

Safety data sheet

Page: 1/15

BASF safety data sheet. This is a translation of the country-specific safety data sheet into a language other than that required by law. This document does not replace the safety data sheet provided according to Regulation (EC) No 1907/2006.

Date / Revised: 12.04.2023 Version: 1.0

Date previous version: not applicable Previous version: none

Date / First version: 12.04.2023 Product: **Rheovis® HS 1980**

(ID no. 30492472/SDS_GEN_GR/EN)

Date of print 23.10.2025

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

1.1. Product identifier

Rheovis® HS 1980

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

Relevant identified uses: Raw material, for industrial use only

1.3. Details of the supplier of the safety data sheet

Company: BASF SE 67056 Ludwigshafen GERMANY Contact address: BASF Hellas S.A. Sindos Industrial Area 57022 Sindos GREECE

Telephone: +30 2310 797-195

E-mail address: gr-psr-hellas@basf.com

1.4. Emergency telephone number

Greek Poison Information Centre 0030 210 7793777 (24 hours / 7 days) International emergency number: Telephone: +49 180 2273-112

to Regulation (EC) No 1907/2006.

Version: 1.0 Previous version: none

Date / Revised: 12.04.2023
Date previous version: not applicable
Date / First version: 12.04.2023
Product: **Rheovis® HS 1980**

(ID no. 30492472/SDS_GEN_GR/EN)

Date of print 23.10.2025

SECTION 2: Hazards Identification

2.1. Classification of the substance or mixture

For the classification of the mixture the following methods have been applied: extrapolation on the concentration levels of the hazardous substances, on basis of test results and after evaluation of experts. The methodologies used are mentioned at the respective test results.

According to Regulation (EC) No 1272/2008 [CLP]

No need for classification according to GHS criteria for this product.

2.2. Label elements

According to Regulation (EC) No 1272/2008 [CLP]

The product does not require a hazard warning label in accordance with GHS criteria.

According to Regulation (EC) No 1272/2008 [CLP]

Labeling of special preparations (GHS):

EUH208: May produce an allergic reaction. Contains: mixture of: 5-chloro-2-methyl-2H-isothiazol-3-one and 2-methyl-2H-isothiazol-3-one

2.3. Other hazards

According to Regulation (EC) No 1272/2008 [CLP]

Product does not contain a substance above legal limits included in the list established in accordance with Article 59(1) of Regulation (EC) No 1907/2006 for having endocrine disrupting properties or is identified to have endocrine disrupting properties in accordance with the criteria set out in Commission Delegated Regulation (EU) 2017/2100 or Commission Regulation (EU) 2018/605. The product is under certain conditions capable of dust explosion.

The product does not contain a substance above legal limits fulfilling the PBT (persistent/bioaccumulative/toxic) criteria or the vPvB (very persistent/very bioaccumulative) criteria.

SECTION 3: Composition/Information on Ingredients

3.1. Substances

Not applicable

3.2. Mixtures

Chemical nature

Polymer based on:acrylic ester, methacrylic ester, copolymer

Regulatory relevant ingredients

mixture of: 5-chloro-2-methyl-2H-isothiazol-3-one and 2-methyl-2H-isothiazol-3-one

to Regulation (EC) No 1907/2006.

Date / Revised: 12.04.2023 Version: 1.0

Date previous version: not applicable Previous version: none

Date / First version: 12.04.2023 Product: **Rheovis® HS 1980**

(ID no. 30492472/SDS_GEN_GR/EN)

Date of print 23.10.2025

Content (W/W): >= 1,5 PPM - < 15 Acute Tox. 3 (oral)

PPM Acute Tox. 2 (Inhalation - mist)

CAS Number: 55965-84-9
REACH registration number: 012120764691-48

Acute Tox. 2 (dermal)
Skin Corr./Irrit. 1C
Eye Dam./Irrit. 1

INDEX-Number: 613-167-00-5

Skin Sens. 1A

Aquatic Acute 1

Aquatic Chronic 1

Aquatic Chronic 1
M-factor acute: 100
M-factor chronic: 100

H301, H317, H314, H310 + H330, H400, H410

EUH071

Specific concentration limit:

Skin Corr./Irrit. 1C: >= 0,6 %

Skin Sens. 1A: >= 0,0015 %

Eye Dam./Irrit. 1: >= 0,6 %

Skin Corr./Irrit. 2: 0,06 - < 0,6 %

Eye Dam./Irrit. 2: 0,06 - < 0,6 %

For the classifications not written out in full in this section, including the hazard classes and the hazard statements, the full text is listed in section 16.

