

**PKT SORB-IT 1G GDT 22x38 8000/ALU/C**

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Substance key: SC0000901386

Revision Date: 12.05.2022

Version : 1 - 3 / EU

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**SECTION 1: Identification of the substance/mixture and of the company/undertaking**

**1.1. Product identifier**

**Trade name**

PKT SORB-IT 1G GDT 22x38 8000/ALU/C

**Material number:** 328115

**REACH - Registration number according to article 20(3):** 01-2119379499-16-0016, 01-2119379499-16-0030, 01-2119379499-16-0279

**CAS number :** 7631-86-9

**1.2. Relevant identified uses of the substance or mixture and uses advised against**

**Relevant identified uses of the substance or mixture**

Type of use : Adsorbing medium for technical applications

**Relevant identified uses of the substance or mixture**

Type of use : Packaging

**1.3. Details of the supplier of the safety data sheet**

**Identification of the company**

Clariant Produkte (Deutschland) GmbH  
Ostenriederstrasse 15  
85368 Moosburg  
Telephone no. : +49 (0)8761/82-0

**Information about the substance/mixture**

BU Adsorbents & Additives  
Product Stewardship  
e-mail: SDS.Europe@clariant.com

**1.4. Emergency telephone number**

00800-5121 5121 (24 h)

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**SECTION 2: Hazards identification**

**2.1 Classification of the substance or mixture**

**Classification (REGULATION (EC) No 1272/2008)**

Not classified, does not meet the criteria for classification.  
Not a hazardous substance or mixture.

**2.2 Label elements**

**Labelling (REGULATION (EC) No 1272/2008)**

Not a hazardous substance or mixture., The product does not require classification and labelling as hazardous according to CLP/GHS.

**2.3 Other hazards**

This substance/mixture contains no components considered to be either persistent, bioaccumulative and toxic (PBT), or very persistent and very bioaccumulative (vPvB) at levels of 0.1% or higher.

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Ecological information: The substance/mixture does not contain components considered to have endocrine disrupting properties according to REACH Article 57(f) or Commission Delegated regulation (EU) 2017/2100 or Commission Regulation (EU) 2018/605 at levels of 0.1% or higher.

Toxicological information: The substance/mixture does not contain components considered to have endocrine disrupting properties according to REACH Article 57(f) or Commission Delegated regulation (EU) 2017/2100 or Commission Regulation (EU) 2018/605 at levels of 0.1% or higher.

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**SECTION 3: Composition/information on ingredients**

**3.1 Substances**

CAS-No. : 7631-86-9

**Components**

| Chemical name                                | CAS-No.<br>EC-No.      | Concentration (%)<br>w/w) | M-Factor, SCL, ATE |
|--|------------------------|---------------------------|--------------------|
| Substances with a workplace exposure limit : |                        |                           |                    |
| Amorphous silicon dioxide                    | 7631-86-9<br>231-545-4 | >= 90 - <= 100            |                    |

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**SECTION 4: First aid measures**

**4.1 Description of first aid measures**

General advice : none

If inhaled : Move to fresh air.

In case of skin contact : Wash off immediately with soap and plenty of water.

In case of eye contact : Rinse immediately with plenty of water, also under the eyelids, for at least 15 minutes.

If swallowed : Rinse mouth.

**4.2 Most important symptoms and effects, both acute and delayed**

Symptoms : There are no acute and delayed symptoms and effects observed.

Risks : No information available.

**4.3 Indication of any immediate medical attention and special treatment needed**

Treatment : Treat symptomatically.

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**SECTION 5: Firefighting measures**

**5.1 Extinguishing media**

Suitable extinguishing media : Use extinguishing measures that are appropriate to local

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circumstances and the surrounding environment.

Unsuitable extinguishing media : High volume water jet

**5.2 Special hazards arising from the substance or mixture**

Specific hazards during firefighting : None known.

**5.3 Advice for firefighters**

Special protective equipment for firefighters : Wear an approved positive pressure self-contained breathing apparatus in addition to standard fire fighting gear.

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**SECTION 6: Accidental release measures**

**6.1 Personal precautions, protective equipment and emergency procedures**

Personal precautions : Do not breathe dust.  
Avoid dust formation.

