The Toro Company

Toro Wheel Horse Red Paint

The Toro Company

Chemwatch: **5230-88** Version No: **4.1** Safety Data Sheet according to OSHA HazCom Standard (2012) requirements

SECTION 1 Identification

Product Identifier

Product name	Toro Wheel Horse Red Paint
Chemical Name	Not Applicable
Synonyms	Part number: 5340, Product Code: N-5831
Proper shipping name	Paint including paint, lacquer, enamel, stain, shellac solutions, varnish, polish, liquid filler and liquid lacquer base; Paint related material including paint thinning, drying, removing, or reducing compound
Chemical formula	Not Applicable
Other means of identification	Not Available

Recommended use of the chemical and restrictions on use

Relevant identified uses Use according to manufacturer's directions.

Name, address, and telephone number of the chemical manufacturer, importer, or other responsible party

Registered company name	The Toro Company
Address	8111 Lyndale Avenue South, Bloomington MN 55420 United States
Telephone	+1-952-888-8801
Fax	+1-952-887-8258
Website	www.toro.com
Email	HealthAndSafety@toro.com

Emergency phone number

Association / Organisation	CHEMTEL	CHEMWATCH EMERGENCY RESPONSE (24/7)
Emergency telephone numbers	1-800-255-3924	+1 855-237-5573
Other emergency telephone numbers	+1-813-248-0585	+61 3 9573 3188

Once connected and if the message is not in your preferred language then please dial 01

Una vez conectado y si el mensaje no está en su idioma preferido, por favor marque 02

SECTION 2 Hazard(s) identification

Classification of the substance or mixture

Chemwatch Hazard Ratings

	Min	Max	
Flammability	2		
Toxicity	1		0 = Minimum
Body Contact	3	1	1 = Low
Reactivity	1		2 = Moderate
Chronic	3	i	3 = High 4 = Extreme

NFPA 704 diamond



Note: The hazard category numbers found in GHS classification in section 2 of this SDSs are NOT to be used to fill in the NFPA 704 diamond. Blue = Health Red = Fire Yellow = Reactivity White = Special (Oxidizer or water reactive substances)

Classification

Flammable Liquids Category 3, Aspiration Hazard Category 1, Skin Corrosion/Irritation Category 2, Sensitisation (Skin) Category 1, Serious Eye Damage/Eye Irritation Category 1, Sensitisation (Respiratory) Category 1, Specific Target Organ Toxicity - Single Exposure (Narcotic Effects) Category 3, Germ Cell Mutagenicity Category 1A, Carcinogenicity Category 1B, Hazardous to the Aquatic Environment Acute Hazard Category 3, Hazardous to the Aquatic Environment Long-Term Hazard Category 3

Label elements



Chemwatch Hazard Alert Code: 3

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Hazard statement(s)

H226	Flammable liquid and vapour.
H304	May be fatal if swallowed and enters airways.
H315	Causes skin irritation.
H317	May cause an allergic skin reaction.
H318	Causes serious eye damage.
H334	May cause allergy or asthma symptoms or breathing difficulties if inhaled.
H336	May cause drowsiness or dizziness.
H340	May cause genetic defects.
H350	May cause cancer.
H412	Harmful to aquatic life with long lasting effects.

Hazard(s) not otherwise classified

Not Applicable

Precautionary statement(s) Prevention

P201	Obtain special instructions before use.	
P210	Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking.	
P233	Keep container tightly closed.	
P261	Avoid breathing mist/vapours/spray.	
P271	Use only outdoors or in a well-ventilated area.	
P280	Wear protective gloves, protective clothing, eye protection and face protection.	

Precautionary statement(s) Response

P301+P310	IF SWALLOWED: Immediately call a POISON CENTER/doctor/physician/first aider.	
P331	Do NOT induce vomiting.	
P304+P340	IF INHALED: Remove person to fresh air and keep comfortable for breathing.	
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.	
P310	Immediately call a POISON CENTER/doctor/physician/first aider.	

Precautionary statement(s) Storage

P403+P235	Store in a well-ventilated place. Keep cool.	
P405	Store locked up.	
P403+P233	Store in a well-ventilated place. Keep container tightly closed.	

Precautionary statement(s) Disposal

P501 Dispose of contents/container to authorised hazardous or special waste collection point in accordance with any local regulation.

SECTION 3 Composition / information on ingredients

Substances

See section below for composition of Mixtures

Mixtures

CAS No	%[weight]	Name
1330-20-7	45.54	xylene
85-44-9	11.64	phthalic anhydride, unregulated
77-85-0	5-10	trimethylolethane
100-41-4	8.03	ethylbenzene
108-65-6	1-5	propylene glycol monomethyl ether acetate, alpha-isomer
1332-37-2	1-5	red iron oxide
64742-95-6	1-5	naphtha petroleum, light aromatic solvent
13586-82-8	0.159	cobalt 2-ethylhexanoate

SECTION 4 First-aid measures

Description of first aid measures		
Eye Contact	 If this product comes in contact with the eyes: Immediately hold eyelids apart and flush the eye continuously with running water. Ensure complete irrigation of the eye by keeping eyelids apart and away from eye and moving the eyelids by occasionally lifting the upper and lower lids. Continue flushing until advised to stop by the Poisons Information Centre or a doctor, or for at least 15 minutes. Transport to hospital or doctor without delay. 	

	Removal of contact lenses after an eye injury should only be undertaken by skilled personnel.
Skin Contact	 If skin contact occurs: Immediately remove all contaminated clothing, including footwear. Flush skin and hair with running water (and soap if available). Seek medical attention in event of irritation. For thermal burns: Decontaminate area around burn. Consider the use of cold packs and topical antibiotics. For first-degree burns (affecting polyayer of skin) Hold burned skin under cool (not cold) running water or immerse in cool water until pain subsides. Use compresses if running water is not available. Cover with sterile non-adhesive bandage or clean cloth. Do NOT apply butter or ointments; this may cause infection. Goive over-the counter pain relievers if pain increases or swelling, redness, fever occur. For second-degree burns (affecting top two layers of skin) Cool the burn by immerse in cold running water for 10-15 minutes. Use compresses if running water is not available. Do NOT apply ice as this may lower body temperature and cause further damage. Do NOT preak blisters or apply butter or ointments; this may cause infection. Protect burn by cover toosely with sterile, nonstick bandage and secure in place with gauze or tape. To prevent shock: (unless the person has a head, neck, or leg injury, or it would cause discomfort): Lay the person flat. Elevate burn area above heart level, if possible. Cover the derial at sistance. For third-degree burns Seek medical area cover loosely with sterile, nonstick bandage or, for large areas, a sheet or other material that will not leave lint in wound. Seek medical area acover loosely with sterile, nonstick bandage or, for large areas, a sheet or other material that will not leave lint in wound. Seek medical area tower loosely with sterile, nonstick bandage or for large areas, a sheet or other material that will not leave lint in wo
Inhalation	 If fumes or combustion products are inhaled remove from contaminated area. Lay patient down. Keep warm and rested. Prostheses such as false teeth, which may block airway, should be removed, where possible, prior to initiating first aid procedures. Apply artificial respiration if not breathing, preferably with a demand valve resuscitator, bag-valve mask device, or pocket mask as trained. Perform CPR if necessary. Transport to hospital, or doctor, without delay.
Ingestion	 If swallowed do NOT induce vomiting. If vomiting occurs, lean patient forward or place on left side (head-down position, if possible) to maintain open airway and prevent aspiration. Observe the patient carefully. Never give liquid to a person showing signs of being sleepy or with reduced awareness; i.e. becoming unconscious. Give water to rinse out mouth, then provide liquid slowly and as much as casualty can comfortably drink. Seek medical advice. Avoid giving milk or oils. Avoid giving alcohol. If spontaneous vomiting appears imminent or occurs, hold patient's head down, lower than their hips to help avoid possible aspiration of vomitus.