SECTION 4: First-Aid Measures

4.1. Description of first aid measures

Remove contaminated clothing.

If inhaled:

If difficulties occur after dust has been inhaled, remove to fresh air and seek medical attention.

On skin contact:

Wash thoroughly with soap and water

On contact with eyes:

Immediately wash affected eyes for at least 15 minutes under running water with eyelids held open, consult an eye specialist.

On ingestion:

Immediately rinse mouth and then drink 200-300 ml of water, seek medical attention.

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

Symptoms: (Further) symptoms and / or effects are not known so far

to Regulation (EC) No 1907/2006.

Date / Revised: 12.04.2023 Version: 1.0

Date previous version: not applicable Previous version: none

Date / First version: 12.04.2023 Product: **Rheovis® HS 1980**

(ID no. 30492472/SDS_GEN_GR/EN)

Date of print 23.10.2025

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

Treatment: Symptomatic treatment (decontamination, vital functions).

SECTION 5: Fire-Fighting Measures

5.1. Extinguishing media

Suitable extinguishing media:

foam, carbon dioxide, dry powder, water spray

5.2. Special hazards arising from the substance or mixture

Advice: No particular hazards known.

5.3. Advice for fire-fighters

Special protective equipment:

Wear a self-contained breathing apparatus.

Further information:

Dispose of fire debris and contaminated extinguishing water in accordance with official regulations.

SECTION 6: Accidental Release Measures

6.1. Personal precautions, protective equipment and emergency procedures

Avoid dust formation. Sources of ignition should be kept well clear.

6.2. Environmental precautions

Do not release untreated into natural waters.

6.3. Methods and material for containment and cleaning up

For small amounts: Pick up in dry form. Dispose of absorbed material in accordance with regulations. For large amounts: Pick up in dry form. Dispose of absorbed material in accordance with regulations.

6.4. Reference to other sections

Information regarding exposure controls/personal protection and disposal considerations can be found in section 8 and 13.

SECTION 7: Handling and Storage

7.1. Precautions for safe handling

Handle in accordance with good industrial hygiene and safety practice.

to Regulation (EC) No 1907/2006.

Version: 1.0 Previous version: none

Date / Revised: 12.04.2023
Date previous version: not applicable
Date / First version: 12.04.2023
Product: Rheovis® HS 1980

(ID no. 30492472/SDS_GEN_GR/EN)

Date of print 23.10.2025

Protection against fire and explosion:

Avoid deposition of dust. Dust can form an explosive mixture with air. Sources of ignition should be kept well clear. Take precautionary measures against static discharges.

7.2. Conditions for safe storage, including any incompatibilities

Further information on storage conditions: Keep container tightly closed and dry; store in a cool place.

7.3. Specific end use(s)

For the relevant identified use(s) listed in Section 1 the advice mentioned in this section 7 is to be observed.

SECTION 8: Exposure Controls/Personal Protection

8.1. Control parameters

Components with occupational exposure limits

No substance specific occupational exposure limits known.

Components with PNEC

55965-84-9: mixture of: 5-chloro-2-methyl-2H-isothiazol-3-one and 2-methyl-2H-isothiazol-3-one

freshwater: 0,00339 mg/l marine water: 0,00339 mg/l

STP: 0,23 mg/l

sediment (freshwater): 0,027 mg/kg sediment (marine water): 0,027 mg/kg

soil: 0,01 mg/kg

Components with DNEL

55965-84-9: mixture of: 5-chloro-2-methyl-2H-isothiazol-3-one and 2-methyl-2H-isothiazol-3-one

worker: Long-term exposure - local effects, Inhalation: 0,02 mg/m3 worker: Short-term exposure - local effects, Inhalation: 0,04 mg/m3 consumer: Short-term exposure - local effects, Inhalation: 0,04 mg/m3 consumer: Long-term exposure - local effects, Inhalation: 0,02 mg/m3 consumer: Long-term exposure- systemic effects, oral: 0,09 mg/kg consumer: Short-term exposure - systemic effects, oral: 0,11 mg/kg

