Auto Klene Solutions

Safety Data Sheet according to WHS and ADG requirements

Chemwatch: 9126549 Issue Date: 27/01/2015 Version No: 3.1.1.1 Print Date: 23/03/2016

Chemwatch Hazard Alert Code: 3

Initial Date: Not Available S.GHS.AUS.EN

SECTION 1 IDENTIFICATION OF THE SUBSTANCE / MIXTURE AND OF THE COMPANY / UNDERTAKING

Product Identifier	
Product name	Auto Klene Ferris Alloy Wheel Cleaner
Synonyms	Not Available
Proper shipping name	CORROSIVE LIQUID, TOXIC, N.O.S. (contains hydrofluoric acid and sulfuric acid)
Other means of identification	Not Available
Relevant identified uses of the	ne substance or mixture and uses advised against
Relevant identified uses	Heavy duty acid suitable for cleaning rust, alloy trays, wheels and parts.

SIGNAL WORD	DANGER
Hazard statement(s)	
H290	May be corrosive to metals.

Chemwatch: 9126549

Issue Date: 27/01/2015 Print Date: 23/03/2016

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H301 Toxic if swallowed. Details of the supplier of the safety data sheet Registered company Auto Klene Solutions name Address 1/83 Merrindale Drive VIC Croydon 3136 Australia Telephone +61 3 8761 1900 Fax +61 3 8761 1955 Website https://www.autoklene.com/msds/ Email Not Available Emergency telephone number Association / Not Available Organisation Emergency telephone 131 126 (Poisons Information Centre) numbers Other emergency 0408 406 968 (Mark Adams mobile) telephone numbers **SECTION 2 HAZARDS IDENTIFICATION**

Classification of the substance or mixture

HAZARDOUS CHEMICAL. DANGEROUS GOODS. According to the WHS Regulations and the ADG Code.

CHEMWATCH HAZARD RATINGS

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

Poisons Schedule	S7
[1] Classification	Metal Corrosion Category 1, Acute Toxicity (Oral) Category 3, Acute Toxicity (Dermal) Category 3, Acute Toxicity (Inhalation) Category 2, Skin Corrosion/Irritation Category 1A, Serious Eye Damage Category 1, Carcinogenicity Category 1A
Legend:	1. Classified by Chernwatch; 2. Classification drawn from HSIS ; 3. Classification drawn from EC Directive 1272/2008 - Annex VI
abel elements	
GHS label elements	
H311	Toxic in contact with skin.
H330	Fatal if inhaled.
H314	Causes severe skin burns and eye damage.
H318	Causes serious eye damage.
H350	May cause cancer.
recautionary statement(s) F	revention
P201	Obtain special instructions before use.
P260	Do not breathe dust/fume/gas/mist/vapours/spray.
P270	Do not eat, drink or smoke when using this product.
P271	Use only outdoors or in a well-ventilated area.
P280	Wear protective gloves/protective clothing/eye protection/face protection.
P281	Use personal protective equipment as required.
P234	Keep only in original container.
Precautionary statement(s) F	Response
P301+P310	IF SWALLOWED: Immediately call a POISON CENTER or doctor/physician.

Page 3 of 12

Chemwatch: 9126549 Version No: 3.1.1.1

Disposal Dispose of contents/container in accordance with local regulations.				
Store locked up.				
Store in a well-ventilated place. Keep container tightly closed.				
Storage				
Wash contaminated clothing before reuse.				
IF exposed or concerned: Get medical advice/attention.				
IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.				
IF INHALED: Remove victim to fresh air and keep at rest in a position comfortable for breathing.				
ON SKIN (or hair): Remove/Take off immediately all contaminated clothing. Rinse skin with water/shower.				
IF SWALLOWED: Rinse mouth. Do NOT induce vomiting.				

SECTION 3 COMPOSITION / INFORMATION ON INGREDIENTS

Substances

See section below for composition of Mixtures

Mixtures

CAS No	%[weight]	Name		
7664-39-3	<5	hydrofluoric acid		
7664-93-9	<4	sulfuric acid		
SECTION 4 FIRST AID MEASURES				

escription of first aid meas	ures
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 there is evidence of severe skin irritation or skin burns: Avoid further contact. Immediately remove contaminated clothing, including footwear. Flush skin under running water for 15 minutes. Avoiding contamination of the hands, massage calcium gluconate gel into affected areas, pay particular attention to creases in skin. Contact the Poisons Information Centre. Continue gel application for at least 15 minutes after burning sensation ceases. If pain recurs, repeat application of calcium gluconate gel or apply every 20 minutes. If no gel is available, continue washing for at least 15 minutes, using soap if available. If patient is conscious, give six calcium gluconate or calcium carbonate tablets in water by mouth. Fransport to hospital, or doctor, urgently.
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. Inhalation of vapours or aerosols (mists, fumes) may cause lung oedema.
	 Corrosive substances may cause lung damage (e.g. lung oedema, fluid in the lungs). As this reaction may be delayed up to 24 hours after exposure, affected individuals need complete rest (preferably in semi-recumbent posture) and must be kept under medical observation even if no symptoms are (yet) manifested. Before any such manifestation, the administration of a spray containing a dexamethasone derivative or beclomethasone derivative may be considered. This must definitely be left to a doctor or person authorised by him/her. (ICSC13719)
Ingestion	 For advice, contact a Poisons Information Centre or a doctor at once. Urgent hospital treatment is likely to be needed. 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. I give water to rinse out mouth, then provide liquid slowly and as much as casualty can comfortably drink. Transport to hospital or doctor without delay.