Most important symptoms and effects, both acute and delayed

See Section 11

Indication of any immediate medical attention and special treatment needed

Any material aspirated during vomiting may produce lung injury. Therefore emesis should not be induced mechanically or pharmacologically. Mechanical means should be used if it is considered necessary to evacuate the stomach contents; these include gastric lavage after endotracheal intubation. If spontaneous vomiting has occurred after ingestion, the patient should be monitored for difficult breathing, as adverse effects of aspiration into the lungs may be delayed up to 48 hours. Treat symptomatically.

SECTION 5 Fire-fighting measures

Extinguishing media

- Foam.
- Dry chemical powder.
- BCF (where regulations permit).
- Carbon dioxide.
- Water spray or fog Large fires only.

Special hazards arising from the substrate or mixture

Fire Incompatibility	Avoid contamination with oxidising agents i.e. nitrates, oxidising acids, chlorine bleaches, pool chlorine etc. as ignition may result
Special protective equipment a	nd precautions for fire-fighters

	Alert Fire Brigade and tell them location and nature of hazard.
	May be violently or explosively reactive.
Eiro Eighting	Wear breathing apparatus plus protective gloves.
rire righting	Prevent, by any means available, spillage from entering drains or water course.
	If safe, switch off electrical equipment until vapour fire hazard removed.
	Use water delivered as a fine spray to control fire and cool adjacent area.

Fire/Explosion Hazard	 Liquid and vapour are flammable. Moderate fire hazard when exposed to heat or flame. Vapour forms an explosive mixture with air. Moderate explosion hazard when exposed to heat or flame. Vapour may travel a considerable distance to source of ignition. Heating may cause expansion or decomposition leading to violent rupture of containers. Combustion products include: carbon dioxide (CO2) other pyrolysis products typical of burning organic material. carbon monoxide (CO)
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SECTION 6 Accidental release measures

Personal precautions, protective equipment and emergency procedures

See section 8

Environmental precautions

See section 12

Methods and material for containment and cleaning up

Minor Spills	 Remove all ignition sources. Clean up all spills immediately. Avoid breathing vapours and contact with skin and eyes. Control personal contact with the substance, by using protective equipment. Contain and absorb small quantities with vermiculite or other absorbent material. Wipe up.
Major Spills	 Clear area of personnel and move upwind. Alert Fire Brigade and tell them location and nature of hazard. May be violently or explosively reactive. Wear breathing apparatus plus protective gloves. Prevent, by any means available, spillage from entering drains or water course. Consider evacuation (or protect in place).

Personal Protective Equipment advice is contained in Section 8 of the SDS.

SECTION 7 Handling and storage

Precautions for safe handling

Safe handling	 Containers, even those that have been emptied, may contain explosive vapours. Do NOT cut, drill, grind, weld or perform similar operations on or near containers. DO NOT allow clothing wet with material to stay in contact with skin Electrostatic discharge may be generated during pumping - this may result in fire. Ensure electrical continuity by bonding and grounding (earthing) all equipment. Restrict line velocity during pumping in order to avoid generation of electrostatic discharge (<=1 m/sec until fill pipe submerged to twice its diameter, then <= 7 m/sec). Avoid splash filling. Do NOT use compressed air for filling discharging or handling operations. Wait 2 minutes after tank filling (for tanks such as those on road tanker vehicles) before opening hatches or manholes. Avoid all personal contact, including inhalation. Wear protective clothing when risk of overexposure occurs. Use in a well-ventilated area. Prevent concentration in hollows and sumps. DO NOT enter confined spaces until atmosphere has been checked. Avoid smoking, naked lights or ignition sources.
Other information	 Store in original containers in approved flammable liquid storage area. Store away from incompatible materials in a cool, dry, well-ventilated area. DO NOT store in pits, depressions, basements or areas where vapours may be trapped. No smoking, naked lights, heat or ignition sources. Storage areas should be clearly identified, well illuminated, clear of obstruction and accessible only to trained and authorised personnel - adequate security must be provided so that unauthorised personnel do not have access. Store according to applicable regulations for flammable materials for storage tanks, containers, piping, buildings, rooms, cabinets, allowable quantities and minimum storage distances.

Conditions for safe storage, including any incompatibilities

Suitable container	 Packing as supplied by manufacturer. Plastic containers may only be used if approved for flammable liquid. Check that containers are clearly labelled and free from leaks. For low viscosity materials (i) : Drums and jerry cans must be of the non-removable head type. (ii) : Where a can is to be used as an inner package, the can must have a screwed enclosure. For materials with a viscosity of at least 2680 cSt. (23 deg. C) For manufactured product having a viscosity of at least 250 cSt. (23 deg. C) Manufactured product that requires stirring before use and having a viscosity of at least 20 cSt (25 deg. C): (i) Removable head packaging; (ii) Cans with friction closures and (iii) low pressure tubes and cartridges may be used. Where combination packages are used, and the inner packages are of glass, there must be sufficient inert cushioning material in contact with inner and outer packages In addition, where inner packagings are glass and contain liquids of packing group I there must be sufficient inert absorbent to absorb any spillage, unless the outer packaging is a close fitting moulded plastic box and the substances are not incompatible with the plastic.
Storage incompatibility	Avoid reaction with oxidising agents

SECTION 8 Exposure controls / personal protection

Control parameters

Occupational Exposure Limits (OEL)

INGREDIENT DATA

	Source	Ingredient	Material name		TWA	STEL	Peak	Notes
	US OSHA Permissible Exposure Limits (PELs) Table Z-1	xylene	Xylenes (o-, m-, p	-isomers)	100 ppm / 435 mg/m3	Not Available	Not Available	Not Available
	US OSHA Permissible Exposure Limits (PELs) Table Z-1	phthalic anhydride, unregulated	Phthalic anhydride	9	2 ppm / 12 mg/m3	Not Available	Not Available	Not Available
	US NIOSH Recommended Exposure Limits (RELs)	phthalic anhydride, unregulated	Phthalic anhydride	9	1 ppm / 6 mg/m3	Not Available	Not Available	Not Available
	US OSHA Permissible Exposure Limits (PELs) Table Z-1	ethylbenzene	Ethyl benzene		100 ppm / 435 mg/m3	Not Available	Not Available	Not Available
	US NIOSH Recommended Exposure Limits (RELs)	ethylbenzene	Ethyl benzene		100 ppm / 435 mg/m3	545 mg/m3 / 125 ppm	Not Available	Not Available
	US OSHA Permissible Exposure Limits (PELs) Table Z-1	red iron oxide	Rouge- Respirable	e fraction	5 mg/m3	Not Available	Not Available	Not Available
	US OSHA Permissible Exposure Limits (PELs) Table Z-1	red iron oxide	Iron oxide- (fume)		10 mg/m3	Not Available	Not Available	Not Available
	US OSHA Permissible Exposure Limits (PELs) Table Z-1	red iron oxide	Rouge- Total dust		15 mg/m3	Not Available	Not Available	Not Available
	US OSHA Permissible Exposure Limits (PELs) Table Z-3	red iron oxide	Inert or Nuisance fraction	Dust: Respirable	5 mg/m3 / 15 mppcf	Not Available	Not Available	Not Available
	US OSHA Permissible Exposure Limits (PELs) Table Z-3	red iron oxide	Inert or Nuisance	Dust: Total Dust	15 mg/m3 / 50 mppcf	Not Available	Not Available	Not Available
	US NIOSH Recommended Exposure Limits (RELs)	red iron oxide	Iron oxide dust an	d fume (as Fe)	5 mg/m3	Not Available	Not Available	Not Available
	US NIOSH Recommended Exposure Limits (RELs)	red iron oxide	Rouge		Not Available	Not Available	Not Available	See Appendix D
	US OSHA Permissible Exposure Limits (PELs) Table Z-1	cobalt 2-ethylhexanoate	Particulates Not C (PNOR)- Respirat	Otherwise Regulated	5 mg/m3	Not Available	Not Available	Not Available
	US OSHA Permissible Exposure Limits (PELs) Table Z-1	cobalt 2-ethylhexanoate	Particulates Not C (PNOR)- Total dus	therwise Regulated	15 mg/m3	Not Available	Not Available	Not Available
	US OSHA Permissible Exposure Limits (PELs) Table Z-3	cobalt 2-ethylhexanoate	Inert or Nuisance fraction	Dust: Respirable	5 mg/m3 / 15 mppcf	Not Available	Not Available	Not Available
	US OSHA Permissible Exposure Limits (PELs) Table Z-3	cobalt 2-ethylhexanoate	Inert or Nuisance	Dust: Total Dust	15 mg/m3 / 50 mppcf	Not Available	Not Available	Not Available
	US NIOSH Recommended Exposure Limits (RELs)	cobalt 2-ethylhexanoate	Particulates not otherwise regulated		Not Available	Not Available	Not Available	See Appendix D
j	Emergency Limits							
	Ingredient	TEEL-1		TEEL-2		TEEL-3		
Ĩ	xylene	Not Available		Not Available		Not Available		

