

SAFETY DATA SHEET

SECTION 1. PRODUCT AND COMPANY IDENTIFICATION

Product identifier	Slow Urethane Reducer	
Other means of identification	ADV 116-5	
Product code		
Recommended use	Solvent	

Manufacturer/Importer/Supplier/Distributor information

Company name Address	INTERNATIONAL AUTOBODY MARKETING GROUP 1505 NORTH HAYDEN RD, SUITE 111 SCOTTSDALE, AZ 85257 UNITED STATES
Website	www.advantagerefinish.com
Telephone	1-87-REFINISH 480.451.4451

Emergency phone number 800-424-9300 ChemTrec EMERGENCY 24 Hrs.

SECTION 2. HAZARDS IDENTIFICATION

GHS Classification	
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Flammable liquids	: Category 2
Skin irritation	: Category 2
Eye irritation	: Category 2A
Germ cell mutagenicity	: Category 1B
Carcinogenicity	: Category 2
Reproductive toxicity	: Category 2
Specific target organ tox- icity - single exposure	: Category 3 (Central nervous system)
Specific target organ tox- icity - repeated exposure	: Category 2 (Liver, Kidney, Central nervous system, Au- ditory system)
Specific target organ tox- icity - repeated exposure (Inhalation)	: Category 2 (Auditory system, Eyes)

Aspiration hazard	: Category 1
GHS Label element Hazard pictograms	
Signal word	: Danger
Hazard statements	 H225 Highly flammable liquid and vapour. H304 May be fatal if swallowed and enters airways. H315 Causes skin irritation. H319 Causes serious eye irritation. H336 May cause drowsiness or dizziness. H340 May cause genetic defects. H351 Suspected of causing cancer. H361 Suspected of damaging fertility or the unborn child. H373 May cause damage to organs (Liver, Kidney, Central nervous system, Auditory system) through prolonged or repeated exposure. H373 May cause damage to organs (Auditory system, Eyes) through prolonged or repeated exposure if inhaled.
Precautionary statements	 Prevention: P201 Obtain special instructions before use. P202 Do not handle until all safety precautions have been read and understood. P210 Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking. P233 Keep container tightly closed. P240 Ground/bond container and receiving equipment. P241 Use explosion-proof electrical/ ventilating/ lighting/ equipment. P242 Use only non-sparking tools. P243 Take precautionary measures against static discharge. P260 Do not breathe dust/ fume/ gas/ mist/ vapours/ spray. P264 Wash skin thoroughly after handling. P271 Use only outdoors or in a well-ventilated area. P280 Wear protective gloves/ eye protection/ face protection. P281 Use personal protective equipment as required. Response: P301 + P310 IF SWALLOWED: Immediately call a POISON CENTER or doctor/ physician.

P303 + P361 + P353 IF ON SKIN (or hair): Remove/ Take off immediately all contaminated clothing. Rinse skin with water/ shower.

P304 + P340 + P312 IF INHALED: Remove victim to fresh air and keep at rest in a position comfortable for breathing. Call a POISON CENTER or doctor/ physician if you feel unwell.

P305 + P351 + P338 IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.

P308 + P313 IF exposed or concerned: Get medical advice/ attention.

P331 Do NOT induce vomiting.

P332 + P313 If skin irritation occurs: Get medical advice/ attention.

P337 + P313 If eye irritation persists: Get medical advice/ attention.

P362 Take off contaminated clothing and wash before reuse.

P370 + P378 In case of fire: Use dry sand, dry chemical or alcohol-resistant foam for extinction.

Storage:

P403 + P233 Store in a well-ventilated place. Keep container tightly closed.

P403 + P235 Store in a well-ventilated place. Keep cool. P405 Store locked up.

Disposal:

P501 Dispose of contents/ container to an approved waste disposal plant.

Potential Health Effects

Carcinogenicity:		
IARC	Group 2B: Possibly carcinogenic to humans	
	64742-49-0	Naphtha (pet), hydrotreated It
	64742-89-8	Solvent naphtha (pet), lt aliph.
	100-41-4	Ethylbenzene
ACGIH	No component of this produ than or equal to 0.1% is ide potential carcinogen by ACC	5
OSHA	No component of this prodution that the product of the product of the product of the potential carcinogen by OSI	

NTP

No component of this product present at levels greater than or equal to 0.1% is identified as a known or anticipated carcinogen by NTP.

Emergency Overview

Appearance	liquid
Colour	clear
Hazard Summary	No information available.

SECTION 3. COMPOSITION/INFORMATION ON INGREDIENTS

Substance / Mixture : Mixture

Hazardous components

CAS-No.	Chemical Name	Concentration (%)
67-64-1	Acetone	30 - 50
64742-49-0	Naphtha (pet), hydrotreated It	0 - 20
64742-89-8	Solvent naphtha (pet), lt aliph.	0 - 20
68410-97-9	Distillates, pet, lt dist hydrotreat process, low-boil	0 - 20
108-65-6	Glycol ether PM acetate	10 - 20
108-88-3	Toluene	10 - 20
110-19-0	Isobutyl acetate	10 - 20
123-86-4	n-Butyl acetate	5 - 10
1330-20-7	Mixed xylenes	5 - 10
100-41-4	Ethylbenzene	1 - 5
142-82-5	Heptane	0.1 - 1

Synonyms

: CP 81-03,

Special Notes: : Functionally equivalent petroleum streams may be found in this preparation at varying concentrations. Mixed Xylenes contains the isomers o-, m-, p- Xylene, and Ethylbenzene. Trace amounts of Toluene and Benzene may also be present as impurities.

SECTION 4. FIRST AID MEASURES

General advice	: Move out of dangerous area.
	Show this safety data sheet to the doctor in attend-
	ance.
	Symptoms of poisoning may appear several hours

If inhaled	later.Do not leave the victim unattended.Consult a physician after significant exposure.If unconscious place in recovery position and seek medical advice.
In case of skin contact	 If skin irritation persists, call a physician. If on skin, rinse well with water. If on clothes, remove clothes.
In case of eye contact	 Immediately flush eye(s) with plenty of water. Remove contact lenses. Protect unharmed eye. Keep eye wide open while rinsing. If eye irritation persists, consult a specialist.
If swallowed	 Keep respiratory tract clear. Do NOT induce vomiting. Do not give milk or alcoholic beverages. Never give anything by mouth to an unconscious person. If symptoms persist, call a physician. Take victim immediately to hospital.

SECTION 5. FIREFIGHTING MEASURES

Suitable extinguishing media	: Alcohol-resistant foam Carbon dioxide (CO2) Dry chemical
Unsuitable extinguishing media	: High volume water jet
Specific hazards during firefighting	: Do not allow run-off from fire fighting to enter drains or water courses.
Hazardous combustion products	: No hazardous combustion products are known
Specific extinguishing methods	: Use a water spray to cool fully closed containers.
Further information	: Collect contaminated fire extinguishing water sepa- rately. This must not be discharged into drains. Fire residues and contaminated fire extinguishing wa- ter must be disposed of in accordance with local regu- lations.

For safety reasons in case of fire, cans should be
stored separately in closed containments.Special protective equip-
ment for firefighters: Wear self-contained breathing apparatus for fire-
fighting if necessary.

NFPA Flammable and Combustible Liquids Classification:

Flammable Liquid Class IB

SECTION 6. ACCIDENTAL RELEASE MEASURES

Personal precautions, protective equipment and emergency procedures	:	Use personal protective equipment. Ensure adequate ventilation. Remove all sources of ignition. Evacuate personnel to safe areas. Beware of vapours accumulating to form explosive concentrations. Vapours can accumulate in low areas.
Environmental precau- tions	:	Prevent product from entering drains. Prevent further leakage or spillage if safe to do so. If the product contaminates rivers and lakes or drains inform respective authorities.
Methods and materials for containment and cleaning up	:	Contain spillage, and then collect with non- combustible absorbent material, (e.g. sand, earth, diatomaceous earth, vermiculite) and place in con- tainer for disposal according to local / national regula- tions (see section 13).

SECTION 7. HANDLING AND STORAGE

Advice on safe handling	 Avoid formation of aerosol. Do not breathe vapours/dust. Avoid exposure - obtain special instructions before use. Avoid contact with skin and eyes. For personal protection see section 8. Smoking, eating and drinking should be prohibited in the application area. Take precautionary measures against static discharg- es. Provide sufficient air exchange and/or exhaust in work rooms. Container may be opened only under exhaust ventila- tion hood. Open drum carefully as content may be under pres- sure.
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	Dispose of rinse water in accordance with local and national regulations.
Conditions for safe stor- age	 No smoking. Keep container tightly closed in a dry and well- ventilated place. Containers which are opened must be carefully re- sealed and kept upright to prevent leakage. Observe label precautions. Electrical installations / working materials must com- ply with the technological safety standards.