8.2. Exposure controls

Personal protective equipment

Respiratory protection:

Respiratory protection in case of vapour/aerosol release. Particle filter with medium efficiency for solid and liquid particles (e.g. EN 143 or 149, Type P2 or FFP2)

Hand protection:

Date / Revised: 12.04.2023 Version: 1.0

Date previous version: not applicable Previous version: none

Date / First version: 12.04.2023
Product: Rheovis® HS 1980

(ID no. 30492472/SDS_GEN_GR/EN)

Date of print 23.10.2025

Chemical resistant protective gloves (EN ISO 374-1)

Suitable materials also with prolonged, direct contact (Recommended: Protective index 6, corresponding > 480 minutes of permeation time according to EN ISO 374-1):
e.g. nitrile rubber (0.4 mm), chloroprene rubber (0.5 mm), polyvinylchloride (0.7 mm) and other Supplementary note: The specifications are based on tests, literature data and information of glove manufacturers or are derived from similar substances by analogy. Due to many conditions (e.g. temperature) it must be considered, that the practical usage of a chemical-protective glove in practice may be much shorter than the permeation time determined through testing. Manufacturer's directions for use should be observed because of great diversity of types.

Eye protection:

Safety glasses with side-shields (frame goggles) (e.g. EN 166)

General safety and hygiene measures

Handle in accordance with good industrial hygiene and safety practice. Wearing of closed work clothing is recommended. Hands and/or face should be washed before breaks and at the end of the shift. Avoid contact with skin and eyes.

SECTION 9: Physical and Chemical Properties

9.1. Information on basic physical and chemical properties

State of matter: solid Form: powder Colour: white

Odour: pungent odour

Odour threshold:

No data available.

Melting point: 233 °C decomposition point: 310 °C

Flammability: not flammable (derived from flash point)

Lower explosion limit: 20 g/m3

Upper explosion limit:

For liquids not relevant for classification and labelling.

Flash point:

No data available.

Auto-ignition temperature: > 200 °C

Thermal decomposition: No decomposition if used correctly.

pH value: 4,5 - 6,0 (DIN ISO 976)

(water, 30 %(m))

Viscosity, dynamic:

not applicable, the product is a solid

Solubility in water: soluble

Solubility (qualitative):

insoluble

Partitioning coefficient n-octanol/water (log Kow):

No data available.

to Regulation (EC) No 1907/2006.

Date / Revised: 12.04.2023 Version: 1.0

Date previous version: not applicable Previous version: none

Date / First version: 12.04.2023 Product: **Rheovis® HS 1980**

(ID no. 30492472/SDS_GEN_GR/EN)

Date of print 23.10.2025

Vapour pressure:

No data available.

Relative density:

No data available.

Density:

No data available.

Relative vapour density (air):

No data available.

Particle characteristics

Particle size distribution: No data available. -

9.2. Other information

Information with regard to physical hazard classes

Explosives

Explosion hazard: not explosive

Oxidizing properties

Fire promoting properties: not fire-propagating

Self-heating substances and mixtures

Self heating ability: It is not a substance capable of

spontaneous heating.

Other safety characteristics

Minimum ignition energy: > 30 - < 100 mJ (DIN EN 13821)

Inductivity: 0,1 mH

Bulk density: 300 - 500 kg/m3 (DIN EN ISO 60)

(20 °C)

Miscibility with water:

completely (e.g. >=90%)

Solids content: >= 97,0 % (DIN EN ISO 3251)

SAPT-Temperature:

Product does not fulfil criteria for polymerizing substances according to

transport regulations.

Evaporation rate:

The product is a non-volatile solid.

SECTION 10: Stability and Reactivity

10.1. Reactivity

No hazardous reactions if stored and handled as prescribed/indicated.

10.2. Chemical stability

The product is stable if stored and handled as prescribed/indicated.

to Regulation (EC) No 1907/2006.

Date / Revised: 12.04.2023 Version: 1.0

Date previous version: not applicable Previous version: none

Date / First version: 12.04.2023 Product: **Rheovis® HS 1980**

(ID no. 30492472/SDS_GEN_GR/EN)

Date of print 23.10.2025

10.3. Possibility of hazardous reactions

Dust explosion hazard.