Ingredient	IEEL-1	TEEL-2		TEEL-3	
xylene	Not Available Not Available			Not Available	
phthalic anhydride, unregulated	18 mg/m3 56 mg/m3			10,000 mg/m3	
ethylbenzene	Not Available	Not Available		Not Available	
propylene glycol monomethyl ether acetate, alpha-isomer	Not Available	Not Available		Not Available	
red iron oxide	15 mg/m3	360 mg/m3		2,200 mg/m3	
naphtha petroleum, light aromatic solvent	1,200 mg/m3	6,700 mg/m3		40,000 mg/m3	
Ingredient	Original IDLH		Revised IDLH		
xylene	900 ppm		Not Available		
phthalic anhydride, unregulated	60 mg/m3		Not Available		
trimethylolethane	Not Available		Not Available		
ethylbenzene	800 ppm		Not Available		
propylene glycol monomethyl ether acetate, alpha-isomer	Not Available		Not Available		
red iron oxide	2,500 mg/m3		Not Available		
naphtha petroleum, light aromatic solvent	Not Available		Not Available		
cobalt 2-ethylhexanoate	Not Available		Not Available		

Occupational Exposure Banding

Ingredient	Occupational Exposure Band Rating	Occupational Exposure Band Limit
naphtha petroleum, light aromatic solvent	E	≤ 0.1 ppm

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Toro Wheel Horse Red Paint

Ingredient	Occupational Exposure Band Rating	Occupational Exposure Band Limit		
Notes:	Occupational exposure banding is a process of assignir adverse health outcomes associated with exposure. The range of exposure concentrations that are expected to p	ng chemicals into specific categories or bands based on a chemical's potency and the e output of this process is an occupational exposure band (OEB), which corresponds to a protect worker health.		
xposure controls				
Appropriate engineering controls	Engineering controls are used to remove a hazard or place a barrier between the worker and the hazard. Well-designed engineering controls can be highly effective in protecting workers and will typically be independent of worker interactions to provide this high level of protection. The basic types of engineering controls are: Process controls which involve changing the way a job activity or process is done to reduce the risk. Enclosure and/or isolation of emission source which keeps a selected hazard "physically" away from the worker and ventilation that strategically "adds" and "removes" air in the work environment. Ventilation can remove or dilute an air contaminant if designed properly. The design of a ventilation system must match the particular process and chemical or contaminant in use. CARE: Use of a quantity of this material in confined space or poorly ventilated area, where rapid build up of concentrated atmosphere may occur, could require increased ventilation and/or protective gear			
Individual protection measures, such as personal protective equipment				
Eye and face protection	 Safety glasses with side shields. Chemical goggles. Contact lenses may pose a special hazard; soft con the wearing of lenses or restrictions on use, should and adsorption for the class of chemicals in use and their removal and suitable equipment should be real 	tact lenses may absorb and concentrate irritants. A written policy document, describing be created for each workplace or task. This should include a review of lens absorption d an account of injury experience. Medical and first-aid personnel should be trained in dily available.		
Skin protection	See Hand protection below			
Hands/feet protection	 Wear crientical protective gloves, e.g. PVC. Wear safety footwear or safety gumboots, e.g. Rubi NOTE: The material may produce skin sensitisation in pred equipment, to avoid all possible skin contact. Contaminated leather items, such as shoes, belts a The selection of suitable gloves does not only depend o manufacturer. Where the chemical is a preparation of se and has therefore to be checked prior to the application. The exact break through time for substances has to be a making a final choice. Personal hygiene is a key element of effective hand car washed and dried thoroughly. 	ber isposed individuals. Care must be taken, when removing gloves and other protective nd watch-bands should be removed and destroyed. In the material, but also on further marks of quality which vary from manufacturer to everal substances, the resistance of the glove material can not be calculated in advance obtained from the manufacturer of the protective gloves and has to be observed when e. Gloves must only be worn on clean hands. After using gloves, hands should be		
Body protection	See Other protection below			
Other protection	 Employees working with confirmed human carcinog (smocks, coveralls, or long-sleeved shirt and pants) national equivalent] Employees engaged in handling operations involvin respirators with filters for dusts, mists and fumes, or be substituted. [AS/NZS 1715 or national equivalen Emergency deluge showers and eyewash fountains level with locations where direct exposure is likely. Prior to each exit from an area containing confirmed clothing and equipment at the point of exit and at th the point of exit for purposes of decontamination or labels. For maintenance and decontamination activi wear clean, impervious garments, including gloves, Prior to removing protective garments the employee garments and hood. Overalls. PVC Apron. PVC Apron. PVC Apron. Ensure there is ready access to a safety shower. Some plastic personal protective equipment (PPE) electricity. For large scale or continuous use wear tight-weave Non sparking safety or conductive footwear should conductive compound chemically bound to the bott static electricity from the body to reduce the possibi 500,000 ohms. Conductive shoes should be stored 	ens should be provided with, and be required to wear, clean, full body protective clothing , shoe covers and gloves prior to entering the regulated area. [AS/NZS ISO 6529:2006 or g carcinogens should be provided with, and required to wear and use half-face filter-type air purifying canisters or cartridges. A respirator affording higher levels of protection may t] , supplied with potable water, should be located near, within sight of, and on the same d human carcinogens, employees should be required to remove and leave protective e last exit of the day, to place used clothing and equipment in impervious containers at disposal. The contents of such impervious containers must be identified with suitable ties, authorized employees entering the area should be provided with and required to boots and continuous-air supplied hood. e should undergo decontamination and be required to shower upon removal of the vere. (e.g. gloves, aprons, overshoes) are not recommended as they may produce static non-static clothing (no metallic fasteners, cuffs or pockets). be considered. Conductive footwear describes a boot or shoe with a sole made from a om components, for permanent control to electrically ground the foot an shall dissipate lity of ignition of volatile compounds. Electrical resistance must range between 0 to in lockers close to the room in which they are worn.		