**6.2 Environmental precautions**

Environmental precautions : No special environmental precautions required.

**6.3 Methods and material for containment and cleaning up**

**6.4 Reference to other sections**

For personal protection see section 8., For disposal considerations see section 13.

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**SECTION 7: Handling and storage**

**7.1 Precautions for safe handling**

Advice on protection against fire and explosion : No special precautions required.

Hygiene measures : Wash off with warm water and soap.

**7.2 Conditions for safe storage, including any incompatibilities**

Requirements for storage areas and containers : Keep container tightly closed and dry.

Advice on common storage : No materials to be especially mentioned.

Further information on storage stability : Stable under recommended storage conditions.

**7.3 Specific end use(s)**

Specific use(s) : Not relevant

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**SECTION 8: Exposure controls/personal protection**

**8.1 Control parameters**

**Further occupational exposure limits**

| Description                     | Value type | Control parameters    | Basis      |
|---------------------------------|------------|-----------------------|------------|
| Sorb-It®Carcinogens or mutagens | TWA        | 0,1 mg/m <sup>3</sup> | 2004/37/EC |

**8.2 Exposure controls**

**Engineering measures**

Use ventilation adequate to keep exposures below recommended exposure limits. See the safety datasheet.

**Personal protective equipment**

Eye protection : Safety glasses

Skin and body protection : not required under normal use

Respiratory protection : Half mask with a particle filter P2 (EN 143)

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**SECTION 9: Physical and chemical properties**

**9.1 Information on basic physical and chemical properties**

Physical state : granular

Colour : various

Odour : none

Odour Threshold : not available

Melting point : not determined

Boiling point : not determined

Flammability : not determined

Upper explosion limit / upper flammability limit : Not applicable

Lower explosion limit / Lower flammability limit : Not applicable

Flash point : does not flash

Auto-ignition temperature : Not applicable

Decomposition temperature : no data available

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pH : 3 - 5 (20 °C)  
Method: aqueous suspension

Viscosity  
Viscosity, dynamic : Not applicable

Viscosity, kinematic : Not applicable

Solubility(ies)  
Water solubility : insoluble

Partition coefficient: n-  
octanol/water : no data available

Vapour pressure : Not applicable

Density : not determined

Bulk density : 680 - 780 kg/m<sup>3</sup>

Relative vapour density : Not applicable

Particle characteristics  
Particle size : no data available

**9.2 Other information**

Metal corrosion rate : Not applicable

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**SECTION 10: Stability and reactivity**

**10.1 Reactivity**

Stable

**10.2 Chemical stability**

Stable

**10.3 Possibility of hazardous reactions**

Hazardous reactions : No dangerous reaction known under conditions of normal use. Stable

**10.4 Conditions to avoid**

Conditions to avoid : None known.

**10.5 Incompatible materials**

Materials to avoid : Water

**10.6 Hazardous decomposition products**

No decomposition if stored and applied as directed.

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**SECTION 11: Toxicological information**

**11.1 Information on hazard classes as defined in Regulation (EC) No 1272/2008**

**Acute toxicity**

**Product:**

- Acute oral toxicity : Remarks: Not applicable
- Acute inhalation toxicity : LC50 (Rat, male and female): > 2,08 mg/l  
Exposure time: 4 h  
Test atmosphere: dust/mist  
Method: OECD Test Guideline 403  
GLP: yes  
Assessment: The substance or mixture has no acute inhalation toxicity
- Acute dermal toxicity : LD50 (Rabbit): > 5.000 mg/kg  
Method: Other  
GLP: no

**Components:**

**Amorphous silicon dioxide:**

- Acute oral toxicity : LD50 (Rat, male and female): > 5.000 mg/kg  
Method: OECD Test Guideline 401  
GLP: yes  
Remarks: No significant adverse effects were reported
- Acute inhalation toxicity : LC50 (Rat, male and female): > 2,08 mg/l  
Exposure time: 4 h  
Test atmosphere: dust/mist  
Method: OECD Test Guideline 403  
GLP: yes  
Assessment: The substance or mixture has no acute inhalation toxicity
- Acute dermal toxicity : LD50 (Rabbit): > 5.000 mg/kg  
Method: Other  
GLP: no

