Product Name Tert BUTYL METHYL ETHER AR 99.5%

Part Number RXSOL-60-6605-659

Company Details:

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### 2. Composition / Information on ingredients

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Chemical Name CAS EC number Weight Formula

216-653-1 Tert-Butyl methyl ether 1634-04-4 2.000 mg/kg (OECD Test

Guideline 401) Symptoms: Nausea, Vomiting, Pulmonary failure possible after aspiration of vomit., Aspiration may cause pulmonary edema and pneumonitis. LC50 Inhalation - Rat - male and female - 4 h - 85 mg/l vapor (OECD Guideline 403) Symptoms: Possible damages:, mucosal irritations LD50 Dermal -Rat - male and female - > 2.000 mg/kg (OECD Test Guideline 402)

Skin corrosion/irritation Skin - Rabbit Result: Skin

irritation - 4 h (OECD Test Guideline 404) Remarks: Drying-out effect resulting in rough and chapped skin.

damage/eye Eyes - Rabbit Result: No eye Serious eye

irritation irritation (OECD Test

Guideline 405)

Respiratory or

skin Maximization Test - Guinea sensitization

pig Result: negative (OECD

Test Guideline 406)

WARNING: THE LC50 Toxicity to Animal

VALUES HEREUNDER ARE ESTIMATED ON THE Acute oral toxicity (LD50): 689 mg/kg [Rat].

Toxic Effects on Human May cause damage to the

> following organs: upper respiratory tract. Other Toxic Effects on Humans: Hazardous in case of skin

contact (irritant, sensitizer), of ingestion, of inhalation lung (lung irritant, sensitizer).

Carcinogenicity

Carcinogenicity No carcinogenic properties suspected. (IUCLID)

Germ cell mutagenicity

No data available Test Type: In vitro mammalian cell gene mutation test Test system: Chinese hamster lung cells Metabolic activation: with without metabolic and activation Method: OECD Test Guideline 476 Result: negative Test Type: Ames test Test system: Salmonella typhimurium Metabolic activation: with and without metabolic activation **OECD** Method: Test 471 Guideline Result: negative Test Type: Mutagenicity (mammal cell test): micronucleus. Test system: mouse lymphoma cells Metabolic activation: without metabolic activation Method: **OECD** Test 473 Guideline Result: Test negative Type: unscheduled DNA synthesis assay Species: Mouse Cell type: Liver cells Application Route: inhalation (vapor) Method: **OECD** Test 486 Guideline Result: Test negative Type: Micronucleus test Species: Mouse Cell type: Bone marrow Application Route: inhalation (vapor) Method: US-EPA Result: negative Test Type: Mutagenicity (mammal cell test): chromosome aberration. Species: Rat Cell type: Bone marrow Application Route: inhalation (vapor) Method: US-EPA Result: negative Test Type: Transgenic rodent somatic cell gene mutation assay Species: Rat Cell type: Bone marrow Application Route: inhalation (vapor) Method: OECD Test Guideline 488 Result: negative

Reproductive toxicity

No data available

Specific target organ toxicity No data available

- single exposure

Special Remarks on other Acute Toxi Effects on Humans

Potential Health Effects: Skin: Causes skin irritation. May cause skin sensitization, an allergic reation, which becomes evident upon re-exposure to this material. Eyes: Causes eye irritation. Ingestion: Causes gastrointestinal (digestive) tract irritation with nausea, vomiting, and diarrhea. May be harmful if swallowed. Inhalation: Causes respiratory tract irritation. May cause chemical pneumonitis and pulmonary edema, inflammation, edema of bronchi and larynx. Chronic Potential Health Effects: Repeated or prolonged skin exposure may cause allergic reactions in sensitive individuals. Repeated or prolonged exposure by inhalation may affect respiration and metabolism.

Additional Information

RTECS: AF3675000 Gastrointestinal disturbance To the best of our knowledge, the chemical, physical, and toxicological properties have not been thoroughly investigated. After uptake of large quantities: muscular symptoms agitation Convulsions Headache Tremors Nausea psychoses

Endocrine disrupting properties

disrupting 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. Repeated dose toxicity - Rat - male and female - Oral - 90 d - NOAEL (No observed adverse effect level) - 3.000 mg/kg Remarks: Subchronic toxicity RTECS: KN5250000 Nausea, Vomiting, Dizziness, Central nervous system depression, Aspiration or inhalation may cause chemical pneumonitis., **MTBE** (methyl-tert-butyl ether) is reported to metabolize to tertbutyl alcohol formaldehyde microsomal demethylation, **MTBE** (methyl-tert?butyl ether) should be considered "potential human carcinogen" due to

increase in leydig interstitial cell tumors of testes in male rats and an increase in lymphomas, leukemias, and uterine sarcomas in female rats., In another unpublished study MTBE was shown to be carcinogenic due to "increased incidence of a rare type of kidney tumor" in male rats and an "increase the incidence hepatocellular adenomas" in female mice. To the best of knowledge, chemical, physical, and toxicological properties have thoroughly not been investigated.