Recommended material(s)

GLOVE SELECTION INDEX

Glove selection is based on a modified presentation of the: **"Forsberg Clothing Performance Index".** The effect(s) of the following substance(s) are taken into account in the *computer*generated selection: Toro Wheel Horse Red Paint

Material	CPI
TEFLON	А
VITON	A

Respiratory protection

Type A Filter of sufficient capacity. (AS/NZS 1716 & 1715, EN 143:2000 & 149:2001, ANSI Z88 or national equivalent)

Where the concentration of gas/particulates in the breathing zone, approaches or exceeds the "Exposure Standard" (or ES), respiratory protection is required. Degree of protection varies with both face-piece and Class of filter; the nature of protection varies with Type of filter.

Required Minimum	Half-Face	Full-Face	Powered Air
Protection Factor	Respirator	Respirator	Respirator
up to 10 x ES	A-AUS	-	A-PAPR-AUS / Class 1

BUTYL	С
BUTYL/NEOPRENE	С
HYPALON	С
NAT+NEOPR+NITRILE	С
NATURAL+NEOPRENE	С
NEOPRENE	С
NEOPRENE/NATURAL	С
NITRILE	С
NITRILE+PVC	С
PE/EVAL/PE	С
PVA	С
PVC	С
PVDC/PE/PVDC	С

* CPI - Chemwatch Performance Index

A: Best Selection

B: Satisfactory; may degrade after 4 hours continuous immersion

C: Poor to Dangerous Choice for other than short term immersion

NOTE: As a series of factors will influence the actual performance of the glove, a final selection must be based on detailed observation. -

* Where the glove is to be used on a short term, casual or infrequent basis, factors such as "feel" or convenience (e.g. disposability), may dictate a choice of gloves which might otherwise be unsuitable following long-term or frequent use. A qualified practitioner should be consulted.

SECTION 9 Physical and chemical properties

Information on basic physical and chemical properties

 up to 50 x ES
 A-AUS / Class 1

 up to 100 x ES
 A-2
 A-PAPR-2^

^ - Full-face

A(All classes) = Organic vapours, B AUS or B1 = Acid gasses, B2 = Acid gas or hydrogen cyanide(HCN), B3 = Acid gas or hydrogen cyanide(HCN), E = Sulfur dioxide(SO2), G = Agricultural chemicals, K = Ammonia(NH3), Hg = Mercury, NO = Oxides of nitrogen, MB = Methyl bromide, AX = Low boiling point organic compounds(below 65 degC)

- Cartridge respirators should never be used for emergency ingress or in areas of unknown vapour concentrations or oxygen content.
- The wearer must be warned to leave the contaminated area immediately on detecting any odours through the respirator. The odour may indicate that the mask is not functioning properly, that the vapour concentration is too high, or that the mask is not properly fitted. Because of these limitations, only restricted use of cartridge respirators is considered appropriate.
- Cartridge performance is affected by humidity. Cartridges should be changed after 2 hr of continuous use unless it is determined that the humidity is less than 75%, in which case, cartridges can be used for 4 hr. Used cartridges should be discarded daily, regardless of the length of time used

Appearance	Flammable liquid with typical odour.		
Physical state	Liquid	Relative density (Water = 1)	0.9846897
Odour	Not Available	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	Not Available
pH (as supplied)	Not Applicable	Decomposition temperature (°C)	Not Available
Melting point / freezing point (°C)	Not Available	Viscosity (cSt)	Not Available
Initial boiling point and boiling range (°C)	136	Molecular weight (g/mol)	Not Applicable
Flash point (°C)	29 (CC)	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Flammable.	Oxidising properties	Not Available
Upper Explosive Limit (%)	12.6	Surface Tension (dyn/cm or mN/m)	Not Available
Lower Explosive Limit (%)	0.8	Volatile Component (%vol)	Not Available
Vapour pressure (kPa)	Not Available	Gas group	Not Available
Solubility in water	Not Applicable	pH as a solution (1%)	Not Available
Vapour density (Air = 1)	>1	VOC g/L	Not Available

SECTION 10 Stability and reactivity

Reactivity	See section 7
Chemical stability	 Unstable in the presence of incompatible materials. Product is considered stable. Hazardous polymerisation will not occur.
Possibility of hazardous reactions	See section 7
Conditions to avoid	See section 7
Incompatible materials	See section 7
Hazardous decomposition products	See section 5

SECTION 11 Toxicological information

Information on toxicological effects

Inhaled

The material can cause respiratory irritation in some persons. The body's response to such irritation can cause further lung damage. Inhalation of vapours may cause drowsiness and dizziness. This may be accompanied by sleepiness, reduced alertness, loss of reflexes, lack of

	co-ordination, and vertigo. Inhalation hazard is increased at higher temperatures. Inhalation of high concentrations of gas/vapour causes lung irritation with coughing and nausea, central nervous depression with headache and dizziness, slowing of reflexes, fatigue and inco-ordination. Inhalation of vapours or aerosols (mists, fumes), generated by the material during the course of normal handling, may be damaging to the health of the individual.			
Ingestion	Swallowing of the liquid may cause aspiration into the lungs with the risk of chemical pneumonitis; serious consequences may result. (ICSC13733) Accidental ingestion of the material may be damaging to the health of the individual.			
Skin Contact	The material may cause moderate inflammation of the skin either following direct contact or after a delay of some time. Repeated exposure can cause contact dermatitis which is characterised by redness, swelling and blistering. Open cuts, abraded or irritated skin should not be exposed to this material Entry into the blood-stream, through, for example, cuts, abrasions or lesions, may produce systemic injury with harmful effects. Examine the skin prior to the use of the material and ensure that any external damage is suitably protected. Skin contact with the material may damage the health of the individual; systemic effects may result following absorption.			
Eye	If applied to the eyes, this material causes severe eye damage. The liquid produces a high level of eye discomfort and is capable of possible permanent impairment of vision, if not promptly and adequa	causing pain and severe conjunctivitis. Corneal injury may develop, with tely treated.		
Chronic	Long-term exposure to respiratory irritants may result in airways disease, involving difficulty breathing and related whole-body problems. Inhaling this product is more likely to cause a sensitisation reaction in some persons compared to the general population. Skin contact with the material is more likely to cause a sensitisation reaction in some persons compared to the general population. There is ample evidence that this material can be regarded as being able to cause cancer in humans based on experiments and other information. Substance accumulation, in the human body, may occur and may cause some concern following repeated or long-term occupational exposure. Chronic solvent inhalation exposures may result in nervous system impairment and liver and blood changes. [PATTYS]			
	τοχιζιτγ	IRRITATION		
Toro Wheel Horse Red Paint	Not Available	Not Available		
	ΤΟΧΙCITY	IRRITATION		
	Dermal (rabbit) D50: >1700 mg/kg ^[2]	Eve (human): 200 ppm irritant		
	Inholation/Pat) LCE0: 5000 ppm/h[2]	Eve (rabbit): 5 mg/24b SEVEPE		
	Oral (Mayres) LDE0: 0440 are/us ^[2]			
xyiene				
		Eye: adverse effect observed (irritating) ^[1]		
		Skin (rabbit):500 mg/24h moderate		
		Skin: adverse effect observed (irritating)[1]		
	ΤΟΧΙΟΙΤΥ	IRRITATION		
phthalic anhydride,	Dermal (rabbit) LD50: >10000 mg/kg ^[2]	Eye (rabbit): 100 mg - SEVERE		
unregulated	Inhalation(Rat) LC50: >2.14 mg/l4h ^[1]	Eye (rabbit): 50 mg/24h - MOD		
	Oral (Cat) LD50; 800 mg/kg ^[2]	Skin (rabbit): 500 mg/24h - mild		
	ΤΟΧΙΟΙΤΥ	IRRITATION		
	Dermal (rabbit) LD50: >10000 mg/kg ^[1]	Eye: no adverse effect observed (not irritating) ^[1]		
trimethylolethane	Inhalation(Rat) LC50: >0.85 mg/l4h ^[1]	Skin: no adverse effect observed (not irritating) ^[1]		
	Oral (Rat) LD50: >2500 mg/kg ^[1]			
	ΤΟΧΙΟΙΤΥ	IRRITATION		
	Dermal (rabbit) LD50: 17800 mg/kg ^[2]	Eye (rabbit): 500 mg - SEVERE		
ethvlbenzene	Inhalation(Rat) LC50: 17.2 mg/l4h ^[2]	Eve: no adverse effect observed (not irritating) ^[1]		
	Oral (Rat) LD50: 3500 mg/kg ^[2]	Skin (rabbit): 15 mg/24h mild		
		Skin: no adverse effect observed (not irritating) ^[1]		
	ΤΟΧΙΟΙΤΥ	IRRITATION		
propylene glycol monomethyl ether acetate, alpha-isomer	dermal (rat) D50: >2000 mg/kg ^[1]	Eve: no adverse effect observed (not irritating) ^[1]		
	Oral (Rat) LD50: 3739 mg/kg ^[2]	Skin: no adverse effect observed (not irritating) ^[1]		
red iron oxide	ΤΟΧΙΟΙΤΥ			
	Oral (Rat) LD50: >5000 mg/kg ^[2]	Eye (rabbit): non-irritant		
		Skin (rabbit): non-irritant 24h		