SECTION 8. EXPOSURE CONTROLS/PERSONAL PROTECTION

CAS-No.	Components	Value type (Form of exposure)	Control parame- ters / Permissi- ble concentra- tion	Basis
67-64-1	Acetone	TWA	500 ppm	ACGIH
		STEL	750 ppm	ACGIH
		TWA	250 ppm 590 mg/m3	NIOSH REL
		TWA	1,000 ppm 2,400 mg/m3	OSHA Z-1
		TWA	750 ppm 1,800 mg/m3	OSHA PO
		STEL	1,000 ppm 2,400 mg/m3	OSHA PO
64742-49-0	Naphtha (pet), hydrotreat- ed It	TWA	500 ppm 2,000 mg/m3	OSHA Z-1
		TWA	400 ppm 1,600 mg/m3	OSHA PO
64742-89-8	Solvent naphtha (pet), lt aliph.	TWA	500 ppm 2,000 mg/m3	OSHA Z-1
		TWA	400 ppm 1,600 mg/m3	OSHA PO
108-65-6	Glycol ether PM acetate	TWA	50 ppm	US WEEL
108-88-3	Toluene	TWA	20 ppm	ACGIH
		TWA	100 ppm 375 mg/m3	NIOSH REL
		ST	150 ppm 560 mg/m3	NIOSH REL
		TWA	200 ppm	OSHA Z-2
		CEIL	300 ppm	OSHA Z-2
		Peak	500 ppm	OSHA Z-2

Components with workplace control parameters

		TWA	100 ppm 375 mg/m3	OSHA PO
		STEL	150 ppm 560 mg/m3	OSHA PO
110-19-0	Isobutyl acetate	TWA	150 ppm	ACGIH
		TWA	150 ppm 700 mg/m3	NIOSH REL
		TWA	150 ppm 700 mg/m3	OSHA Z-1
		TWA	150 ppm 700 mg/m3	OSHA PO
123-86-4	n-Butyl acetate	TWA	150 ppm	ACGIH
		STEL	200 ppm	ACGIH
		ST	200 ppm 950 mg/m3	NIOSH REL
		TWA	150 ppm 710 mg/m3	NIOSH REL
		TWA	150 ppm 710 mg/m3	OSHA Z-1
		TWA	150 ppm 710 mg/m3	OSHA PO
		STEL	200 ppm 950 mg/m3	OSHA PO
1330-20-7	Mixed xylenes	TWA	100 ppm	ACGIH
		STEL	150 ppm	ACGIH
		TWA	100 ppm 435 mg/m3	OSHA Z-1
100-41-4	Ethylbenzene	TWA	20 ppm	ACGIH
-		STEL	125 ppm	ACGIH
		TWA	100 ppm 435 mg/m3	NIOSH REL
		ST	125 ppm 545 mg/m3	NIOSH REL
		TWA	100 ppm 435 mg/m3	OSHA Z-1
		TWA	100 ppm 435 mg/m3	OSHA PO
		STEL	125 ppm 545 mg/m3	OSHA PO
142-82-5	Heptane	TWA	85 ppm 350 mg/m3	NIOSH REL
		С	440 ppm 1,800 mg/m3	NIOSH REL
		TWA	500 ppm 2,000 mg/m3	OSHA Z-1
		TWA	400 ppm 1,600 mg/m3	OSHA PO
		STEL	500 ppm 2,000 mg/m3	OSHA PO

Components	CAS-No.	Control parame- ters	Biological specimen	Sam- pling time	Permissi- ble con- centration	Basis
Acetone	67-64-1	Acetone	Urine	End of shift (As soon as possible after expo- sure ceases)	50 mg/l	ACGIH BEI
Toluene	108-88- 3	Toluene	In blood	Prior to last shift of work- week	0.02 mg/l	ACGIH BEI
		Toluene	Urine	End of shift (As soon as possible after expo- sure ceases)	0.03 mg/l	ACGIH BEI
		o-Cresol	Urine	End of shift (As soon as possible after expo- sure ceases)	0.3 mg/g Creatinine	ACGIH BEI
Ethylbenzene	100-41- 4	Sum of mandelic acid and phenyl glyoxylic acid	Urine	End of shift at end of work- week	0.7 g/g creatinine	ACGIH BEI

Biological occupational exposure limits

Personal protective equipment

Respiratory protection : No personal respiratory protective equipment normally required. In the case of vapour formation use a respirator with an approved filter.

Hand protection

Remarks	: The suitability for a specific workplace should be dis- cussed with the producers of the protective gloves.
Eye protection	 Eye wash bottle with pure water Tightly fitting safety goggles Wear face-shield and protective suit for abnormal pro- cessing problems.
Skin and body protection	: impervious clothing Choose body protection according to the amount and concentration of the dangerous substance at the work place.
Hygiene measures	: When using do not eat or drink. When using do not smoke. Wash hands before breaks and at the end of workday.

SECTION 9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance	: liquid	
Colour	: clear	
Odour	: No data available	
Odour Threshold	: No data available	
рН	: No data available	
Freezing Point	: No data available	
Boiling Point (Boiling point/boiling range)	: 56 - 245 °C (133 - 473 °F)	
Flash point	: < -18 °C (-0.40 °F)	
Evaporation rate	: 1 Ethyl Ether	
Flammability (solid, gas)	: No data available	
Burning rate	: No data available	
Upper explosion limit	: 12.8 %(V) Calculated Explosive Limit	
Lower explosion limit	: 1 %(V) Calculated Explosive Limit	

Vapour pressure	: No data available
Relative vapour density	: > 1(Air = 1.0)
Relative density	: 0.827 @ 77.00 °F (77.00 °F)
Density	: 0.827 g/cm3 @ 25 °C (77 °F)
Bulk density	: No data available
Water solubility	: No data available
Solubility in other sol- vents	: No data available
Partition coefficient: n- octanol/water	: No data available
Auto-ignition temperature	: No data available
Thermal decomposition	: No data available

SECTION 10. STABILITY AND REACTIVITY

Reactivity	: No dangerous reaction known under conditions of normal use.
Chemical stability	: Stable under normal conditions.
Possibility of hazardous reactions	: Product will not undergo hazardous polymerization. Vapours may form explosive mixture with air.
Conditions to avoid	 Heat, flames and sparks. Exposure to air. Exposure to moisture. Extremes of temperature and direct sunlight.
Incompatible materials	: Acids alkalis Amines Ammonia halogens Peroxides Reducing agents Strong oxidizing agents Oxygen aluminum

nitrates organic absorbents such as sawdust, peat moss, ground corn cobs, etc. Bases metal salts

SECTION 11. TOXICOLOGICAL INFORMATION

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Product:

Acute oral toxicity	: Acute toxicity estimate : > 5,000 mg/kg Method: Calculation method
Acute inhalation toxicity	 Acute toxicity estimate : > 30000 ppm Exposure time: 4 h Test atmosphere: gas Method: Calculation method
Acute dermal toxicity	: Acute toxicity estimate : > 5,000 mg/kg Method: Calculation method
Components:	
67-64-1: Acute oral toxicity	: LD50 (rat): 5,800 mg/kg
Acute inhalation toxicity	: LC50 (rat): 76.0 mg/l Exposure time: 4 h
Acute dermal toxicity	: LD50 : > 7,426 mg/kg
64742-49-0:	
Acute oral toxicity	: LD50 (rat, male and female): > 5,000 mg/kg Method: OECD Test Guideline 401 GLP: yes
Acute inhalation toxicity	: Remarks: No data available
Acute dermal toxicity	: LD50 (rabbit, male and female): > 2,000 mg/kg Method: OECD Test Guideline 402 GLP: yes
64742-89-8: Acute oral toxicity	: LD50 (rat, male and female): > 5,000 mg/kg Method: OECD Test Guideline 401 GLP: yes