**Skin corrosion/irritation**

**Product:**

- Species : Rabbit  
Exposure time : 4 h  
Method : OECD Test Guideline 404  
Result : No skin irritation  
GLP : yes

**Components:**

**Amorphous silicon dioxide:**

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Species : Rabbit  
Exposure time : 4 h  
Method : OECD Test Guideline 404  
Result : No skin irritation  
GLP : yes

**Serious eye damage/eye irritation**

**Product:**

Species : Rabbit  
Exposure time : 24 h  
Method : OECD Test Guideline 405  
Result : No eye irritation  
GLP : yes

**Components:**

**Amorphous silicon dioxide:**

Species : Rabbit  
Exposure time : 24 h  
Method : OECD Test Guideline 405  
Result : No eye irritation  
GLP : yes

**Respiratory or skin sensitisation**

**Product:**

Remarks : no data available

**Components:**

**Amorphous silicon dioxide:**

Remarks : no data available

**Germ cell mutagenicity**

**Product:**

Genotoxicity in vitro : Test Type: Ames test  
Test system: Salmonella typhimurium  
Concentration: 667 - 10000 µg/plate  
Metabolic activation: with and without metabolic activation  
Method: OECD Test Guideline 471  
Result: negative  
GLP: yes

Test Type: In vitro gene mutation study in mammalian cells  
Test system: Chinese hamster ovary cells  
Concentration: 10 - 500 µg/ml  
Metabolic activation: with and without metabolic activation  
Method: OECD Test Guideline 476  
Result: negative  
GLP: yes

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Test Type: Chromosome aberration test in vitro  
Test system: Chinese hamster ovary cells  
Concentration: 38 - 1000 µg/ml  
Metabolic activation: with and without metabolic activation  
Method: OECD Test Guideline 473  
Result: negative  
GLP: yes

Genotoxicity in vivo : Test Type: Cytogenetic assay  
Species: Rat (male)  
Strain: Fischer F344  
Application Route: Inhalation  
Exposure time: 13 w, 6 h/d, 5 d/wk  
Dose: ca. 50 mg/m<sup>3</sup>  
Method: Other  
Result: negative  
GLP: No information available.

Germ cell mutagenicity-  
Assessment : In vitro tests did not show mutagenic effects, In vivo tests did not show mutagenic effects

**Components:**

**Amorphous silicon dioxide:**

Genotoxicity in vitro : Test Type: Ames test  
Test system: Salmonella typhimurium  
Concentration: 667 - 10000 µg/plate  
Metabolic activation: with and without metabolic activation  
Method: OECD Test Guideline 471  
Result: negative  
GLP: yes

Test Type: In vitro gene mutation study in mammalian cells  
Test system: Chinese hamster ovary cells  
Concentration: 10 - 500 µg/ml  
Metabolic activation: with and without metabolic activation  
Method: OECD Test Guideline 476  
Result: negative  
GLP: yes

Test Type: Chromosome aberration test in vitro  
Test system: Chinese hamster ovary cells  
Concentration: 38 - 1000 µg/ml  
Metabolic activation: with and without metabolic activation  
Method: OECD Test Guideline 473  
Result: negative  
GLP: yes

Genotoxicity in vivo : Test Type: Cytogenetic assay  
Species: Rat (male)  
Strain: Fischer F344  
Application Route: Inhalation



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Exposure time: 13 w, 6 h/d, 5 d/wk  
Dose: ca. 50 mg/m<sup>3</sup>  
Method: Other  
Result: negative  
GLP: No information available.

Germ cell mutagenicity-  
Assessment : In vitro tests did not show mutagenic effects, In vivo tests did not show mutagenic effects

**Carcinogenicity**

**Product:**

Species : Rat, male and female  
Application Route : oral (feed)  
Exposure time : 103 w  
Dose : 1,25 - 2,5 - 5 % in diet  
Control Group : yes  
Frequency of Treatment : daily  
NOAEL : ca. 1.800 - 3.000 mg/kg bw/day  
Method : OECD Test Guideline 453  
Result : negative  
GLP : No information available.