naphtha petroleum, light aromatic solvent ΤΟΧΙΟΙΤΥ

Dermal (rabbit) LD50: >1900 mg/kg^[1]

Inhalation(Rat) LC50: >4.42 mg/L4h^[1]

Oral (Rat) LD50: >4500 mg/kg^[1]

Eye: no adverse effect observed (not irritating)^[1] Skin: adverse effect observed (irritating)^[1]

IRRITATION

Mutagenicity

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Toro Wheel Horse Red Paint

	TOXICITY IRRITATION			
	dermal (rat) LD50: >2000 mg/kg ^[1] Eye: adverse effect observed (irritating) ^[1]		fect observed (irritating) ^[1]	
cobait 2-ethylnexanoate	Inhalation(Rat) LC50: >2.5 mg/L4h ^[2] Skin: no adverse effect observed (not irritating) ^[1]			
	Oral (Rat) LD50: 3129 mg/kg ^[1]			
Legend:	 d: 1. Value obtained from Europe ECHA Registered Substances - Acute toxicity 2. Value obtained from manufacturer's SDS. Unless otherwise specified data extracted from RTECS - Register of Toxic Effect of chemical Substances 			
	Poproductive offector in rate			
XYLENE	The substance is classified by IARC as Group 3: NOT classifiable as to its carcinogenicity to humans. Evidence of carcinogenicity may be inadequate or limited in animal testing.			
PHTHALIC ANHYDRIDE, UNREGULATED	Allergic reactions involving the respiratory tract are usually due to interactions between IgE antibodies and allergens and occur rapidly. Allergic potential of the allergen and period of exposure often determine the severity of symptoms. Some people may be genetically more prone than others, and exposure to other irritants may aggravate symptoms. Allergy causing activity is due to interactions with proteins. Attention should be paid to atopic diathesis, characterised by increased susceptibility to nasal inflammation, asthma and eczema. Exogenous allergic alveolitis is induced essentially by allergen specific immune-complexes of the IgG type; cell-mediated reactions (T lymphocytes) may be involved. Such allergy is of the delayed type with onset up to four hours following exposure.			
ETHYLBENZENE	Liver changes, utheral tract, effects on fertility, foetotoxicity, specific developmental abnormalities (musculoskeletal system) recorded. Ethylbenzene is readily absorbed when inhaled, swallowed or in contact with the skin. It is distributed throughout the body, and passed out through urine. It may irritate the skin, eyes and may cause hearing loss if exposed to high doses. Long Term exposure may cause damage to the kidney, liver and lungs, including a tendency to cancer formation, according to animal testing. There is no research on its effect on sex organs and unborn babies. NOTE: Substance has been shown to be mutagenic in at least one assay, or belongs to a family of chemicals producing damage or change to cellular DNA.			
	WARNING: This substance has been classified by the IARC as	Group 2B: Possibly Carcin	ogenic to Humans.	
PROPYLENE GLYCOL MONOMETHYL ETHER ACETATE, ALPHA-ISOMER	A BASF report (in ECETOC) showed that inhalation exposure to 545 ppm PGMEA (beta isomer) was associated with a teratogenic response in rabbits; but exposure to 145 ppm and 36 ppm had no adverse effects. The beta isomer of PGMEA comprises only 10% of the commercial material, the remaining 90% is alpha isomer. Hazard appears low but emphasizes the need for care in handling this chemical. [I.C.I] *Shin-Etsu SDS			
NAPHTHA PETROLEUM, LIGHT AROMATIC SOLVENT	 * [Devoe]. For trimethylbenzenes: Absorption of 1,2,4-trimethylbenzene occurs after exposure by swallowing, inhalation, or skin contact. In the workplace, inhalation and skin contact are the most important routes of absorption; whole-body toxic effects from skin absorption are unlikely to occur as the skin irritation caused by the chemical generally leads to quick removal. The substance is fat-soluble and may accumulate in fatty tissues. It is also bound to red blood cells in the bloodstream. It is excreted from the body both by exhalation and in the urine. Acute toxicity: Direct contact with liquid 1,2,4-trimethylbenzene is irritating to the skin, and breathing the vapour is irritating to the airway, causing lung inflammation. Breathing high concentrations of the chemical vapour causes headache, fatigue and drowsiness. For C9 aromatics (typically trimethylbenzenes – TMBs) Acute toxicity: Animal testing shows that semi-lethal concentrations and doses vary amongst this group. The semilethal concentrations for inhalation range from 6000 to 10000 mg/cubic metre for C9 aromatic naphtha and 18000-24000 mg/cubic metre for 1,2,4- and 1,3,5-TMB, respectively. Irritation and sensitization: Results from animal testing indicate that C9 aromatic hydrocarbon solvents are mildly to moderately irritating to the skin, minimally irritating to the eye, and have the potential to irritate the airway and cause depression of breathing rate. There is no evidence that it sensitizes skin. Repeated dose toxicity: Animal studies show that chronic inhalation toxicity for C9 aromatic hydrocarbon solvents is slight. Similarly, oral exposure does not appear to pose a high toxicity hazard for pure trimethylbenzene isomers. Mutation-causing ability: No evidence of mutation-causing ability and genetic toxicity was found in animal and laboratory testing. Reproductive and developmental toxicity: No definitive effects on reproduction were seen, although reduc			
COBALT 2-ETHYLHEXANOATE	No significant acute toxicological data identified in literature se	irch.		
XYLENE & PHTHALIC ANHYDRIDE, UNREGULATED & ETHYLBENZENE	The material may produce severe irritation to the eye causing pronounced inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis. The material may cause skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of vesicles, scaling and thickening of the skin.			
PHTHALIC ANHYDRIDE, UNREGULATED & COBALT 2-ETHYLHEXANOATE	The following information refers to contact allergens as a group and may not be specific to this product. Contact allergies quickly manifest themselves as contact eczema, more rarely as urticaria or Quincke's oedema. The pathogenesis of contact eczema involves a cell-mediated (T lymphocytes) immune reaction of the delayed type. Other allergic skin reactions, e.g. contact urticaria, involve antibody-mediated immune reactions. The significance of the contact allergen is not simply determined by its sensitisation potential: the distribution of the substance and the opportunities for contact with it are equally important. A weakly sensitising substance which is widely distributed can be a more important allergen than one with stronger sensitising potential with which few individuals come into contact.			
PHTHALIC ANHYDRIDE, UNREGULATED & NAPHTHA PETROLEUM, LIGHT AROMATIC SOLVENT	Asthma-like symptoms may continue for months or even years after exposure to the material ends. This may be due to a non-allergic condition known as reactive airways dysfunction syndrome (RADS) which can occur after exposure to high levels of highly irritating compound. Main criteria for diagnosing RADS include the absence of previous airways disease in a non-atopic individual, with sudden onset of persistent asthma-like symptoms within minutes to hours of a documented exposure to the irritant. Other criteria for diagnosis of RADS include a reversible airflow pattern on lung function tests, moderate to severe bronchial hyperreactivity on methacholine challenge testing, and the lack of minimal lymphocytic inflammation, without eosinophila. RADS (or asthma) following an irritating inhalation is an infrequent disorder with rates related to the concentration of and duration of exposure to the irritating substance. On the other hand, industrial bronchitis is a disorder that occurs as a result of exposure due to high concentrations of irritating substance (often particles) and is completely reversible after exposure ceases.			
Acute Toxicity	×	Carcinogenicity	✓	
Skin Irritation/Corrosion	✓ Reproductivity X		×	
Serious Eye Damage/Irritation	*	STOT - Single Exposure	✓	
Respiratory or Skin sensitisation	✓ S1	OT - Repeated Exposure	×	