Acute inhalation toxicity	: Remarks: No data available
Acute dermal toxicity	: LD50 (rabbit, male and female): > 2,000 mg/kg Method: OECD Test Guideline 402 GLP: yes
68410-97-9: Acute oral toxicity	: LD50 (rat): > 5,000 mg/kg
Acute inhalation toxicity	: Remarks: No data available
Acute dermal toxicity	: LD50 (rabbit): > 2,000 mg/kg
108-65-6: Acute oral toxicity	: LD50 (rat): 8,532 mg/kg
Acute inhalation toxicity	: Remarks: No data available
Acute dermal toxicity	: LD50 (rabbit): > 5,000 mg/kg Method: OECD Test Guideline 402
108-88-3: Acute oral toxicity	: LD50 (rat, male): > 5,580 mg/kg
Acute inhalation toxicity	: LC50 (rat, male and female): 28.1 mg/l Exposure time: 4 h Test atmosphere: vapour Method: OECD Test Guideline 403
Acute dermal toxicity	: LD50 (rabbit): > 5,000 mg/kg
110-19-0: Acute oral toxicity	: LD50 (rat): 13,413 mg/kg Assessment: The substance or mixture has no acute oral toxicity
Acute inhalation toxicity	: LC50 (rat): 23.4 mg/l Exposure time: 4 h Assessment: The substance or mixture has no acute inhalation toxicity Remarks: Information given is based on data obtained from similar substances.
Acute dermal toxicity	: LD50 (rabbit): > 17,400 mg/kg Assessment: The substance or mixture has no acute dermal toxicity
123-86-4: Acute oral toxicity	: LD50 (rat): > 5,000 mg/kg

	Method: OECD Test Guideline 423 GLP: no
Acute inhalation toxicity	: LC50 (rat, male and female): > 21 mg/l Exposure time: 4 h Test atmosphere: vapour Method: OECD Test Guideline 403 GLP: yes
Acute dermal toxicity	: LD50 (rabbit, male and female): > 5,000 mg/kg Method: OECD Test Guideline 402 GLP: yes
1330-20-7: Acute oral toxicity	: LD50 (rat, male): 3,523 mg/kg Method: EU Method B.1 (Acute Toxicity, Oral) GLP: no
Acute inhalation toxicity	: LC50 (rat, male): 6700 ppm Exposure time: 4 h Method: Directive 67/548/EEC, Annex V, B.2. Assessment: The component/mixture is moderately toxic after short term inhalation.
Acute dermal toxicity	: LD50 (rabbit): 1,100 mg/kg Assessment: The component/mixture is moderately toxic after single contact with skin.
100-41-4: Acute inhalation toxicity	: LC50 (Mouse, Male): 10 mg/l Exposure time: 4 h Assessment: The component/mixture is moderately toxic after short term inhalation.
Acute dermal toxicity	: LD50 (rabbit): 15,433 mg/kg
142-82-5: Acute oral toxicity	: LD50 (rat, male and female): 5,000 mg/kg Method: OECD Test Guideline 401 Symptoms: Salivation GLP: yes Remarks: Information given is based on data obtained from similar substances.
Acute inhalation toxicity	: LC50 (rat, male and female): 73.5 mg/l Exposure time: 4 h Test atmosphere: vapour Method: OECD Test Guideline 403
Acute dermal toxicity	: LD50 (rabbit, male and female): > 2,000 mg/kg

Method: OECD Test Guideline 402 GLP: yes Remarks: Information given is based on data obtained from similar substances.

Skin corrosion/irritation

Product:

Remarks: Irritating to skin.

Components:

67-64-1:

Species: rabbit Exposure time: 24 h Method: In vivo Result: Mild skin irritation

64742-49-0:

Species: rabbit Result: Irritating to skin.

64742-89-8:

Species: rabbit Exposure time: 4 h Result: Irritating to skin.

68410-97-9:

Species: rabbit Result: Irritating to skin.

108-65-6:

Species: rabbit Method: OECD Test Guideline 404 Result: No skin irritation

108-88-3:

Species: rabbit Exposure time: 4 h Result: Irritating to skin.

110-19-0:

Species: rabbit Result: No skin irritation

123-86-4:

Species: rabbit Method: OECD Test Guideline 404 Result: No skin irritation GLP: no

1330-20-7:

Species: rabbit Exposure time: 24 h Result: Irritating to skin.

100-41-4: Species: rabbit Result: Mild skin irritation

142-82-5:

Species: rabbit Exposure time: 24 h Method: OECD Test Guideline 404 Result: Irritating to skin. GLP: yes Remarks: Based on a similar product formulation.

Serious eye damage/eye irritation

Product: Remarks: Irritating to eyes.

Components:

67-64-1: Species: rabbit Result: Irritating to eyes. Exposure time: 24 h

64742-49-0:

Species: rabbit Result: Irritating to eyes.

64742-89-8:

Species: rabbit Result: Irritating to eyes.

68410-97-9:

Species: rabbit Result: Irritating to eyes.

108-65-6:

Species: rabbit Result: No eye irritation Method: OECD Test Guideline 405

108-88-3:

Species: rabbit Result: Irritating to eyes. Method: OECD Test Guideline 405

110-19-0:

Species: rabbit Result: No eye irritation

123-86-4:

Species: rabbit Result: No eye irritation GLP: yes

1330-20-7:

Species: rabbit Result: Irritating to eyes.

100-41-4:

Species: rabbit Result: Mild eye irritation

142-82-5:

Species: rabbit Result: Irritating to eyes. Method: OECD Test Guideline 405 GLP: yes Remarks: Information given is based on data obtained from similar substances.

Respiratory or skin sensitisation

Components:

67-64-1: Test Type: Maximization test Species: guinea pig Result: Did not cause sensitisation on laboratory animals.

64742-49-0:

Test Type: Buehler Test Species: guinea pig Result: Did not cause sensitisation on laboratory animals.

64742-89-8:

Test Type: Buehler Test Species: guinea pig Result: Did not cause sensitisation on laboratory animals.

108-65-6:

Test Type: Maximization test Species: guinea pig Method: OECD Test Guideline 406 Result: Did not cause sensitisation on laboratory animals. GLP: no

108-88-3:

Test Type: Maximisation Test (GPMT) Species: guinea pig Result: Did not cause sensitisation on laboratory animals. GLP: yes

110-19-0:

Test Type: Maximization test Species: guinea pig Result: Did not cause sensitisation on laboratory animals.

123-86-4:

Species: guinea pig Result: Did not cause sensitisation on laboratory animals.

1330-20-7:

Remarks: No data available

100-41-4: Remarks: No data available

142-82-5:

Test Type: Maximization test Species: guinea pig Method: OECD Test Guideline 406 Result: Does not cause skin sensitisation. Remarks: Based on a similar product formulation.

Germ cell mutagenicity

Components:

67-64-1: Genotoxicity in vitro

: Test Type: Mammalian cell gene mutation assay Test species: Mouse lymphoma cells Metabolic activation: Without metabolic activation Method: OECD Test Guideline 476 Result: negative

: Test Type: Ames test Metabolic activation: with and without metabolic activation Method: OECD Test Guideline 471 Result: negative

: Test Type: Chromosome aberration test in vitro Test species: Chinese hamster ovary (CHO) Metabolic activation: with and without metabolic activation Method: OECD Test Guideline 473 Result: negative

Genotoxicity in vivo	: Test Type: In vivo micronucleus test Test species: mouse Application Route: Oral Exposure time: 13 wk Dose: 5,000, 10,000, 20,000 ppm Result: negative
Germ cell mutagenicity- Assessment	: Tests on bacterial or mammalian cell cultures did not show mutagenic effects.
64742-49-0: Germ cell mutagenicity- Assessment	: Mutagenicity classification not possible from current data
64742-89-8: Germ cell mutagenicity- Assessment	: Mutagenicity classification not possible from current data
68410-97-9: Genotoxicity in vitro	: Test Type: Mammalian cell gene mutation assay Test species: mouse lymphoma cells Result: positive
Genotoxicity in vivo	: Test Type: In vivo micronucleus test Test species: mouse Method: OECD Test Guideline 474 Result: positive
Germ cell mutagenicity- Assessment	: Positive result(s) from in vivo heritable germ cell mu- tagenicity tests in mammals
108-65-6: Genotoxicity in vitro	: Test Type: DNA damage and/or repair Test species: rat hepatocytes Metabolic activation: Without metabolic activation Method: OECD Test Guideline 482 Result: negative GLP: yes
Germ cell mutagenicity- Assessment	: Tests on bacterial or mammalian cell cultures did not show mutagenic effects.
108-88-3: Genotoxicity in vitro	: Test Type: Mammalian cell gene mutation assay Test species: Mouse lymphoma cells Metabolic activation: with and without metabolic acti- vation Method: OECD Test Guideline 476 Result: negative