Carcinogenicity -  
Assessment : Not classifiable as a human carcinogen.

**Components:**

**Amorphous silicon dioxide:**

Species : Rat, male and female  
Application Route : oral (feed)  
Exposure time : 103 w  
Dose : 1,25 - 2,5 - 5 % in diet  
Control Group : yes  
Frequency of Treatment : daily  
NOAEL : ca. 1.800 - 3.000 mg/kg bw/day  
Method : OECD Test Guideline 453  
Result : negative  
GLP : No information available.

Carcinogenicity -  
Assessment : Not classifiable as a human carcinogen.

**Reproductive toxicity**

**Product:**

Effects on fertility : Species: Rat, male and female  
Strain: Sprague-Dawley  
Application Route: oral (feed)  
Dose: 497 (m), 509 (f) mg/kg  
General Toxicity - Parent: NOAEL: 497 mg/kg body weight  
General Toxicity F1: NOAEL: 497 mg/kg body weight  
Method: OECD Test Guideline 415

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GLP: no

Effects on foetal development : Test Type: Pre-natal  
Species: Rat  
Strain: wistar  
Application Route: oral (gavage)  
Dose: 13,5 - 62,7 - 292 - 1350mg/kg  
General Toxicity Maternal: NOAEL: 1.350 mg/kg body weight  
Teratogenicity: NOAEL: 1.350 mg/kg body weight  
Method: OECD Test Guideline 414  
GLP: no

Reproductive toxicity - Assessment : No evidence of adverse effects on sexual function and fertility, or on development, based on animal experiments.  
No teratogenic effects to be expected.

**Components:**

**Amorphous silicon dioxide:**

Effects on fertility : Test Type: One generation study  
Species: Rat, male and female  
Strain: Sprague-Dawley  
Application Route: oral (feed)  
Dose: 497 (m), 509 (f) mg/kg  
General Toxicity - Parent: NOAEL: 497 mg/kg body weight  
General Toxicity F1: NOAEL: 497 mg/kg body weight  
Method: OECD Test Guideline 415  
GLP: no

Effects on foetal development : Test Type: Pre-natal  
Species: Rat  
Strain: wistar  
Application Route: oral (gavage)  
Dose: 13,5 - 62,7 - 292 - 1350mg/kg  
General Toxicity Maternal: NOAEL: 1.350 mg/kg body weight  
Teratogenicity: NOAEL: 1.350 mg/kg body weight  
Method: OECD Test Guideline 414  
GLP: no

Reproductive toxicity - Assessment : No evidence of adverse effects on sexual function and fertility, or on development, based on animal experiments.  
No teratogenic effects to be expected.

**STOT - single exposure**

**Product:**

Assessment : The substance or mixture is not classified as specific target organ toxicant, single exposure.

**Components:**

**Amorphous silicon dioxide:**

Assessment : The substance or mixture is not classified as specific target

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organ toxicant, single exposure.

**STOT - repeated exposure**

**Product:**

Assessment : The substance or mixture is not classified as specific target organ toxicant, repeated exposure.

**Components:**

**Amorphous silicon dioxide:**

Assessment : The substance or mixture is not classified as specific target organ toxicant, repeated exposure.

**Repeated dose toxicity**

**Product:**

Species : Rat, male and female  
NOAEL : 4000 - 4500 mg/kg bw/day  
Application Route : oral (feed)  
Exposure time : 13 w  
Number of exposures : continuously  
Dose : 0,5 - 2 - 6,7 % SI in diet  
Control Group : yes  
Method : OECD Test Guideline 408  
GLP : yes

Species : Rat, male and female  
NOAEL : 1,3 mg/m<sup>3</sup>  
LOAEL : 0,0059 mg/l  
Application Route : Inhalation  
Exposure time : 13 w  
Number of exposures : 6 hr/day; 5 days a week  
Dose : 1,3 - 5,9 - 31 mg/m<sup>3</sup>  
Control Group : yes  
Method : OECD Test Guideline 413  
GLP : yes

Application Route : Skin contact  
Remarks : This information is not available.