Aspiration Hazard

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Legena:

Toro Wheel Horse Red Paint

SECTION 12 Ecological information

Foxicity					
	Endpoint	Test Duration (hr)	Species	Value	Source
Toro Wheel Horse Red Paint	Not Available	Not Available	Not Available	Not Available	Not Available
	Endpoint	Test Duration (hr)	Species	Value	Source
	LC50	96h	Fish	2.6mg/l	2
xylene	EC50	72h	Algae or other aquatic plants	4.6mg/l	2
	EC50	48h	Crustacea	1.8mg/l	2
	NOEC(ECx)	73h	Algae or other aquatic plants	0.44mg/	2
	Endpoint	Test Duration (hr)	Species	Value	Source
	EC50(ECx)	96h	Algae or other aquatic plants	0.06-0.35mg/	4
nbthalic anhydride	EC50	96h	Algae or other aquatic plants	0.06-0.35mg/	4
unregulated	EC50	72h	Algae or other aquatic plants	68mg/l	2
	LC50	96h	Fish	>99ma/l	2
	EC50	48h	Crustacea	71mg/l	2
	Endpoint	Test Duration (hr)	Species	Value	Source
	EC0(ECx)	48h	Crustacea	>=102mg/	2
trimethylolethane	EC50	72h	Algae or other aquatic plants	>1000mg/	2
unnetnylolethane	1.050	96b	Fish	>1000mg/	2
	EC50	48h	Crustacea	13000mg/	2
					-
ethylbenzene	Endpoint	Test Duration (hr)	Species	Value	Source
	LC50	96h	Fish	3.381-4.075mg/L	4
	EC50	72h	Algae or other aquatic plants	2.4-9.8mg/l	4
	EC50	48h	Crustacea	1.37-4.4mg/l	4
	EC50(ECx)	24h	Algae or other aquatic plants	0.02-938mg/l	4
	EC50	960	Algae or other aquatic plants	1.7-7.6mg/i	4
	Endpoint	Test Duration (hr)	Species	Value	Source
	LC50	96h	Fish	100mg/l	1
propylene glycol monomethyl	EC50	72h	Algae or other aquatic plants	>1000mg/	2
ether acetate, alpha-isomer	EC50	48h	Crustacea	373mg/l	2
	NOEC(ECx)	336h	Fish	47.5mg/l	2
	EC50	96h	Algae or other aquatic plants	Algae or other aquatic plants >1000mg	
	Endpoint	Test Duration (hr)	Species	Value	Source
	LC50	96h	Fish	0.05mg/l	2
red iron oxide	EC50	72h	Algae or other aquatic plants	18mg/l	2
	EC50	48h	Crustacea	>100mg/	2
	NOEC(ECx)	504h	Fish	0.52mg/l	2
	Endpoint	Test Duration (hr)	Species	Value	Source
	NOEC(ECx)	72h	Algae or other aquatic plants	1mg/l	1
naphtha petroleum, light	EC50	72h	Algae or other aquatic plants	19mg/l	1
aromatic solvent	EC50	96h	Algae or other aquatic plants	64mg/l	2
	EC50	48h	Crustacea	6.14mg/	1
	Endpoint	Test Duration (hr)	Species	Value	Source
	ErC50	72h	Algae or other aquatic plants	0.6542mg/l	2
	LC50	96h	Fish	0.8mg/l	2
cobalt 2-ethylhexanoate	EC50	72h	Algae or other aquatic plants	0.0288ma/l	2
	EC50	48h	Crustacea	0.241mg/l	2
	EC50	96h	Algae or other aquatic plants	10.8mg/l	2
	EC10(ECx)	168h	Algae or other aquatic plants	0.00123mg/	2

Extracted from 1. IUCLID Toxicity Data 2. Europe ECHA Registered Substances - Ecotoxicological Information - Aquatic Toxicity 4. US EPA,

Ecotox database - Aquatic Toxicity Data 5. ECETOC Aquatic Hazard Assessment Data 6. NITE (Japan) - Bioconcentration Data 7. METI (Japan) - Bioconcentration Data 8. Vendor Data

Harmful to aquatic organisms, may cause long-term adverse effects in the aquatic environment. **DO NOT** discharge into sewer or waterways.

Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
xylene	HIGH (Half-life = 360 days)	LOW (Half-life = 1.83 days)
phthalic anhydride, unregulated	LOW (Half-life = 0.02 days)	HIGH (Half-life = 201.96 days)
trimethylolethane	LOW	LOW
ethylbenzene	HIGH (Half-life = 228 days)	LOW (Half-life = 3.57 days)
propylene glycol monomethyl ether acetate, alpha-isomer	LOW	LOW

Bioaccumulative potential

Ingredient	Bioaccumulation
xylene	MEDIUM (BCF = 740)
phthalic anhydride, unregulated	HIGH (BCF = 3169)
trimethylolethane	LOW (LogKOW = -0.3022)
ethylbenzene	LOW (BCF = 79.43)
propylene glycol monomethyl ether acetate, alpha-isomer	LOW (LogKOW = 0.56)

Mobility in soil

Ingredient	Mobility
phthalic anhydride, unregulated	LOW (KOC = 10.84)
trimethylolethane	HIGH (KOC = 1)
ethylbenzene	LOW (KOC = 517.8)
propylene glycol monomethyl ether acetate, alpha-isomer	HIGH (KOC = 1.838)