Genotoxicity in vivo	 Test Type: Dominant lethal assay Test species: mouse (male) Application Route: inhalation (vapour) Exposure time: 6 h/d, 5 d/wk for 8 wks Dose: 0, 100, 400 ppm Method: OECD Test Guideline 478 Result: negative
Germ cell mutagenicity- Assessment	: Tests on bacterial or mammalian cell cultures did not show mutagenic effects.
110-19-0: Genotoxicity in vitro	: Test Type: Chromosome aberration test in vitro Test species: Chinese hamster lung fibroblasts Metabolic activation: with and without metabolic acti- vation Result: negative
Genotoxicity in vivo	: Test Type: In vivo micronucleus test Test species: mouse Application Route: Oral Result: negative
Germ cell mutagenicity- Assessment	: Tests on bacterial or mammalian cell cultures did not show mutagenic effects.
123-86-4: Genotoxicity in vitro	: Test Type: Chromosome aberration test in vitro Test species: Chinese hamster lung fibroblasts Metabolic activation: Without metabolic activation Method: OECD Test Guideline 473 Result: negative GLP: No data available
Genotoxicity in vivo	 Test Type: In vivo micronucleus test Test species: mouse (male and female) Application Route: Oral Dose: 500, 1000, 2000 mg/kg bw Method: OECD Test Guideline 474 Result: negative GLP: yes Test substance: Information given is based on data obtained from similar substances.
Germ cell mutagenicity- Assessment	: Tests on bacterial or mammalian cell cultures did not show mutagenic effects.
1330-20-7: Genotoxicity in vitro	: Test Type: Chromosome aberration test in vitro Test species: Chinese hamster ovary (CHO)

	Metabolic activation: with and without metabolic acti- vation Method: Mutagenicity (in vitro mammalian cytogenetic test) Result: negative
	 Test Type: Sister chromatid exchange assay in mam- malian cells Test species: Chinese hamster ovary (CHO) Metabolic activation: with and without metabolic acti- vation Result: negative
Genotoxicity in vivo	: Test Type: Dominant lethal assay Test species: mouse Application Route: Subcutaneous Exposure time: 8 wk Dose: 1.0 mL/kg Method: OECD Test Guideline 478 Result: negative GLP: no
Germ cell mutagenicity- Assessment	: Animal testing did not show any mutagenic effects.
100-41-4: Genotoxicity in vitro	: Test Type: Chromosome aberration test in vitro Test species: Chinese hamster ovary (CHO) Metabolic activation: with and without metabolic acti- vation Method: OECD Test Guideline 473 Result: negative GLP: no
	: Test Type: Mammalian cell gene mutation assay Test species: mouse lymphoma cells Metabolic activation: with and without metabolic acti- vation Method: OECD Test Guideline 476 Result: negative GLP: yes
Genotoxicity in vivo	: Test Type: In vivo micronucleus test Test species: mouse (male) Application Route: Oral Method: OECD Test Guideline 474 Result: negative GLP: yes
	Test Type: DNA damage and/or repair

	Test species: mouse (male and female) Application Route: Inhalation Method: OECD Test Guideline 486 Result: negative GLP: yes
Germ cell mutagenicity- Assessment	: In vivo tests did not show mutagenic effects
142-82-5: Genotoxicity in vitro	: Test Type: Chromosome aberration test in vitro Test species: Rat liver Metabolic activation: Without metabolic activation Method: OECD Test Guideline 473 Result: negative
	: Test Type: Ames test Metabolic activation: with and without metabolic acti- vation Method: OECD Test Guideline 471 Result: negative
Germ cell mutagenicity- Assessment	: Did not show mutagenic effects in animal experi- ments.

Carcinogenicity

Components:

67-64-1: Species: mouse, (female) Application Route: Dermal Exposure time: 365 d (90%) or 424 d (100%) Dose: 0.1ml 90(71mg) or 100% (79mg) Frequency of Treatment: 3 times per wk NOAEL: 79

Result: did not display carcinogenic properties

Carcinogenicity - As- sessment	:	Carcinogenicity classification not possible from current data.
64742-49-0: Carcinogenicity - As- sessment	:	Not classifiable as a human carcinogen.
64742-89-8: Carcinogenicity - As- sessment	:	Not classifiable as a human carcinogen.

68410-97-9:

Species: mouse NOAEL: 50 mg/kg bw/day

Method: OECD Test Guideline 451 Result: evidence of carcinogenic activity

Carcinogenicity - As- : Possible human carcinogen sessment

108-65-6:

Species: rat, (male and female) Application Route: inhalation (vapour) Exposure time: 2 yr Dose: 0, 300, 1000, 3000 ppm Frequency of Treatment: 6 hr/d, 5 d/wk NOAEL: No observed adverse effect level: 3,000 ppm

Method: OECD Test Guideline 453 Result: did not display carcinogenic properties GLP: yes

Carcinogenicity - As- : No evidence of carcinogenicity in animal studies. sessment

108-88-3:

Species: rat, (male and female) Application Route: inhalation (vapour) Exposure time: 103 wks Dose: 0, 600, 1200 ppm Frequency of Treatment: 6.5 h/d, 5 d/wk NOAEL: No observed adverse effect level: 1,200 ppm

Method: OECD Test Guideline 453 Result: did not display carcinogenic properties Symptoms: Erosion of nasal epithelium GLP: yes

Carcinogenicity - As- : Not classifiable as a human carcinogen. sessment

110-19-0:

Remarks: This information is not available.

Carcinogenicity - As- : No evidence of carcinogenicity in animal studies. sessment

123-86-4:

Remarks: This information is not available.

Carcinogenicity - As- : No evidence of carcinogenicity in animal studies.

sessment

1330-20-7:

Species: mouse, (male and female) Application Route: Oral Exposure time: 103 wk Dose: 0, 500 or 1000 mg/kg Frequency of Treatment: 5 days/week Method: Directive 67/548/EEC, Annex V, B.32. Result: did not display carcinogenic properties GLP: No data available

Carcinogenicity - As- : Animal testing did not show any carcinogenic effects. sessment

100-41-4:

Species: mouse, (male and female) Application Route: Inhalation Exposure time: 103 wk Activity duration: 6 h Dose: 0, 75, 250, 750 ppm Frequency of Treatment: 5 days/week NOAEL: 250 ppm

Method: OECD Test Guideline 453 Result: evidence of carcinogenic activity Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes

Carcinogenicity - As- : Suspected human carcinogens sessment

142-82-5:

Remarks: This information is not available.

Carcinogenicity - As-	:	Carcinogenicity classification not possible from current
sessment		data.

Reproductive toxicity

Components: 67-64-1

: Species: rat, male
Application Route: oral
Dose: 0, 5000, 10000 mg/L
Frequency of Treatment: 7 days/week
General Toxicity - Parent: LOAEL: 10,000
Fertility: 10,000

Effects on foetal devel- opment	: Species: rat Application Route: Inhalation Dose: 0, 440, 2200, 11000 ppm Frequency of Treatment: 7 days/week General Toxicity Maternal: NOAEC: 2,200 ppm Teratogenicity: NOAEC: 11,000 ppm Embryo-foetal toxicity.: NOAEC: 2,200 ppm Method: OECD Test Guideline 414 Result: No teratogenic potential. GLP: No data available
Reproductive toxicity - Assessment	 No evidence of adverse effects on sexual function and fertility, and on development, based on animal exper- iments.
64742-49-0: Reproductive toxicity - Assessment	: Fertility classification not possible from current data. Embryotoxicity classification not possible from current data.
64742-89-8: Reproductive toxicity - Assessment	: Fertility classification not possible from current data. Embryotoxicity classification not possible from current data.
68410-97-9: Reproductive toxicity - Assessment	: Fertility classification not possible from current data. Embryotoxicity classification not possible from current data.
108-65-6: Effects on fertility	 Species: rat Application Route: Oral Dose: 0, 100, 300, 1000 mg/kg General Toxicity - Parent: NOAEL: 1,000 mg/kg bw General Toxicity F1: NOAEL: 1,000 mg/kg bw Method: OECD Test Guideline 422 Result: Animal testing did not show any effects on fertility. GLP: yes Remarks: Information given is based on data obtained from similar substances.
Effects on foetal devel- opment	: Species: rat Application Route: Inhalation Dose: 0, 500, 2000, 4000 ppm Duration of Single Treatment: 9 d Frequency of Treatment: 6 hr/day General Toxicity Maternal: NOAEL: 500 ppm Teratogenicity: NOAEL: > 4,000 ppm