10.4. Conditions to avoid

Avoid extreme temperatures. Avoid dust formation.

10.5. Incompatible materials

Substances to avoid:

No substances known that should be avoided.

10.6. Hazardous decomposition products

Hazardous decomposition products:

No data available.

SECTION 11: Toxicological Information

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

Acute toxicity

Assessment of acute toxicity:

Virtually nontoxic after a single ingestion. Virtually nontoxic after a single skin contact. Virtually nontoxic by inhalation. The product has not been tested. The statement has been derived from the properties of the individual components.

Experimental/calculated data:

ATE other (oral): > 5.000 mg/kg (calculated) ATE (dermal): > 5.000 mg/kg (calculated)

<u>Irritation</u>

Assessment of irritating effects:

Not irritating to eyes and skin. The product has not been tested. The statement has been derived from the properties of the individual components.

Experimental/calculated data:

Skin corrosion/irritation

rabbit:

not determined

Serious eye damage/irritation

rabbit:

not determined

to Regulation (EC) No 1907/2006.

Date / Revised: 12.04.2023 Version: 1.0

Date previous version: not applicable Previous version: none

Date / First version: 12.04.2023 Product: **Rheovis® HS 1980**

(ID no. 30492472/SDS_GEN_GR/EN)

Date of print 23.10.2025

Respiratory/Skin sensitization

Assessment of sensitization:

A sensitizing effect on particularly sensitive individuals cannot be excluded. The product has not been tested. The statement has been derived from the properties of the individual components.

Experimental/calculated data:

not determined

Germ cell mutagenicity

Assessment of mutagenicity:

The chemical structure does not suggest a specific alert for such an effect.

Carcinogenicity

Assessment of carcinogenicity:

The whole of the information assessable provides no indication of a carcinogenic effect.

Reproductive toxicity

Assessment of reproduction toxicity:

Not expected to cause reproductive toxicity (based on composition).

Developmental toxicity

Assessment of teratogenicity:

Based on the ingredients, there is no suspicion of a teratogenic effect.

Specific target organ toxicity (single exposure)

Remarks: No data available.

Repeated dose toxicity and Specific target organ toxicity (repeated exposure)

Experimental/calculated data:

Not expected to cause chronic toxic effects.

Aspiration hazard

No aspiration hazard expected.

Interactive effects

No data available.

11.2. Information on other hazards

Endocrine disrupting properties

Product does not contain a substance above legal limits included in the list established in accordance with Article 59(1) of Regulation (EC) No 1907/2006 for having endocrine disrupting properties or is identified to have endocrine disrupting properties in accordance with the criteria set out in Commission Delegated Regulation (EU) 2017/2100 or Commission Regulation (EU) 2018/605.

to Regulation (EC) No 1907/2006.

Version: 1.0 Previous version: none

Date / Revised: 12.04.2023
Date previous version: not applicable
Date / First version: 12.04.2023
Product: **Rheovis® HS 1980**

(ID no. 30492472/SDS_GEN_GR/EN)

Date of print 23.10.2025

Other information

Other relevant toxicity information

Based on our experience and the information available, no adverse health effects are expected if handled as recommended with suitable precautions for designated uses.

SECTION 12: Ecological Information

12.1. Toxicity

Assessment of aquatic toxicity:

There is a high probability that the product is not acutely harmful to aquatic organisms.

Toxicity to fish: LC50 (96 h), Fish (other) not determined

Aquatic invertebrates: LC50 (48 h), daphnia (other) not determined

Aquatic plants: EC50 (72 h), algae (other) not determined

Microorganisms/Effect on activated sludge: EC50 (0,5 h), bacteria (other) not determined

Chronic toxicity to fish: No data available.

Chronic toxicity to aquatic invertebrates: No data available.

Assessment of terrestrial toxicity: No data available concerning terrestrial toxicity.

12.2. Persistence and degradability

Assessment biodegradation and elimination (H2O):

The product can be virtually eliminated from water by abiotic processes e.g. adsorption onto activated sludge.

to Regulation (EC) No 1907/2006.