SECTION 13 Disposal considerations

Waste treatment methods	
Product / Packaging disposal	 Containers may still present a chemical hazard/ danger when empty. Return to supplier for reuse/ recycling if possible. Otherwise: If container can not be cleaned sufficiently well to ensure that residuals do not remain or if the container cannot be used to store the same product, then puncture containers, to prevent re-use, and bury at an authorised landfill. Where possible retain label warnings and SDS and observe all notices pertaining to the product. Legislation addressing waste disposal requirements may differ by country, state and/ or territory. Each user must refer to laws operating in their area. In some areas, certain wastes must be tracked. A Hierarchy of Controls seems to be common - the user should investigate: Reduction Reuse Recycling Disposal (if all else fails) This material may be possible to reclaim the product by filtration, distillation or some other means. Shelf life considerations should also be applied in making decisions of this type. D NOT allow wash water from cleaning or process equipment to enter drains. It may be necessary to collect all wash water for treatment before disposal. In all cases disposal to sever may be subject to local laws and regulations and these should be considered first. Where in doubt contact the responsible authority. Recycle wherever possible. Consult manufacturer for recycling options or consult local or regional waste management authority for disposal if no suitable treatment or disposal facility can be identified. Dispose of by: burial in a land-fill specifically licensed to accept chemical and / or pharmaceutical wastes or Incineration in a licensed apparatus (after admixture with suitable combustible material). Decontaminate empty containers. Observe all label safeguards until containers are cleaned and destroyed.

SECTION 14 Transport information

Labels Required



Marine Pollutant

Land transport (DOT)

UN number or ID number	1263		
UN proper shipping name	Paint including paint, lacquer, enamel, stain, shellac solutions, varnish, polish, liquid filler and liquid lacquer base; Paint related material including paint thinning, drying, removing, or reducing compound		
Transport hazard class(es)	Class Subsidiary risk	Class 3 Subsidiary risk Not Applicable	
Packing group			
Environmental hazard	Not Applicable		
Special precautions for user	Hazard Label	3 367, B1, B52, B131, IB3, T2, TP1, TP29	

Air transport (ICAO-IATA / DGR)

UN number	1263			
UN proper shipping name	Paint related material (including paint thinning or reducing compounds); Paint (including paint, lacquer, enamel, stain, shellac, varnish, polish, liquid filler and liquid lacquer base)			
Transport hazard class(es)	ICAO/IATA Class ICAO / IATA Subrisk ERG Code	3 Not Applicable 3L		
Packing group	III			
Environmental hazard	Not Applicable			
Special precautions for user	Special provisions Cargo Only Packing Instructions Cargo Only Maximum Qty / Pack Passenger and Cargo Packing Instructions		A3 A72 A192 366 220 L 355	
	Passenger and Cargo Maximum Qty / Pack		60 L	
	Passenger and Cargo Limited Quantity Packing Instructions		Y344	
	Passenger and Cargo Limited Maximum Qty / Pack		10 L	

Sea transport (IMDG-Code / GGVSee)

UN number	1263		
UN proper shipping name	PAINT (including paint, lacquer, enamel, stain, shellac, varnish, polish, liquid filler and liquid lacquer base) or PAINT RELATED MATERIAL (including paint thinning or reducing compound)		
Transport hazard class(es)	IMDG Class 3 IMDG Subrisk N	lot Applicable	
Packing group	Ш		
Environmental hazard	Not Applicable		
Special precautions for user	EMS Number Special provisions Limited Quantities	F-E, S-E 163 223 367 955 5 L	

Transport in bulk according to Annex II of MARPOL and the IBC code Not Applicable

Transport in bulk in accordance with MARPOL Annex V and the IMSBC Code

Product name	Group
xylene	Not Available
phthalic anhydride, unregulated	Not Available
trimethylolethane	Not Available
ethylbenzene	Not Available
propylene glycol monomethyl ether acetate, alpha-isomer	Not Available
red iron oxide	Not Available
naphtha petroleum, light aromatic solvent	Not Available
cobalt 2-ethylhexanoate	Not Available

Product name	Ship Type
xylene	Not Available
phthalic anhydride, unregulated	Not Available
trimethylolethane	Not Available
ethylbenzene	Not Available
propylene glycol monomethyl ether acetate, alpha-isomer	Not Available
red iron oxide	Not Available
naphtha petroleum, light aromatic solvent	Not Available
cobalt 2-ethylhexanoate	Not Available

SECTION 15 Regulatory information

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

xylene is found on the following regulatory lists

US DOE Temporary Emergency Exposure Limits (TEELs)

cobalt 2-ethylhexanoate is found on the following regulatory lists

International Agency for Research on Cancer (IARC) - Agents Classified by the IARC US Declamporary Emergency Exposure Limits (TEELs) Wonographs - Not Classified as Carcinogenic US Declamporary Emergency Exposure Limits (TEELs) US - California Hazardous Air Pollutants Identified as Toxic Air Contaminants US Declamporary Emergency Exposure Limits (TEELs) US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs) US SOFIA Permissible Exposure Limits (TEELs) US Clean Air Act - Hazardous Air Pollutants Identified as Toxic Air Contaminants US EPCRA Section 313 Chemical List US - California Hazardous Air Pollutants Identified as Toxic Air Contaminants US EPCRA Section 313 Chemical List US Clean Air Act - Hazardous Air Pollutants Identified as Toxic Air Contaminants US EPCRA Section 313 Chemical List US Clean Air Act - Hazardous Air Pollutants Identified as Toxic Air Contaminants US PCRA Section 313 Chemical List US DoE Temporary Emergency Exposure Limits (TEELs) US NOSH Recommended Exposure Limits (RELs) US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory US TSCA Chemical Substance Inventory - Interim List of Active Substance International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Informational Agency for Research on Cancer (IARC) - Agents Classified by the IARC Informational Agency for Research on Cancer (IARC) - Agents Classified by the IARC Informational Agency for Research on Cancer (IARC) - Agents Classified by the IARC Informationa Proposition 65 - Carcin	tory inces ory inces
Monographs - Not Classified as Carcinogenic UB - Califormia Hazardous Air Pollutants Identified as Toxic Air Contaminants US EPCRA Section 313 Chemical List US - Califormia Hazardous Air Pollutants Identified as Toxic Air Contaminants US SOHA Permissible Exposure Limits (FELs) Table Z-1 US Clean Air Act - Hazardous Air Pollutants Identified as Toxic Air Contaminants US EPCRA Section 313 Chemical List US - California Hazardous Air Pollutants Identified as Toxic Air Contaminants US EPCRA Section 313 Chemical List US - California Hazardous Air Pollutants Identified as Toxic Air Contaminants US EPCRA Section 313 Chemical List US - California Hazardous Air Pollutants Identified as Toxic Air Contaminants US EPCRA Section 313 Chemical List US Clean Air Act - Hazardous Air Pollutants US NOSH Recommended Exposure Limits (RELs) US Doi Temporary Emergency Exposure Limits (TEELs) US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory US Toxic Substance Inventory - Interim List of Active Substance US Toxic Substances Inventory US Toxic Substance Inventory - Interim List of Active Substance US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory US Toxic Substance Inventory - Interim List of Active Substance US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory US CWA (Clean Water Act	tory ances ory ances
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US - Massachusetts - Right To Know Listed Chemicals US OSHA Permissible Exposure Limits (PELs) Table Z-1 US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs) US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory - Interim List of Active Substance Iphthalic anhydride, unregulated is found on the following regulatory lists US Cean Air Act - Hazardous Air Pollutants Identified as Toxic Air Contaminants US EPCRA Section 313 Chemical List US ODE Temporary Emergency Exposure Limits (TEELs) US OSHA Permissible Exposure Limits (PELs) Table Z-1 US DOE Temporary Emergency Exposure Limits (TEELs) US OSHA Permissible Exposure Limits (PELs) Table Z-1 US DOE Temporary Emergency Exposure Limits (TEELs) US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory - Interim List of Active Substance US TSCA Chemical Substance Inventory - Interim List of Active Substance trimethylolethane is found on the following regulatory lists US TSCA Chemical Substance Inventory - Interim List of Active Substance US - California Hazardous Air Pollutants Identified as Toxic Air Contaminants US CWA (Clean Water Act) - List of Hazardous Substances US - California Hazardous Air Pollutants Identified as Toxic Air Contaminants US CWA (Clean Water Act) - Priority Pollutants US - California Hazardous Air Pollutants Identified as Toxic Air Contaminants US California Proposition 65 - Ansignificant Risk Levels	tory ances ory inces
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International Agency for Research on Cancer (IARC) - Agents Classified by the IARC US NIOSH Recommended Exposure Limits (RELs)	
Monographs - Not Classified as Carcinogenic US OSHA Permissible Exposure Limits (PELs) Table Z-1	
International WHO List of Proposed Occupational Exposure Limit (OEL) Values for US OSHA Permissible Exposure Limits (PELs) Table Z-3	
Manufactured Nanomaterials (MNMS) US Toxic Substances Control Act (TSCA) - Chemical Substance Inventor	ory
US - Alaska Air Quality Control - Concentrations Triggering an Air Quality Episode for Air Pollutants Other Than PM-2.5	inces
US - Massachusetts - Right To Know Listed Chemicals	
US DOE Temporary Emergency Exposure Limits (TEELs)	
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International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs - Not Classified as Carcinogenic	inces