**Components:**

**Amorphous silicon dioxide:**

Species : Rat, male and female  
NOAEL : 4000 - 4500 mg/kg bw/day  
Application Route : oral (feed)  
Exposure time : 13 w  
Number of exposures : continuously  
Dose : 0,5 - 2 - 6,7 % SI in diet  
Control Group : yes  
Method : OECD Test Guideline 408  
GLP : yes

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Species : Rat, male and female  
NOAEL : 1,3 mg/m<sup>3</sup>  
LOAEL : 0,0059 mg/l  
Application Route : Inhalation  
Exposure time : 13 w  
Number of exposures : 6 hr/day; 5 days a week  
Dose : 1,3 - 5,9 - 31 mg/m<sup>3</sup>  
Control Group : yes  
Method : OECD Test Guideline 413  
GLP : yes

Application Route : Skin contact  
Remarks : This information is not available.

**Aspiration toxicity**

**Product:**

No aspiration toxicity classification

**Components:**

**Amorphous silicon dioxide:**

No aspiration toxicity classification

**11.2 Information on other hazards**

**Endocrine disrupting properties**

**Product:**

Assessment : The substance/mixture does not contain components considered to have endocrine disrupting properties according to REACH Article 57(f) or Commission Delegated regulation (EU) 2017/2100 or Commission Regulation (EU) 2018/605 at levels of 0.1% or higher.

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**SECTION 12: Ecological information**

**12.1 Toxicity**

**Product:**

Toxicity to fish : LL0 (Brachydanio rerio (zebrafish)): 10.000 mg/l  
End point: mortality  
Exposure time: 96 h  
Test Type: static test  
Analytical monitoring: no  
Method: OECD Test Guideline 203  
GLP: yes  
Remarks: The details of the toxic effect relate to the nominal concentration.

Toxicity to daphnia and other aquatic invertebrates : EL50 (Daphnia magna (Water flea)): > 1.000 mg/l  
End point: Immobilization

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Exposure time: 24 h  
Test Type: static test  
Analytical monitoring: no  
Method: OECD Test Guideline 202  
GLP: yes  
Remarks: The details of the toxic effect relate to the nominal concentration.

Toxicity to algae/aquatic plants : EL50 (Desmodesmus subspicatus (green algae)): > 10.000 mg/l  
End point: Growth rate  
Exposure time: 72 h  
Test Type: static test  
Analytical monitoring: no  
Method: OECD Test Guideline 201  
GLP: yes  
Remarks: By analogy with a product of similar composition  
The details of the toxic effect relate to the nominal concentration.

Toxicity to fish (Chronic toxicity) : NOEC: 86,03 mg/l  
Exposure time: 30 d  
Method: Other  
GLP: no  
Remarks: The value is given based on a SAR/AAR approach using OECD Toolbox, DEREK, VEGA QSAR models (CAESAR models), etc.

Toxicity to daphnia and other aquatic invertebrates (Chronic toxicity) : NOEC: 34,223 mg/l  
Exposure time: 30 d  
Method: Other  
GLP: no  
Remarks: The value is given based on a SAR/AAR approach using OECD Toolbox, DEREK, VEGA QSAR models (CAESAR models), etc.

Sediment toxicity : LC50: 148,41 mg/l  
Duration: 14 d  
Method: Other  
GLP: no  
Remarks: The value is given based on a SAR/AAR approach using OECD Toolbox, DEREK, VEGA QSAR models (CAESAR models), etc.

**Ecotoxicology Assessment**

Acute aquatic toxicity : This product has no known ecotoxicological effects.

Chronic aquatic toxicity : This product has no known ecotoxicological effects.