	GLP: yes
Reproductive toxicity - Assessment	 No evidence of adverse effects on sexual function and fertility, and on development, based on animal exper- iments.
108-88-3: Effects on fertility	 Test Type: Two-generation study Species: rat, male and female Application Route: Inhalation Dose: 0, 100, 500, 2000 ppm Frequency of Treatment: 7 days/week General Toxicity - Parent: NOAEC: 500 ppm General Toxicity F1: NOAEC: 500 ppm Fertility: NOAEC: 2,000 ppm Symptoms: Reduced maternal body weight gain. Re- duced offspring weight gain. Method: OECD Test Guideline 416 Result: Animal testing did not show any effects on fertility. GLP: yes
	Test Type: Fertility Species: rat, male and female Application Route: inhalation (vapour) Dose: 0, 600, 1200 ppm Frequency of Treatment: 7 days/week General Toxicity - Parent: NOAEC: 600 ppm Symptoms: Decreased sperm count Result: Animal testing did not show any effects on fertility.
Effects on foetal devel- opment	: Species: rat Application Route: inhalation (vapour) Dose: 0, 250, 750, 1500, 3000 ppm Duration of Single Treatment: 10 d Frequency of Treatment: 6 hr/day General Toxicity Maternal: NOAEC: 750 ppm Developmental Toxicity: NOAEC: 750 ppm Symptoms: Maternal toxicity, Reduced body weight, Skeletal malformations. GLP: yes
Reproductive toxicity - Assessment	: Some evidence of adverse effects on sexual function and fertility, and/or on development, based on animal experiments.
110-19-0: Effects on fertility	: Test Type: Two-generation study Species: rat Application Route: Inhalation

	Duration of Single Treatment: 6 h Frequency of Treatment: 7 days/week General Toxicity - Parent: NOAEL: 2,500 ppm Method: OECD Test Guideline 416	
Reproductive toxicity - Assessment	No evidence of adverse effects on sexual function ar fertility, and on development, based on animal experiments.	
123-86-4: Effects on fertility	Species: rat, male and female Application Route: Inhalation Dose: 0, 750, 1500, 2000 ppm Duration of Single Treatment: 6 h Frequency of Treatment: 7 days/week General Toxicity - Parent: NOAEC: 750 ppm General Toxicity F1: NOAEC: 750 ppm Fertility: NOAEC: 2,000 ppm Early Embryonic Development: NOAEC: 750 ppm Symptoms: Effect on reproduction capacity. Method: OECD Test Guideline 416 GLP: yes	
Effects on foetal devel- opment	Species: rat, male and female Application Route: vapour Dose: 500, 1500, 3000 ppm Duration of Single Treatment: 6 h Frequency of Treatment: 5 days/week GLP: yes	
Reproductive toxicity - Assessment	Fertility classification not possible from current data. Embryotoxicity classification not possible from curredata.	
1330-20-7: Effects on fertility	Test Type: Two-generation study Species: rat, male and female Application Route: Inhalation Dose: 0, 25, 100 and 500 ppm Duration of Single Treatment: 6 h Frequency of Treatment: 7 days/week General Toxicity - Parent: NOAEC: > 500 ppm General Toxicity F1: NOAEC: > 500 ppm Early Embryonic Development: NOAEC: > 500 ppm Result: No reproductive effects.	
Effects on foetal devel- opment	Species: rat Application Route: Inhalation Dose: 0, 100, 500, 1000 or 2000 ppm Duration of Single Treatment: 14 d	

	Frequency of Treatment: 6 hr/day General Toxicity Maternal: NOAEC: 500 ppm Teratogenicity: NOAEC: > 2,000 Developmental Toxicity: NOAEC: 100 ppm Result: No teratogenic effects., Developmental toxicity occurred at maternal toxicity dose levels
Reproductive toxicity - Assessment	: Animal testing did not show any effects on fertility. Damage to fetus not classifiable
100-41-4: Effects on fertility	 Test Type: One generation study Species: rat, male and female Application Route: Inhalation Dose: 0, 100, 500 and 1000 ppm Duration of Single Treatment: 6 h General Toxicity - Parent: NOAEC: 1,000 ppm General Toxicity F1: NOAEC: 100 ppm Symptoms: Reduced foetal weight. Reduced offspring weight gain. Method: OECD Test Guideline 415 Result: No reproductive effects. GLP: yes
Effects on foetal devel- opment	: Species: rat Application Route: Inhalation Dose: 0, 100, 500, 1000, 2000 ppm Duration of Single Treatment: 15 d General Toxicity Maternal: NOAEC: 500 ppm Teratogenicity: NOAEC: 2,000 ppm Developmental Toxicity: NOAEC: 500 ppm Symptoms: Reduced body weight Method: OECD Test Guideline 414 Result: Developmental toxicity occurred at maternal toxicity dose levels GLP: No data available
Reproductive toxicity - Assessment	: Fertility classification not possible from current data. Embryotoxicity classification not possible from current data.
142-82-5: Effects on fertility	: Test Type: Two-generation study Species: rat, male and female Application Route: vapour Dose: 0, 900, 3000, 9000 ppm Frequency of Treatment: 5 days/week General Toxicity - Parent: NOAEC: 3,000 ppm General Toxicity F1: NOAEC: 3,000 ppm Fertility: NOAEC: 9,000 ppm Symptoms: Reduced maternal body weight gain. Re-

	duced offspring weight gain. Method: OECD Test Guideline 416 Result: No reproductive effects. GLP: yes Remarks: Information given is based on data obtained from similar substances.
Effects on foetal devel- opment	 Species: mouse Application Route: inhalation (vapour) Dose: 0, 900, 3000, 9000 ppm Duration of Single Treatment: 10 d Frequency of Treatment: 6 hr/day General Toxicity Maternal: NOAEC: 900 ppm Developmental Toxicity: NOAEC: 3,000 ppm Symptoms: Skeletal malformations. Method: OECD Test Guideline 414 GLP: yes Remarks: Information given is based on data obtained from similar substances.
Reproductive toxicity - Assessment	Animal testing did not show any effects on fertility. Embryotoxicity classification not possible from current data.

STOT - single exposure

Product:No data available

Components: 67-64-1:

Exposure routes:	Target Organs:	Assessment:	Remarks:
Inhalation	Central nervous system	May cause drowsi- ness or dizziness., The substance or mixture is classified as specific target organ toxicant, sin- gle exposure, cate- gory 3 with narcotic effects.	

64742-49-0:

Exposure routes:	Target Organs:	Assessment:	Remarks:
Inhalation	Central nervous	May cause drowsi-	
	system	ness or dizziness.,	
		The substance or	
		mixture is classified	
		as specific target	
		organ toxicant, sin-	
		gle exposure, cate-	

	gory 3 with narcotic	
	effects.	

64742-89-8:No data available

68410-97-9:

Exposure routes:	Target Organs:	Assessment:	Remarks:
Inhalation	Central nervous system	May cause drowsi- ness or dizziness., The substance or mixture is classified as specific target organ toxicant, sin- gle exposure, cate- gory 3 with narcotic effects.	

108-65-6:No data available

108-88-3:

Exposure routes:	Target Organs:	Assessment:	Remarks:
Inhalation	Central nervous system	May cause drowsi- ness or dizziness., The substance or mixture is classified as specific target organ toxicant, sin- gle exposure, cate- gory 3 with narcotic effects.	

110-19-0:

Exposure routes:	Target Organs:	Assessment:	Remarks:
Inhalation	Central nervous system	The substance or mixture is classified as specific target organ toxicant, sin- gle exposure, cate- gory 3 with narcotic effects., May cause drowsiness or diz- ziness.	

123-86-4:	

120 00 11				
Exposure routes:	Target Organs:	Assessment:	Remarks:	

Inhalation	Central nervous	May cause drowsi-	
	system	ness or dizziness.,	
	-	The substance or	
		mixture is classified	
		as specific target	
		organ toxicant, sin-	
		gle exposure, cate-	
		gory 3 with narcotic	
		effects.	

1330-20-7:

Exposure routes:	Target Organs:	Assessment:	Remarks:
Inhalation	Respiratory system	May cause respira- tory irritation., The substance or mix- ture is classified as specific target or- gan toxicant, single exposure, category 3 with respiratory tract irritation.	

100-41-4:No data available

142-82-5:

Exposure routes:	Target Organs:	Assessment:	Remarks:
Inhalation	Central nervous system	May cause drowsi- ness or dizziness., The substance or mixture is classified as specific target organ toxicant, sin- gle exposure, cate- gory 3 with narcotic effects.	

STOT - repeated exposure

Product:No data available

Components:

67-64-1:No data available

64742-49-0:No data available

64742-89-8:No data available

68410-97-9:No data available

108-65-6:No data available

108-88-3:

Exposure routes:	Target Organs:	Assessment:	Remarks:
Inhalation	Auditory system, Eyes	May cause damage to organs through prolonged or re- peated exposure., The substance or mixture is classified as specific target organ toxicant, re- peated exposure, category 2.	

110-19-0:No data available

123-86-4:No data available

1330-20-7:

Exposure routes:	Target Organs:	Assessment:	Remarks:
	Liver, Kidney, Cen- tral nervous system	May cause damage to organs through prolonged or re- peated exposure., The substance or mixture is classified as specific target organ toxicant, re- peated exposure, category 2.	