## 12. Ecological information

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Persistence and degradability

Bioaccumulative potential

Toxicity to fish

Toxicity to daphnia and other aquatic invertebrates

Toxicity to bacteria

Toxicity to algae

**Products Biodegradation** 

Mobility in soil

Results of PBT and vPvB assessment

Endocrine disrupting properties

Other adverse effects

Biodegradability aerobic - Exposure time 28 d Result: 0% - Not readily biodegradable. (OECD Test Guideline 301D)

Bioaccumulation Cyprinus carpio (Carp) - 28 d at 25 °C(tert-butyl methyl ether) Bioconcentration factor (BCF): 1,5

Semi-static test LC50 - Menidia beryllina - 574 mg/l - 96 h (OECD Test Guideline 203)

Flow-through test EC50 - Americamysis bahia (Mysid) - 187 mg/l - 96 h (US-EPA OPPTS 850.1035)

Static test EC10 - Pseudomonas putida - 710 mg/l - 18 h Remarks: (ECHA)

Static test IC50 - Pseudokirchneriella subcapitata (green algae) - 491 mg/l - 96 h

The methods for determining the biological degradability are not applicable to inorganic substances.

No Information available

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.

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.

No data available

# 13. Disposal considerations

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Disposal methods

Contaminated packaging

Waste treatment Method

The material can be disposed of by removal to a licensed chemical destruction plant or by controlled incineration with flue gas scrubbing. Do not contaminate water, foodstuffs, feed or seed by storage or disposal. Do not discharge to sewer systems.

Containers can be triply rinsed (or equivalent) and offered for recycling or reconditioning. Alternatively, the packaging can be punctured to make it unusable for other purposes and then be disposed of in a sanitary landfill. Controlled incineration with flue gas scrubbing is possible for combustible packaging materials.

See www.retrologistik.com for processes regarding the return of chemicals and containers, or contact us there if you have further

# 14. Transport information

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UN number ADR/RID:2398 IMDG:2398 IATA:2398

UN proper shipping name ADR/RID:Methyl tert-butyl ether IMDG:Methyl tert-butyl ether

IATA:Methyl tert-butyl ether

Transport hazard class(es)

ADR/RID: 3 IMDG: 3 IATA: 3

Packaging group

ADR/RID: II IMDG: II IATA: II

Environmental hazards

ADR/RID: No IMDG: No IATA: No

Special precautions for user Tunnel restriction code: (C/E)

Further information Not classified as dangerous in the meaning of transport regulations.

No data available

### 15. Regulatory information

Transport in bulk according to IMOinstruments

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Inventory status

proposition 65.

Safety, health and environmental regulations/legislation specific for the This material safety data sheet complies with the requirements of

Safety, health and environmental regulations specific for the product in European Inventory of Existing Commercial Chemical Substances question (EINECS) -Listed.EC Inventory Listed. United States Toxic Substances

National legislation

substance or mixture

Other regulations Chemical Safety Assessment Components are on the following inventories: Polymaleic acid: - US TSCA, Canadian DSL, EU EINECS, Australian AICS, Korean, Philippine PICCS and Chinese Xi irritant R 36/38 Irritant to eyes & skin R 41 Risk of serious damage to eyes S24/25 Avoid contact with skin and eyes S26/28 In case of contact eyes & skin, rinse with plenty water and seek medical advice Section 312/313: Not listed. Not listed under California proposition 65.

This material safety data sheet complies with the requirements of Regulation (EC) No. 1907/2006.

European Inventory of Existing Commercial Chemical Substances (EINECS) -Listed.EC Inventory Listed. United States Toxic Substances Control Act (TSCA) Inventory- Listed. China Catalog of Hazardous chemicals 2015 Not- Listed. New Zealand Inventory of Chemicals (NZIoC)Listed. PICCS -Listed. Vietnam National Chemical Inventory -Listed. IECSC Listed. Korea Existing Chemicals List (KECL) -Listed.

Seveso III: Directive 2012/18/EU of the European Parliament and of the Council on the control of major-accident hazards involving dangerous substances. : FLAMMABLE LIQUIDS

Take note of Dir 94/33/EC on the protection of young people at work. A Chemical Safety Assessment has been carried out for this substance.

#### 16. Other information

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Other Information

The information above is believed to be accurate and represents the best information currently available to us. However, we make no warranty of merchantability or any other warranty, express or implied, with respect to such information, and weassume no liability resulting from its use. Users should make their own investigations to determine the suitability of the information for their particular purposes. In no event shall be liable for any claims, losses, or damages of any third party or for lost profits or any special, indirect, incidental, consequential or exemplary damages, howsoever arising, even if Rx Marine International has been advised of the possibility of such damages.

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