Chemical Footprint Project - Chemicals of High Concern List

International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs - Not Classified as Carcinogenic International WHO List of Proposed Occupational Exposure Limit (OEL) Values for

Manufactured Nanomaterials (MNMS)

- US Alaska Air Quality Control Concentrations Triggering an Air Quality Episode for Air Pollutants Other Than PM-2.5
- US California Hazardous Air Pollutants Identified as Toxic Air Contaminants
- US Clean Air Act Hazardous Air Pollutants
- US EPA Integrated Risk Information System (IRIS)

US EPCRA Section 313 Chemical List

Federal Regulations

Superfund Amendments and Reauthorization Act of 1986 (SARA)

Section 311/312 hazard categories

US List of Active Substances Exempt from the TSCA Inventory Notifications (Active-Inactive) Rule

US National Toxicology Program (NTP) 15th Report Part B. Reasonably Anticipated to be a Human Carcinogen

US NIOSH Recommended Exposure Limits (RELs)

- US OSHA Permissible Exposure Limits (PELs) Table Z-1
- US OSHA Permissible Exposure Limits (PELs) Table Z-3

US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory

US TSCA Chemical Substance Inventory - Interim List of Active Substances

Flammable (Gases, Aerosols, Liquids, or Solids)	Yes
Gas under pressure	No
Explosive	No
Self-heating	No
Pyrophoric (Liquid or Solid)	No
Pyrophoric Gas	No
Corrosive to metal	No
Oxidizer (Liquid, Solid or Gas)	No
Organic Peroxide	No
Self-reactive	No
In contact with water emits flammable gas	No
Combustible Dust	No
Carcinogenicity	Yes
Acute toxicity (any route of exposure)	No
Reproductive toxicity	No
Skin Corrosion or Irritation	Yes
Respiratory or Skin Sensitization	Yes
Serious eye damage or eye irritation	Yes
Specific target organ toxicity (single or repeated exposure)	Yes
Aspiration Hazard	Yes
Germ cell mutagenicity	Yes
Simple Asphyxiant	No
Hazards Not Otherwise Classified	No

US. EPA CERCLA Hazardous Substances and Reportable Quantities (40 CFR 302.4)

Name	Reportable Quantity in Pounds (Ib)	Reportable Quantity in kg
xylene	100	45.4
phthalic anhydride, unregulated	5000	2270
ethylbenzene	1000	454

State Regulations

US. California Proposition 65

WARNING: This product can expose you to chemicals including ethylbenzene, which is known to the State of California to cause cancer. For more information, go to www.P65Warnings.ca.gov.

National Inventory Status

National Inventory	Status
Australia - AIIC / Australia Non-Industrial Use	Yes
Canada - DSL	Yes
Canada - NDSL	No (xylene; phthalic anhydride, unregulated; trimethylolethane; ethylbenzene; propylene glycol monomethyl ether acetate, alpha-isomer; red iron oxide; naphtha petroleum, light aromatic solvent; cobalt 2-ethylhexanoate)
China - IECSC	Yes
Europe - EINEC / ELINCS / NLP	Yes
Japan - ENCS	Yes
Korea - KECI	Yes
New Zealand - NZIoC	Yes
Philippines - PICCS	Yes
USA - TSCA	Yes

National Inventory	Status	
Taiwan - TCSI	TCSI Yes	
Mexico - INSQ	No (trimethylolethane)	
Vietnam - NCI	Yes	
Russia - FBEPH Yes		
Legend:	Yes = All CAS declared ingredients are on the inventory No = One or more of the CAS listed ingredients are not on the inventory. These ingredients may be exempt or will require registration.	

SECTION 16 Other information

Revision Date	10/12/2021
Initial Date	02/12/2016

SDS Version Summary

Version	Date of Update	Sections Updated
3.1	01/11/2019	One-off system update. NOTE: This may or may not change the GHS classification
4.1	10/12/2021	Classification change due to full database hazard calculation/update.

Other information

Classification of the preparation and its individual components has drawn on official and authoritative sources as well as independent review by the Chernwatch Classification committee using available literature references.

The SDS is a Hazard Communication tool and should be used to assist in the Risk Assessment. Many factors determine whether the reported Hazards are Risks in the workplace or other settings. Risks may be determined by reference to Exposures Scenarios. Scale of use, frequency of use and current or available engineering controls must be considered.

Definitions and abbreviations

PC-TWA: Permissible Concentration-Time Weighted Average
PC-STEL: Permissible Concentration-Short Term Exposure Limit
IARC: International Agency for Research on Cancer
ACGIH: American Conference of Governmental Industrial Hygienists
STEL: Short Term Exposure Limit
TEEL: Temporary Emergency Exposure Limit。
IDLH: Immediately Dangerous to Life or Health Concentrations
ES: Exposure Standard
OSF: Odour Safety Factor
NOAEL :No Observed Adverse Effect Level
LOAEL: Lowest Observed Adverse Effect Level
TLV: Threshold Limit Value
LOD: Limit Of Detection
OTV: Odour Threshold Value
BCF: BioConcentration Factors
BEI: Biological Exposure Index
AIIC: Australian Inventory of Industrial Chemicals
DSL: Domestic Substances List
NDSL: Non-Domestic Substances List
IECSC: Inventory of Existing Chemical Substance in China
EINECS: European INventory of Existing Commercial chemical Substances
ELINCS: European List of Notified Chemical Substances
NLP: No-Longer Polymers
ENCS: Existing and New Chemical Substances Inventory
KECI: Korea Existing Chemicals Inventory
NZIoC: New Zealand Inventory of Chemicals
PICCS: Philippine Inventory of Chemicals and Chemical Substances
TSCA: Toxic Substances Control Act
TCSI: Taiwan Chemical Substance Inventory
INSQ: Inventario Nacional de Sustancias Químicas
NCI: National Chemical Inventory
FBEPH: Russian Register of Potentially Hazardous Chemical and Biological Substances

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