100-41-4:

Exposure routes: Target Organs:	Assessment:	Remarks:	
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Auditory system	May cause damage to organs through prolonged or re- peated exposure., The substance or mixture is cleasified	
	mixture is classified as specific target organ toxicant, re- peated exposure, category 2.	

142-82-5:No data available

Repeated dose toxicity

Components:

67-64-1:

Species: mouse, male NOAEL: 20000 Application Route: Oral Exposure time: 13 wk Number of exposures: daily Dose: 1250, 2500, 5000, 10000, 20000 Method: OECD Test Guideline 408 GLP: No data available

Species: mouse, female NOAEL: 20000 LOAEL: 50000 Application Route: Oral Exposure time: 13 wk Number of exposures: daily Dose: 2500, 5000, 10000, 20000, 5000 Method: OECD Test Guideline 408 GLP: No data available

Repeated dose toxicity - : Causes mild skin irritation., Causes serious eye irrita-Assessment tion.

64742-89-8:

Species: rat, male and female NOAEL: 1402 Application Route: inhalation (vapour) Test atmosphere: vapour Exposure time: 13 weeks Number of exposures: 6 hours/day, 5 days/week Dose: 322, 1402, 9869 mg/m3 GLP: yes Target Organs: Kidney Symptoms: Nasal and ocular discharge

108-65-6:

Species: rat, male and female NOAEL: > 1,000 mg/kg Application Route: Oral Dose: 0, 100, 300, 1000 mg/kg Method: OECD Test Guideline 422

108-88-3:

Species: rat, male and female NOAEL: 300 Application Route: inhalation (vapour) Exposure time: 6, 12, or 18 mths Number of exposures: 6 h/d, 5 d/wk Dose: 0, 30, 100, 300 ppm Method: OECD Test Guideline 453

Repeated dose toxicity - : Causes skin irritation. Assessment

110-19-0:

Species: rat NOAEL: 316 mg/kg Application Route: Oral Exposure time: 92 d

123-86-4:

Species: rat, male and female NOAEL: 500 Application Route: inhalation (vapour) Exposure time: 13 wk Number of exposures: 6 h/d, 5d/wk Dose: 500, 1500, 3000 ppm GLP: yes Symptoms: oral or nasal discharge

1330-20-7:

Species: rat, male and female NOAEL: 250 mg/kg Application Route: Oral Exposure time: 103 wk Number of exposures: 5 d/wk Dose: 0, 250 or 500 mg/kg Assessment: The substance or mixture is classified as specific target organ toxicant, repeated exposure, category 2.

100-41-4:

Species: rat, male and female

NOAEL: 75 mg/kg Application Route: Oral Exposure time: 28 d Dose: 75, 250 and 750 mg/kg bw/day Method: OECD Test Guideline 407 GLP: yes Symptoms: Increased kidney and liver weights

142-82-5:

Species: rat, male NOAEL: 12470 mg/m3 Application Route: inhalation (vapour) Exposure time: 16 wks Number of exposures: 12 h/d, 7 d/wk Dose: 0, 12470 mg/3

Repeated dose toxicity - : Causes skin irritation. Assessment

Aspiration toxicity

Components:

64742-49-0:

May be fatal if swallowed and enters airways.

64742-89-8:

May be fatal if swallowed and enters airways.

68410-97-9:

May be fatal if swallowed and enters airways.

108-88-3:

Aspiration Toxicity - Category 1

1330-20-7:

May be fatal if swallowed and enters airways.

100-41-4:

May be fatal if swallowed and enters airways.

142-82-5: Aspiration Toxicity - Category 1

Further information

Product:

Remarks: Symptoms of overexposure may be headache, dizziness, tiredness, nausea and vomiting., Concentrations substantially above the TLV value may cause narcotic effects., Solvents may degrease the skin.

SECTION 12. ECOLOGICAL INFORMATION

Ecotoxicity		
<u>Components:</u> 67-64-1:		
Toxicity to fish	:	LC50 (Oncorhynchus mykiss (rainbow trout)): 6,100 mg/l Exposure time: 48 h
Toxicity to daphnia and other aquatic inverte- brates	:	EC50 (Daphnia magna (Water flea)): 7,630 mg/l Exposure time: 48 h Test substance: Acetone
Toxicity to algae	:	Remarks: No data available
64742-49-0:		
Toxicity to fish	:	LC50 (Oncorhynchus mykiss (rainbow trout)): 10 mg/l Exposure time: 96 h
Toxicity to daphnia and other aquatic inverte- brates	:	EC50 (Daphnia magna (Water flea)): 4.5 mg/l Exposure time: 48 h
Toxicity to algae	:	EC50 (Pseudokirchneriella subcapitata (green algae)): 3.71 mg/l Exposure time: 96 h
Ecotoxicology Assessment Acute aquatic toxicity	:	Toxic to aquatic life.
Chronic aquatic toxicity	:	Toxic to aquatic life with long lasting effects.
64742-89-8:		
Toxicity to fish	:	LC50 (Oncorhynchus mykiss (rainbow trout)): 8.2 mg/l Exposure time: 96 h Test Type: semi-static test
Toxicity to daphnia and other aquatic inverte- brates	:	EC50 (Daphnia magna (Water flea)): 4.5 mg/l Exposure time: 48 h Test Type: Immobilization Analytical monitoring: yes
Toxicity to algae	:	EC50 (Pseudokirchneriella subcapitata (green algae)):

	3.7 mg/l Exposure time: 96 h Test Type: static test
Ecotoxicology Assessment Acute aquatic toxicity	: Toxic to aquatic life.
Chronic aquatic toxicity	: Toxic to aquatic life with long lasting effects.
68410-97-9:	
Toxicity to fish	: LC50 (Pimephales promelas (fathead minnow)): 8.2 mg/l Exposure time: 96 h
Toxicity to daphnia and other aquatic inverte- brates	: EC50 (Daphnia magna (Water flea)): 4.5 mg/l Exposure time: 48 h
Toxicity to algae	 EC50 (Pseudokirchneriella subcapitata (green algae)): 3.1 mg/l Exposure time: 72 h Method: OECD Test Guideline 201
Ecotoxicology Assessment Acute aquatic toxicity	: Toxic to aquatic life.
Chronic aquatic toxicity	: Toxic to aquatic life with long lasting effects.
108-65-6:	
Toxicity to fish	 LC50 (Oncorhynchus mykiss (rainbow trout)): > 100 mg/l Exposure time: 96 h Test Type: static test Method: OECD Test Guideline 203
Toxicity to daphnia and other aquatic inverte- brates	: EC50 (Daphnia magna (Water flea)): 500 mg/l Exposure time: 48 h Test Type: Immobilization
Toxicity to algae	: EC50 (Selenastrum capricornutum (green algae)): > 1,000 mg/l End point: Growth rate Exposure time: 96 h Test Type: static test
108-88-3:	
Toxicity to fish	: LC50 (Oncorhynchus mykiss (rainbow trout)): 5.5 mg/l Exposure time: 96 h Test Type: flow-through test

Toxicity to daphnia and other aquatic inverte- brates	: EC50 (Ceriodaphnia dubia): 3.78 mg/l Exposure time: 48 h Test Type: Renewal
Toxicity to algae	: EC50 (Chlorella vulgaris (Fresh water algae)): 134 mg/l Exposure time: 3 h Test Type: static test
Toxicity to bacteria	: IC50 (Bacteria): 84 mg/l Exposure time: 24 h Test Type: Static
Ecotoxicology Assessment Acute aquatic toxicity	: Toxic to aquatic life.
Chronic aquatic toxicity	: Toxic to aquatic life with long lasting effects.
110-19-0:	
Toxicity to fish	: LC50 (Oryzias latipes (Japanese medaka)): 17 mg/l Exposure time: 96 h
Toxicity to daphnia and other aquatic inverte- brates	: (Daphnia magna (Water flea)): 25 mg/l Exposure time: 48 h Test Type: static test
Toxicity to algae	: EC50 (Pseudokirchneriella subcapitata): 370 mg/l Exposure time: 72 h Test Type: static test
Ecotoxicology Assessment Acute aquatic toxicity	: This product has no known ecotoxicological effects.
Chronic aquatic toxicity	: This product has no known ecotoxicological effects.
123-86-4:	
Toxicity to fish	: LC50 (Pimephales promelas (fathead minnow)): 18 mg/l Exposure time: 96 h Test Type: flow-through test Method: OECD Test Guideline 203 GLP: no
Toxicity to daphnia and other aquatic inverte- brates	: EC50 (Daphnia magna (Water flea)): 44 mg/l Exposure time: 48 h Test Type: static test
Toxicity to algae	: EC50 (Desmodesmus subspicatus (green algae)): 674.7 mg/l