Page 4 of 12

Chemwatch: 9126549

Version No: 3.1.1.1

Following acute or short term repeated exposure to hydrofluoric acid:

Subcutaneous injections of Calcium Gluconate may be necessary around the burnt area. Continued application of Calcium Gluconate Gel or subcutaneous Calcium Gluconate should then continue for 3-4 days at a frequency of 4-6 times per day. If a "burning" sensation recurs, apply more frequently.

Systemic effects of extensive hydrofluoric acid burns include renal damage, hypocalcaemia and consequent cardiac arrhythmias. Monitor haematological, respiratory, renal, cardiac and electrolyte status at least daily. Tests should include FBE, blood gases, chest X-ray, creatinine and electrolytes, urine output, Ca ions, Mg ions and phosphate ions. Continuous ECG monitoring may be required.

- Where serum calcium is low, or clinical, or ECG signs of hypocalcaemia develop, infusions of calcium gluconate, or if less serious, oral Sandocal, should be given. Hydrocortisone 500 mg in a four to six hourly infusion may help.
- Antibiotics should not be given as a routine, but only when indicated.
- Eye contact pain may be excruciating and 2-3 drops of 0.05% pentocaine hydrochloride may be instilled, followed by further irrigation

BIOLOGICAL EXPOSURE INDEX - BEI

These represent the determinants observed in specimens collected from a healthy worker exposed at the Exposure Standard (ES or TLV):

Determinant	Index	Sampling Time	Comments		
1. Methaemoglobin in blood	1.5% of haemoglobin	During or end of shift	B, NS, SQ		
B: Background levels occur in specimens collected from subjects NOT exposed.					

NS: Non-specific determinant; Also seen after exposure to other materials

SQ: Semi-quantitative determinant - Interpretation may be ambiguous; should be used as a screening test or confirmatory test.

SECTION 5 FIREFIGHTING MEASURES

Extinguishing media

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

Special hazards arising from the substrate or mixture

Fire Incompatibility	► Avoid strong bases.
Advice for firefighters	
Fire Fighting	 Alert Fire Brigade and tell them location and nature of hazard. Wear full body protective clothing with breathing apparatus. Prevent, by any means available, spillage from entering drains or water course. Use fire fighting procedures suitable for surrounding area. Do not approach containers suspected to be hot. Cool fire exposed containers with water spray from a protected location. If safe to do so, remove containers from path of fire.
Fire/Explosion Hazard	 Non combustible. Not considered to be a significant fire risk. Acids may react with metals to produce hydrogen, a highly flammable and explosive gas. Heating may cause expansion or decomposition leading to violent rupture of rigid containers. May emit acrid smoke. May emit corrosive and poisonous fumes.

Personal precautions, protective equipment and emergency procedures

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 full body protective clothing with breathing apparatus. Prevent, by any means available, spillage from entering drains or water course. Consider evacuation (or protect in place). Stop leak if safe to do so.
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Personal Protective Equipment advice is contained in Section 8 of the SDS.

SECTION 7 HANDLING AND STORAGE

Precautions for safe handling Minor Spills • Drains for storage or use areas should have retention basins for pH adjustments and dilution of spills before discharge or disposal of material. • Check regularly for spills and leaks. 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 spill with sand, earth, inert material or vermiculite. • Wipe up. • Place in a suitable, labelled container for waste disposal.

Issue Date: 27/01/2015 Print Date: 23/03/2016

Auto Klene F	Ferris Alloy	Wheel	Cleaner
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Page 5 of 12

Chemwatch: 9126549 Version No: 3.1.1.1 Issue Date: 27/01/2015 Print Date: 23/03/2016

Safe handling	 DO NOT allow clothing wet with material to stay in contact with skin Avoid all personal contact, including inhalation. Wear protective clothing when risk of exposure occurs. Use in a well-ventilated area. WARNING: To avoid violent reaction, ALWAYS add material to water and NEVER water to material. Avoid smoking, naked lights or ignition sources. Avoid contact with incompatible materials. When handling, DO NOT eat, drink or smoke.
Other information Conditions for safe storage,	 Store in original containers. Keep containers securely sealed. Store in a cool, dry, well-ventilated area. Store away from incompatible materials and foodstuff containers. Protect containers against physical damage and check regularly for leaks. Observe manufacturer's storage and handling recommendations contained within this SDS.
Suitable container	Plastic container Plastic carboy Polyethylene or polypropylene container. Packing as recommended by manufacturer. Pcheck all containers are clearly labelled and free from leaks. DO NOT use aluminium or galvanised containers Material is corrosive to most metals, glass and other siliceous materials.
Storage incompatibility	 Inorganic acids are generally soluble in water with the release of hydrogen ions. The resulting solutions have pH's of less than 7.0. Inorganic acids neutralise chemical bases (for example: amines and inorganic hydroxides) to form salts - neutralisation can generate dangerously large amounts of heat in small spaces. The dissolution of inorganic acids in water or the dilution of their concentrated solutions with additional water may generate significant heat. The addition of water to inorganic acids often generates sufficient heat in the small region of mixing to cause some of the water to boil explosively. The resulting "bumping" can spatter the acid. Inorganic acids react with active metals, including such structural metals as aluminum and iron, to release hydrogen, a flammable gas. Reacts with mild steel, galvanised steel / zinc producing hydrogen gas which may form an explosive mixture with air. Avoid contamination of water, foodstuffs, feed or seed.