Date / Revised: 12.04.2023 Version: 1.0

Date previous version: not applicable Previous version: none

Date / First version: 12.04.2023 Product: **Rheovis® HS 1980**

(ID no. 30492472/SDS_GEN_GR/EN)

Date of print 23.10.2025

12.3. Bioaccumulative potential

Assessment bioaccumulation potential: The product has not been tested.

12.4. Mobility in soil

Assessment transport between environmental compartments: Volatility: No data available.

12.5. Results of PBT and vPvB assessment

According to Annex XIII of Regulation (EC) No.1907/2006 concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals (REACH): The product does not contain a substance fulfilling the PBT (persistent/bioaccumulative/toxic) criteria or the vPvB (very persistent/very bioaccumulative) criteria.

12.6. Endocrine disrupting properties

Product does not contain a substance above legal limits included in the list established in accordance with Article 59(1) of Regulation (EC) No 1907/2006 for having endocrine disrupting properties or is identified to have endocrine disrupting properties in accordance with the criteria set out in Commission Delegated Regulation (EU) 2017/2100 or Commission Regulation (EU) 2018/605.

12.7. Other adverse effects

The product does not contain substances that are listed in Regulation (EC) 1005/2009 on substances that deplete the ozone layer.

12.8. Additional information

Add. remarks environm. fate & pathway:

Treatment in biological waste water treatment plants has to be performed according to local and administrative regulations.

Other ecotoxicological advice:

Do not release untreated into natural waters. The inhibition of the degradation activity of activated sludge is not anticipated when introduced to biological treatment plants in appropriate low concentrations. The local regulations on waste-water treatment must be followed.

to Regulation (EC) No 1907/2006.

Date / Revised: 12.04.2023 Version: 1.0

Date previous version: not applicable Previous version: none

Date / First version: 12.04.2023 Product: **Rheovis® HS 1980**

(ID no. 30492472/SDS_GEN_GR/EN)

Date of print 23.10.2025

SECTION 13: Disposal Considerations

13.1. Waste treatment methods

Must be sent to a suitable incineration plant, observing local regulations.

A waste code in accordance with the European waste catalog (EWC) cannot be specified, due to dependence on the usage.

Observe national and local legal requirements.

SECTION 14: Transport Information

Land transport

ADR

Not classified as a dangerous good under transport regulations

UN number or ID number:
UN proper shipping name:
Transport hazard class(es):
Packing group:
Environmental hazards:

Not applicable
Not applicable
Not applicable
Not applicable

Special precautions for

user

None known

RID

Not classified as a dangerous good under transport regulations

UN number or ID number: Not applicable UN proper shipping name: Not applicable Transport hazard class(es): Not applicable Packing group: Not applicable

Environmental hazards: Not applicable Special precautions for None known

user

Inland waterway transport

ADN

Not classified as a dangerous good under transport regulations

UN number or ID number:
UN proper shipping name:
Transport hazard class(es):
Packing group:
Environmental hazards:
Special precautions for

Not applicable
Not applicable
Not applicable
Not applicable
Not applicable

user:

to Regulation (EC) No 1907/2006.

Date / Revised: 12.04.2023 Version: 1.0

Date previous version: not applicable Previous version: none

Date / First version: 12.04.2023 Product: **Rheovis® HS 1980**

(ID no. 30492472/SDS_GEN_GR/EN)

Date of print 23.10.2025

Transport in inland waterway vessel

Not evaluated

Sea transport

IMDG

Not classified as a dangerous good under transport regulations

UN number or ID number:
UN proper shipping name:
Transport hazard class(es):
Packing group:
Environmental hazards:
Special precautions for

Not applicable
Not applicable
Not applicable
Not applicable
Not applicable

user

Air transport

IATA/ICAO

Not classified as a dangerous good under transport regulations

UN number or ID number:
UN proper shipping name:
Transport hazard class(es):
Packing group:
Environmental hazards:
Special precautions for

Not applicable
Not applicable
Not applicable
Not applicable
Not applicable
Not applicable

user

14.1. UN number or ID number

See corresponding entries for "UN number or ID number" for the respective regulations in the tables above.

14.2. UN proper shipping name

See corresponding entries for "UN proper shipping name" for the respective regulations in the tables above.

14.3. Transport hazard class(es)

See corresponding entries for "Transport hazard class(es)" for the respective regulations in the tables above.

14.4. Packing group

See corresponding entries for "Packing group" for the respective regulations in the tables above.

to Regulation (EC) No 1907/2006.