**Components:**

**Amorphous silicon dioxide:**

Toxicity to fish : LL0 (Brachydanio rerio (zebrafish)): 10.000 mg/l

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- End point: mortality  
Exposure time: 96 h  
Test Type: static test  
Analytical monitoring: no  
Method: OECD Test Guideline 203  
GLP: yes  
Remarks: The details of the toxic effect relate to the nominal concentration.
- Toxicity to daphnia and other aquatic invertebrates : EL50 (Daphnia magna (Water flea)): > 1.000 mg/l  
End point: Immobilization  
Exposure time: 24 h  
Test Type: static test  
Analytical monitoring: no  
Method: OECD Test Guideline 202  
GLP: yes  
Remarks: The details of the toxic effect relate to the nominal concentration.
- Toxicity to algae/aquatic plants : EL50 (Desmodesmus subspicatus (green algae)): > 10.000 mg/l  
End point: Growth rate  
Exposure time: 72 h  
Test Type: static test  
Analytical monitoring: no  
Method: OECD Test Guideline 201  
GLP: yes  
Remarks: By analogy with a product of similar composition  
The details of the toxic effect relate to the nominal concentration.
- Toxicity to fish (Chronic toxicity) : NOEC: 86,03 mg/l  
Exposure time: 30 d  
Method: Other  
GLP: no  
Remarks: The value is given based on a SAR/AAR approach using OECD Toolbox, DEREK, VEGA QSAR models (CAESAR models), etc.
- Toxicity to daphnia and other aquatic invertebrates (Chronic toxicity) : NOEC: 34,223 mg/l  
Exposure time: 30 d  
Method: Other  
GLP: no  
Remarks: The value is given based on a SAR/AAR approach using OECD Toolbox, DEREK, VEGA QSAR models (CAESAR models), etc.
- Sediment toxicity : LC50: 148,41 mg/l  
Duration: 14 d  
Method: Other  
GLP: no  
Remarks: The value is given based on a SAR/AAR approach using OECD Toolbox, DEREK, VEGA QSAR models (CAESAR models), etc.

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**Ecotoxicology Assessment**

Acute aquatic toxicity : This product has no known ecotoxicological effects.

Chronic aquatic toxicity : This product has no known ecotoxicological effects.

**12.2 Persistence and degradability**

**Product:**

Biodegradability : Remarks: Not applicable

**Components:**

**Amorphous silicon dioxide:**

Biodegradability : Remarks: Not applicable

**12.3 Bioaccumulative potential**

no data available

**12.4 Mobility in soil**

no data available

**12.5 Results of PBT and vPvB assessment**

**Product:**

Assessment : This substance/mixture contains no components considered to be either persistent, bioaccumulative and toxic (PBT), or very persistent and very bioaccumulative (vPvB) at levels of 0.1% or higher.

**Components:**

**Amorphous silicon dioxide:**

Assessment : The substance is not identified as a PBT or as a vPvB substance.

**12.6 Endocrine disrupting properties**

**Product:**

Assessment : The substance/mixture does not contain components considered to have endocrine disrupting properties according to REACH Article 57(f) or Commission Delegated regulation (EU) 2017/2100 or Commission Regulation (EU) 2018/605 at levels of 0.1% or higher.

**12.7 Other adverse effects**

**Product:**

Environmental fate and pathways : not available

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**Components:**

**Amorphous silicon dioxide:**

Environmental fate and pathways : not available

Additional ecological information : Do not allow to enter ground water, waterways or waste water.

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**SECTION 13: Disposal considerations**

**13.1 Waste treatment methods**

Product : Can be landfilled, when in compliance with local regulations. Contact waste disposal services.

Contaminated packaging : Dispose of as unused product.

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**SECTION 14: Transport information**

**Section 14.1. to 14.5.**

|             |                |
|-------------|----------------|
| <b>ADR</b>  | not restricted |
| <b>ADN</b>  | not restricted |
| <b>RID</b>  | not restricted |
| <b>IATA</b> | not restricted |
| <b>IMDG</b> | not restricted |

**14.6. Special precautions for user**

See sections 6 to 8 of this Safety Data Sheet.

**14.7. Maritime transport in bulk according to IMO instruments**

No transport as bulk according IBC - Code.

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**SECTION 15: Regulatory information**

**15.1 Safety, health and environmental regulations/legislation specific for the substance or mixture**

REACH - Restrictions on the manufacture, placing on the market and use of certain dangerous substances, mixtures and articles (Annex XVII) : Not applicable

REACH - Candidate List of Substances of Very High Concern for Authorisation (Article 59). : Not applicable

Regulation (EC) No 1005/2009 on substances that deplete the ozone layer : Not applicable

Regulation (EU) 2019/1021 on persistent organic pollutants (recast) : Not applicable



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Council Regulation (EC) No 111/2005 laying down rules for the monitoring of trade between the Community and third countries in drug precursors : Neither banned nor restricted

Regulation (EC) No 649/2012 of the European Parliament and the Council concerning the export and import of dangerous chemicals : Not applicable

REACH - List of substances subject to authorisation (Annex XIV) : Not applicable

**Other regulations:**

Apart from the data/regulations specified in this chapter, no further information is available concerning safety, health and environmental protection.

