 🔗

Composition of assessed substance

+ Add...
X Delete
↑ Move up
↓ Move down
> Go to link target

PBT status of the assessed substance
 ... ▼

Remark for assessed substance

Results of detailed PBT / vPvB assessment ^

Persistence ^

Evidence of non-P / non-vP properties ^

Screening criteria ^

Not P and not vP based on: readily biodegradable
 Remark

Not P and not vP based on: other screening test(s) (e.g. enhanced ready biodegradability, inherent biodegradability test) under valid conditions
 Remark

Criteria based on Annex XIII of REACH ^

Not P and not vP based on
 $T_{1/2} \leq 60$ days in marine water

and $T_{1/2} \leq 40$ days in fresh- or estuarine water

and $T_{1/2} \leq 180$ days in marine sediment

and $T_{1/2} \leq 120$ days in fresh- or estuarine sediment

and $T_{1/2} \leq 120$ days in soil

P but not vP based on

T½ ≤ 60 days in marine, fresh- or estuarine water
C

and T½ ≤ 180 days in marine, fresh- or estuarine sediment
C

and T½ ≤ 180 days in soil
C

Other evidence of non-P / non-vP properties ^
Remark
C

Further information for the PBT assessment is necessary ^
Remark
C

Evidence of P or vP properties ^
Remark
C

Conclusion ^
Conclusion on P / vP properties
C ... ▾
Remark
C

Bioaccumulation ^

Evidence of non-B / non-vB properties ^
Screening criteria ^

C Not B and not vB based on: Log Kow ≤ 4.5
Remark
C

Criteria based on Annex XIII of REACH ^

C Not B and not vB based on: BCF ≤ 2,000 L/kg
Remark
C

C B but not vB based on: 2,000 < BCF ≤ 5,000 L/kg
Remark
C

Other evidence of non-B / non-vB properties ^
Remark
C

<p>Further information for the PBT assessment is necessary ^</p> <p>Remark</p> <p>C</p>
<p>Evidence of B or vB properties ^</p> <p>Remark</p> <p>C</p>
<p>Conclusion ^</p> <p>Conclusion on B / vB properties</p> <p>C ... ▾</p> <p>Remark</p> <p>C</p>
<p>Toxicity ^</p> <p>Evidence of non-T properties ^</p> <p>Criteria based on Annex XIII of REACH ^</p> <p>C <input type="checkbox"/> Not T based on:</p> <p>EC10 or NOEC \geq 0.01 mg/L for marine / freshwater organisms (long-term toxicity)</p> <p>C</p> <p>and substance is not classified as carcinogenic (category 1 or 2), mutagenic (category 1 or 2), or toxic for reproduction (category 1, 2 or 3) according to Directive 67/548/EEC or carcinogenic (category 1A or 1B), germ cell mutagenic (category 1A or 1B), or toxic for reproduction (category 1A, 1B or 2) according to Regulation EC No 1272/2008</p> <p>C</p> <p>and no other evidence of chronic toxicity, as identified by the classifications T, R48 or Xn, R48 according to Directive 67/548/EEC or specific tar organ toxicity after repeated exposure (STOT RE category 1 or 2) according to Regulation EC No 1272/2008</p> <p>C</p> <p>Other evidence of non-T properties ^</p> <p>Remark</p> <p>C</p>
<p>Further information for the PBT assessment is necessary ^</p> <p>Remark</p> <p>C</p>
<p>Evidence of T properties ^</p> <p>C <input type="checkbox"/> Screening criteria: L(E)C50 < 0.01 mg/L</p> <p>C <input type="checkbox"/> other evidence</p> <p>Remark</p> <p>C</p>
<p>Conclusion ^</p> <p>Conclusion on T properties</p> <p>C ... ▾</p> <p>Remark</p> <p>C</p>

EUROPEAN CHEMICALS AGENCY
ANNANKATU 18, P.O. BOX 400,
00121 HELSINKI, FINLAND
ECHA.EUROPA