	End point: Growth rate Exposure time: 72 h
Toxicity to daphnia and other aquatic inverte-brates (Chronic toxicity)	: NOEC (Daphnia magna (Water flea)): 23 mg/l Exposure time: 21 d
Toxicity to bacteria	: EC 50 (Tetrahymena pyriformis (Ciliate)): 356 mg/l Exposure time: 40 h Test Type: Static
Ecotoxicology Assessment Acute aquatic toxicity	: Harmful to aquatic life.
Chronic aquatic toxicity	: Harmful to aquatic life with long lasting effects.
1330-20-7:	
Toxicity to fish	: LC50 (Oncorhynchus mykiss (rainbow trout)): 2.6 mg/l Exposure time: 96 h Method: OECD Test Guideline 203
Toxicity to daphnia and other aquatic inverte- brates	: EC50 (Daphnia magna (Water flea)): 1 mg/l Exposure time: 24 h Test Type: static test Method: OECD Test Guideline 202
Toxicity to algae	: EC50 (Pseudokirchneriella subcapitata): 4.36 mg/l End point: Growth rate Exposure time: 73 h Test Type: static test Analytical monitoring: yes Method: OECD Test Guideline 201 GLP: yes
Ecotoxicology Assessment Acute aquatic toxicity	: Toxic to aquatic life.
Chronic aquatic toxicity	: Toxic to aquatic life with long lasting effects.
100-41-4:	
Toxicity to fish	: LC50 (Oncorhynchus mykiss (rainbow trout)): 4.2 mg/l Exposure time: 96 h Test Type: semi-static test
Toxicity to daphnia and other aquatic inverte- brates	: EC50 (Daphnia magna (Water flea)): 1.8 mg/l Exposure time: 48 h Test Type: static test

Toxicity to algae	: EC50 (Pseudokirchneriella subcapitata): 5.4 mg/l Exposure time: 72 h Test Type: static test
Toxicity to bacteria	: Remarks: No data available
Ecotoxicology Assessment Acute aquatic toxicity	: Toxic to aquatic life.
Chronic aquatic toxicity	: Toxic to aquatic life with long lasting effects.
142-82-5: Toxicity to fish	: LC50 (Carassius auratus (goldfish)): 4 mg/l Exposure time: 24 h Remarks: Very toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment.
Toxicity to daphnia and other aquatic inverte- brates	 EC50 (Daphnia magna (Water flea)): 1.5 mg/l Exposure time: 48 h Test Type: static test Remarks: Very toxic to aquatic organisms.
Toxicity to algae	: Remarks: No data available
Ecotoxicology Assessment Acute aquatic toxicity	: Very toxic to aquatic life.
Chronic aquatic toxicity	: Very toxic to aquatic life with long lasting effects.

Persistence and degradability

Components:67-64-1:Biodegradability:	Remarks: Readily biodegradable
64742-49-0:	
Biodegradability :	aerobic Inoculum: activated sludge Concentration: 20 mg/l Biodegradation: 74.30 % Exposure time: 56 d GLP: yes Remarks: Inherently biodegradable.
64742-89-8:	
Biodegradability :	Concentration: 49.2 mg/l Result: Readily biodegradable. Biodegradation: 77 % Testing period: 2 d

	Exposure time: 28 d GLP: yes
108-65-6:	
Biodegradability	: aerobic Inoculum: activated sludge Concentration: 76.4 mg/l Result: Readily biodegradable. Biodegradation: 90 % Exposure time: 28 d GLP: yes
Biochemical Oxygen De- mand (BOD)	: 0.36 mg/l
Chemical Oxygen De- mand (COD)	: 1.74 mg/l
108-88-3:	
Biodegradability	: Inoculum: Sewage Biodegradation: 100 % Remarks: Readily biodegradable
110-19-0:	
Biodegradability	: aerobic Inoculum: Sewage Result: Readily biodegradable. Biodegradation: 81 % Exposure time: 20 d
123-86-4:	
Biodegradability	: Biodegradation: 83 % Exposure time: 28 d Method: OECD Test Guideline 301D
Chemical Oxygen De- mand (COD)	: 0.00169 mg/g
BOD/COD	: BOD/COD: 72 %
Theoritical Oxygen De- mand (ThOD)	: 0.0022 mg/g
1330-20-7:	
Biodegradability	: Inoculum: activated sludge Result: Readily biodegradable. Biodegradation: 72 % Exposure time: 20 d

100-41-4:

Biodegradability	: Inoculum: activated sludge
	Concentration: 22 mg/l
	Result: Readily biodegradable.
	Biodegradation: 70 %
	Exposure time: 28 d
	GLP: yes

142-82-5:

Biodegradability : Primary biodegradation Inoculum: activated sludge Concentration: 100 mg/l Biodegradation: 100 % Testing period: 2 d Exposure time: 25 d Remarks: Readily biodegradable

Bioaccumulative potential

Components:

67	-64-	1.	
07	-04-	.	

Partition coefficient: n- octanol/water	: log Pow: -0.24
64742-49-0: Partition coefficient: n- octanol/water	: Remarks: No data available
64742-89-8: Partition coefficient: n- octanol/water	: log Pow: 2.13 - 4.85 (25 °C)
108-65-6: Partition coefficient: n- octanol/water	: log Pow: 0.43
108-88-3: Partition coefficient: n- octanol/water	: log Pow: 2.73
110-19-0: Partition coefficient: n- octanol/water	: log Pow: 1.78
123-86-4: Bioaccumulation	: Species: Fish Bioconcentration factor (BCF): 15
Partition coefficient: n-	: log Pow: 1.82

octanol/water

1330-20-7:

Partition coefficient: n- : log Pow: 2.77 - 3.15 octanol/water

100-41-4:

Partition coefficient: n- : log Pow: 2.92 octanol/water

Mobility in soil

No data available

Other adverse effects

Product:

Regulation	40 CFR Protection of Environment; Part 82 Protection of Stratospheric Ozone - CAA Section 602 Class I Substances
Remarks	This product neither contains, nor was manufactured with a Class I or Class II ODS as defined by the U.S. Clean Air Act Section 602 (40 CFR 82, Subpt. A, App.A + B).
Additional ecological in- : formation	An environmental hazard cannot be excluded in the event of unprofessional handling or disposal., Toxic to aquatic life with long lasting effects.
<u>Components:</u>	
100-41-4:	
Results of PBT and vPvB : assessment	This substance is not considered to be persistent, bio- accumulating nor toxic (PBT). This substance is not considered to be very persistent nor very bioaccumu- lating (vPvB).

SECTION 13. DISPOSAL CONSIDERATIONS

Disposal methods Waste from residues	: Dispose of in accordance with all applicable local, state and federal regulations.
Contaminated packaging	: Empty remaining contents.

Dispose of as unused product. Do not re-use empty containers. Do not burn, or use a cutting torch on, the empty drum.

SECTION 14. TRANSPORT INFORMATION

IATA (International Air Transport Association): UN1263, PAINT RELATED MATERIAL, 3, II, Flash Point:-18 °C(-0.40 °F)

IMDG (International Maritime Dangerous Goods): UN1263, PAINT RELATED MATERIAL, 3, II

DOT (Department of Transportation): UN1263, PAINT RELATED MATERIAL, 3, II

SECTION 15. REGULATORY INFORMATION

OSHA Hazards	: Flammable liquid, Carcinogen, Harmful by skin absorption., Moderate skin irritant, Moderate eye irritant, Moderate respiratory irritant, Teratogen, Reproductive hazard, Mutagen
WHMIS Classification	: Flammable Liquid D2A: Very Toxic Material Causing Other Toxic Effects D2B: Toxic Material Causing Other Toxic Effects

EPCRA - Emergency Planning and Community Right-to-Know Act

CERCLA Reportable Quantity

Components	CAS-No.	Component RQ (lbs)	Calculated product RQ (lbs)
Mixed xylenes	1330-20-7	100	1901

SARA 304 Extremely Hazardous Substances Reportable Quantity

This material does not contain any components with a section 304 EHS RQ.