Date / Revised: 12.04.2023 Version: 1.0

Date previous version: not applicable Previous version: none

Date / First version: 12.04.2023 Product: **Rheovis® HS 1980**

(ID no. 30492472/SDS_GEN_GR/EN)

Date of print 23.10.2025

14.5. Environmental hazards

See corresponding entries for "Environmental hazards" for the respective regulations in the tables above.

14.6. Special precautions for user

See corresponding entries for "Special precautions for user" for the respective regulations in the tables above.

14.7. Maritime transport in bulk according to IMO instruments

Maritime transport in bulk is not intended.

SECTION 15: Regulatory Information

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

Prohibitions, Restrictions and Authorizations

Annex XVII of Regulation (EC) No 1907/2006: Number on List: 75

Directive 2012/18/EU - Control of Major Accident Hazards involving dangerous substances (EU): Listed in above regulation: no

If other regulatory information applies that is not already provided elsewhere in this safety data sheet, then it is described in this subsection.

15.2. Chemical Safety Assessment

Chemical Safety Assessment not required

SECTION 16: Other Information

Any other intended applications should be discussed with the manufacturer.

Full text of the classifications, including the hazard classes and the hazard statements, if mentioned

in section 2 or 3:

Acute Tox. Acute toxicity

Skin Corr./Irrit. Skin corrosion/irritation

Eye Dam./Irrit. Serious eye damage/eye irritation

Skin Sens. Skin sensitization

Aquatic Acute Hazardous to the aquatic environment - acute Aquatic Chronic Hazardous to the aquatic environment - chronic

H301 Toxic if swallowed.

H317 May cause an allergic skin reaction.

H314 Causes severe skin burns and eye damage.

H310 + H330 Fatal in contact with skin or if inhaled

H400 Very toxic to aquatic life.

to Regulation (EC) No 1907/2006. Date / Revised: 12.04.2023

Version: 1.0 Previous version: none

Date previous version: not applicable
Date / First version: 12.04.2023
Product: Rheovis® HS 1980

(ID no. 30492472/SDS_GEN_GR/EN)

Date of print 23.10.2025

Very toxic to aquatic life with long lasting effects.EUH071 Corrosive to the respiratory tract.

Abbreviations

ADR = The European Agreement concerning the International Carriage of Dangerous Goods by Road. ADN = The European Agreement concerning the International Carriage of Dangerous Goods by Inland waterways. ATE = Acute Toxicity Estimates. CAO = Cargo Aircraft Only. CAS = Chemical Abstract Service. CLP = Classification, Labelling and Packaging of substances and mixtures. DIN = German national organization for standardization. DNEL = Derived No Effect Level. EC50 = Effective concentration median for 50% of the population. EC = European Community. EN = European Standards. IARC = International Agency for Research on Cancer. IATA = International Air Transport Association. IBC-Code = Intermediate Bulk Container code. IMDG = International Maritime Dangerous Goods Code. ISO = International Organization for Standardization. STEL = Short-Term Exposure Limit. LC50 = Lethal concentration median for 50% of the population. LD50 = Lethal dose median for 50% of the population. TLV = Threshold Limit Value. MARPOL = The International Convention for the Prevention of Pollution from Ships. NEN = Dutch Norm. NOEC = No Observed Effect Concentration. OEL = Occupational Exposure Limit. OECD = Organization for Economic Cooperation and Development. PBT = Persistent, Bioaccumulative and Toxic. PNEC = Predicted No Effect Level. PPM = Parts per million. RID = The European Agreement concerning the International Carriage of Dangerous Goods by Rail, TWA = Time Weight Average. UN-number = UN number at transport. vPvB = very Persistent and very Bioaccumulative.

The data contained in this safety data sheet are based on our current knowledge and experience and describe the product only with regard to safety requirements. This safety data sheet is neither a Certificate of Analysis (CoA) nor technical data sheet and shall not be mistaken for a specification agreement. Identified uses in this safety data sheet do neither represent an agreement on the corresponding contractual quality of the substance/mixture nor a contractually designated use. It is the responsibility of the recipient of the product to ensure any proprietary rights and existing laws and legislation are observed.

Vertical lines in the left hand margin indicate an amendment from the previous version.