**The components of this product are reported in the following inventories:**

TCSI : On the inventory, or in compliance with the inventory  
AIIC : On the inventory, or in compliance with the inventory  
ENCS : On the inventory, or in compliance with the inventory  
ISHL : On the inventory, or in compliance with the inventory  
KECI : On the inventory, or in compliance with the inventory  
PICCS : On the inventory, or in compliance with the inventory  
IECSC : On the inventory, or in compliance with the inventory  
NZIoC : Not in compliance with the inventory  
TECI : On the inventory, or in compliance with the inventory

**15.2 Chemical safety assessment**

A Chemical Safety Assessment is not required for this substance.

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**SECTION 16: Other information**

**Full text of other abbreviations**

2004/37/EC : Europe. Directive 2004/37/EC on the protection of workers from the risks related to exposure to carcinogens or mutagens at work  
2004/37/EC / TWA : Long term exposure limit

ADN - European Agreement concerning the International Carriage of Dangerous Goods by Inland Waterways; ADR - Agreement concerning the International Carriage of Dangerous Goods by Road; AIIC - Australian Inventory of Industrial Chemicals; ASTM - American Society for the Testing of Materials; bw - Body weight; CLP - Classification Labelling Packaging Regulation;

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Regulation (EC) No 1272/2008; CMR - Carcinogen, Mutagen or Reproductive Toxicant; DIN - Standard of the German Institute for Standardisation; DSL - Domestic Substances List (Canada); ECHA - European Chemicals Agency; EC-Number - European Community number; ECx - Concentration associated with x% response; ELx - Loading rate associated with x% response; EmS - Emergency Schedule; ENCS - Existing and New Chemical Substances (Japan); ErCx - Concentration associated with x% growth rate response; GHS - Globally Harmonized System; GLP - Good Laboratory Practice; IARC - International Agency for Research on Cancer; IATA - International Air Transport Association; IBC - International Code for the Construction and Equipment of Ships carrying Dangerous Chemicals in Bulk; IC50 - Half maximal inhibitory concentration; ICAO - International Civil Aviation Organization; IECSC - Inventory of Existing Chemical Substances in China; IMDG - International Maritime Dangerous Goods; IMO - International Maritime Organization; ISHL - Industrial Safety and Health Law (Japan); ISO - International Organisation for Standardization; KECI - Korea Existing Chemicals Inventory; LC50 - Lethal Concentration to 50 % of a test population; LD50 - Lethal Dose to 50% of a test population (Median Lethal Dose); MARPOL - International Convention for the Prevention of Pollution from Ships; n.o.s. - Not Otherwise Specified; NO(A)EC - No Observed (Adverse) Effect Concentration; NO(A)EL - No Observed (Adverse) Effect Level; NOELR - No Observable Effect Loading Rate; NZIoC - New Zealand Inventory of Chemicals; OECD - Organization for Economic Co-operation and Development; OPPTS - Office of Chemical Safety and Pollution Prevention; PBT - Persistent, Bioaccumulative and Toxic substance; PICCS - Philippines Inventory of Chemicals and Chemical Substances; (Q)SAR - (Quantitative) Structure Activity Relationship; REACH - Regulation (EC) No 1907/2006 of the European Parliament and of the Council concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals; RID - Regulations concerning the International Carriage of Dangerous Goods by Rail; SADT - Self-Accelerating Decomposition Temperature; SDS - Safety Data Sheet; SVHC - Substance of Very High Concern; TCSI - Taiwan Chemical Substance Inventory; TECI - Thailand Existing Chemicals Inventory; TRGS - Technical Rule for Hazardous Substances; TSCA - Toxic Substances Control Act (United States); UN - United Nations; vPvB - Very Persistent and Very Bioaccumulative

**Further information**

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