SARA 311/312	: Fire Hazard
Hazards	Chronic Health Hazard
	Acute Health Hazard

Clean Air Act

The following chemical(s) are listed as HAP under the U.S. Clean Air Act, Section 12 (40 CFR 61):

108-88-3	Toluene	10.54 %
100-41-4	Ethylbenzene	1.5983 %
71-43-2	Benzene	0.0281 %
110-54-3	Hexane	0.0031 %

67-56-1	Methanol	0.0022 %
91-20-3	Naphthalene	0.0003 %
98-82-8	Cumene	0.000 %
-	-	ed under the U.S. Clean Air Act
		(40 CFR 68.130, Subpart F).
		S. Clean Air Act Section 111 SOCMI
Intermediate or Final V		
67-64-1	Acetone	37.3599 %
108-88-3	Toluene	10.54 %
110-19-0	Isobutyl acetate	10.5389 %
123-86-4	n-Butyl acetate	5.3421 %
1330-20-7	Mixed xylenes	5.2608 %
100-41-4	Ethylbenzene	1.5983 %
110-82-7	Cyclohexane	0.3931 %
71-43-2	Benzene	0.0281 %
67-56-1	Methanol	0.0022 %
98-82-8	Cumene	0.000 %
90-02-0	Cumene	0.000 %
Clean Water Act		
The following Hazardou	is Substances are listed u	nder the U.S. CleanWater Act, Sec-
tion 311, Table 116.4A		,
108-88-3	Toluene	10.54 %
110-19-0	Isobutyl acetate	10.5389 %
123-86-4	n-Butyl acetate	5.3421 %
1330-20-7	Mixed xylenes	5.2608 %
	•	
100-41-4	Ethylbenzene	1.5983 %
110-82-7	Cyclohexane	0.3931 %
71-43-2	Benzene	0.0281 %
91-20-3	Naphthalene	0.0003 %
-	is Chemicals are listed un	der the U.S. CleanWater Act, Section
311, Table 117.3:		
108-88-3	Toluene	10.54 %
123-86-4	n-Butyl acetate	5.3421 %
1330-20-7	Mixed xylenes	5.2608 %
100-41-4	Ethylbenzene	1.5983 %
110-82-7	Cyclohexane	0.3931 %
71-43-2	Benzene	0.0281 %
91-20-3	Naphthalene	0.0003 %
This product contains t		ts listed under the U.S. Clean Water
Act Section 307	2 .	
108-88-3	Toluene	10.54 %
100-41-4	Ethylbenzene	1.5983 %
US State Regulations		
05 State Regulations	•	
Massachusetts Right	To Know	
67-64-1	Acetone	30 - 50 %
108-88-	3 Toluene	10 - 20 %
110-19-		10 - 20 %
	,	5 - 10 %
123-86-	4 n-Butyl acetate	5 - 10 %

	1330-20-7	Mixed xylenes	5 - 10 %
	100-41-4	Ethylbenzene	1 - 5 %
	71-43-2	Benzene	0 - 0.1 %
Pennsylv	vania Right To Kn	ow	
	67-64-1	Acetone	30 - 50 %
	64742-49-0	Naphtha (pet), hydrotreated It	0 - 20 %
	64742-89-8	Solvent naphtha (pet), lt aliph.	0 - 20 %
	68410-97-9	Distillates, pet, lt dist hydrotreat process, low-boil	0 - 20 %
	108-65-6	Glycol ether PM acetate	10 - 20 %
	108-88-3	Toluene	10 - 20 %
	110-19-0	Isobutyl acetate	10 - 20 %
	123-86-4	n-Butyl acetate	5 - 10 %
	1330-20-7	Mixed xylenes	5 - 10 %
	100-41-4	Ethylbenzene	1 - 5 %
	110-82-7	Cyclohexane	0.1 - 1 %
	71-43-2	Benzene	0 - 0.1 %
New Jers	sey Right To Know	N	
	67-64-1	Acetone	30 - 50 %
	64742-49-0	Naphtha (pet), hydrotreated It	0 - 20 %
	64742-89-8	Solvent naphtha (pet), lt aliph.	0 - 20 %
	68410-97-9	Distillates, pet, lt dist hydrotreat process, low-boil	0 - 20 %
	108-65-6	Glycol ether PM acetate	10 - 20 %
	108-88-3	Toluene	10 - 20 %
	110-19-0	Isobutyl acetate	10 - 20 %

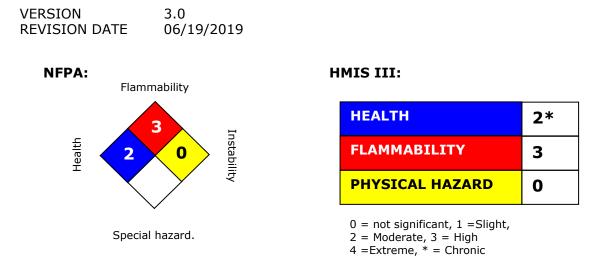
123-86-4n-Butyl acetate5 - 10 %1330-20-7Mixed xylenes5 - 10 %100-41-4Ethylbenzene1 - 5 %

California Prop 65	WARNING! This product contains a chemical known to the State of California to cause cancer.
100-41-4	Ethylbenzene
71-43-2	Benzene
91-20-3	Naphthalene
98-82-8	Cumene
	WARNING: This product contains a chemical known to the State of California to cause birth defects or other reproductive harm.
108-88-3	Toluene
71-43-2 67-56-1	Benzene Methanol

The components of this product are reported in the following inventories:Switzerland. New notified substances and declared:y (positive listing)

preparations		(The formulation contains substances listed on the Swiss Inventory)
United States TSCA Inventory	:	y (positive listing) (On TSCA Invento- ry)
Canadian Domestic Substances List (DSL)	:	y (positive listing) (All components of this product are on the Canadian DSL.)
Australia Inventory of Chemical Substances (AICS)	:	y (positive listing) (On the inventory, or in compliance with the inventory)
New Zealand. Inventory of Chemical Substances	:	n (Negative listing) (Not in compliance with the inventory)
Japan. ENCS - Existing and New Chemical Substances Inventory	:	n (Negative listing) (Not in compliance with the inventory)
Japan. ISHL - Inventory of Chemical Substances (METI)	:	n (Negative listing) (Not in compliance with the inventory)
Korea. Korean Existing Chemicals Inventory (KECI)	:	y (positive listing) (On the inventory, or in compliance with the inventory)
Philippines Inventory of Chemicals and Chemical Substances (PICCS)	:	y (positive listing) (On the inventory, or in compliance with the inventory)
China. Inventory of Existing Chemical Substances in China (IECSC)	:	y (positive listing) (On the inventory, or in compliance with the inventory)

SECTION 16. OTHER INFORMATION



The information accumulated is based on the data of which we are aware and is believed to be correct as of the date hereof. Since this information may be applied under conditions beyond our control and with which we may be unfamiliar and since data made become available subsequently to the date hereof, we do not assume any responsibility for the results of its use. Recipients are advised to confirm in advance of need that the information is current, applicable, and suitable to their circumstances.

Legecy MSDS: R0329927

Material number: 16069388, 547005, 146398

Key or leg	Key or legend to abbreviations and acronyms used in the safety data sheet				
ACGIH	American Conference of Gov-	LD50	Lethal Dose 50%		
	ernment Industrial Hygienists				
AICS	Australia, Inventory of Chem-	LOAEL	Lowest Observed Adverse Effect		
	ical Substances		Level		
DSL	Canada, Domestic Substanc-	NFPA	National Fire Protection Agency		
	es List				
NDSL	Canada, Non-Domestic Sub-	NIOSH	National Institute for Occupational		
	stances List		Safety & Health		
CNS	Central Nervous System	NTP	National Toxicology Program		
CAS	Chemical Abstract Service	NZIoC	New Zealand Inventory of Chemicals		
EC50	Effective Concentration	NOAEL	No Observable Adverse Effect Level		
EC50	Effective Concentration 50%	NOEC	No Observed Effect Concentration		
EGEST	EOSCA Generic Exposure	OSHA	Occupational Safety & Health Admin-		

	Scenario Tool		istration
EOSCA	European Oilfield Specialty Chemicals Association	PEL	Permissible Exposure Limit
EINECS	European Inventory of Exist- ing Chemical Substances	PICCS	Philipines Inventory of Commercial Chemical Substances
MAK	Germany Maximum Concen- tration Values	PRNT	Presumed Not Toxic
GHS	Globally Harmonized System	RCRA	Resource Conservation Recovery Act
>=	Greater Than or Equal To	STEL	Short-term Exposure Limit
IC50	Inhibition Concentration 50%	SARA	Superfund Amendments and Reau- thorization Act.
IARC	International Agency for Re- search on Cancer	TLV	Threshold Limit Value
IECSC	Inventory of Existing Chemi- cal Substances in China	TWA	Time Weighted Average
ENCS	Japan, Inventory of Existing and New Chemical Substanc- es	TSCA	Toxic Substance Control Act
KECI	Korea, Existing Chemical In- ventory	UVCB	Unknown or Variable Compositon, Complex Reaction Products, and Biological Materials
<=	Less Than or Equal To	WHMIS	Workplace Hazardous Materials In- formation System
LC50		Lethal Concentration 50%	