SECTION 8 EXPOSURE CONTROLS / PERSONAL PROTECTION

Control parameters

OCCUPATIONAL EXPOSURE LIMITS (OEL)

INGREDIENT DATA						
Source	Ingredient	Material name	TWA	STEL	Peak	Notes
Australia Exposure Standards	hydrofluoric acid	Hydrogen fluoride (as F)	Not Available	Not Available	2.6 mg/m3 / 3 ppm	Not Available
Australia Exposure Standards	sulfuric acid	Sulphuric acid	1 mg/m3	3 mg/m3	Not Available	Not Available

EMERGENCY LIMITS

Ingredient	Material name	TE	EL-1	TEEL-2	TEEL-3	
hydrofluoric acid	Hydrogen fluoride; (Hydrofluoric acid)	No	ot Available	Not Available	Not Available	
sulfuric acid	Sulfuric acid	No	ot Available	Not Available	Not Available	
Ingredient	Original IDLH	Original IDLH		Revised IDLH		
hydrofluoric acid	30 ppm	30 ppm		30 [Unch] ppm		
sulfuric acid	80 mg/m3 15 mg/m3					
Exposure controls			1			

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. Employers may need to use multiple types of controls to prevent employee overexposure.

Heads No. 13.1.1 Description Personal protection Image: Comparison of the protection of subable protection below Hands/Feer protection Image: Protection below Hands/Feer protection Subable protection below Other protection Subable protection below Image: Protection of subable protection protection of subable protection of subable protection of subable protection below Hands/Feer protection Subable protection below Image: Protection Suba		Auto Klene Ferris Alloy Wheel Cleaner	
Personal protection Image: Image	Chemwatch: 9126549	Page 6 of 12	Issue Date: 27/01/2015
Hands/feet protection Chemical goggles. Full face shield may be required for supplementary but never for primary protection of eyes. Contact lenses may pose a special hazard; soft contact lenses may absorb and concentrate inflants. A written policy document, describing the wearing of lenses or restrictions on use, should be created for each workplace or task. This should include a review of lens absorption and adsorption for the class of chemicals in use and an account of hirury experience. Medical and first-aid personnel in their removal and suitable equipment should be readily available. In the event of chemical exposure, begin eye irrigation immediately and remove contact lens as soon as practicable. Skin protection See Hand protection below Wear chemical protective gloves, e.g. PVC. Wear safety footwear or safety gumboots, e.g. Rubber When handling corrosive liquids, wear trousers or overails outside of boots, to avoid spills entering boots. The selection of suitable goves does not only depend on the material, but also on further marks of quality which vary from manufacturer to manufacturer. Where the chemical is a peparation of several substances, the resistance of the glove material can not be calculated in advance and has therefore to be checked prior to the application. The selection of suitable gove material, explication. The evact break through time for substances has to be obtained from the manufacturer of the protective gloves and has therefore to be checked prior to the application. The evact break through time for substances has to be obtained from the manufacturer of the glove material, eligiove thickness and substances and astreaked protocub material, eligiove thicknes and conce	Version No: 3.1.1.1		Print Date: 23/03/2016
Full face shield may be required for supplementary but never for primary protection of eyes. Contact lenses may pose a special hazard; soft contact lenses may absorb and concentrate initiants. A written policy document, describing the veating of lenses or restrictions on use, should be created for each workplace or task. This should include a review of lens absorption and adsorption for the class of chemicals in use and an account of injury experience. Medical and first-aid personnel should be trained in their removal and suitable equipment should be trained in their removal and suitable equipment should be readily available. In the event of chemical exposure, begin eye irrigation immediately and remove contact lens as soon as practicable. Skin protection See Hand protection below Wear shelly footwear or safety gumboots, e.g. PVC. Wear shelly footwear or safety gumboots, e.g. Rubber Wear chemical is protection of suitable gloves does not only depend on the material, but also on further marks of quality which vary from manufacturer to manufacturer. Where the chemical is a preparation of several substances, the resistance of the gloves and has to be observed when making a final choice. Hands/feet protection Jif exact theak through time for substances has to be obtained from the manufacturer of the protective gloves and has to be observed when making a final choice. Hands/feet protection Jif equipment should be concern and yeapenet and ye	Personal protection		
Hands/feet protection • Wear chemical protective gloves, e.g. PVC. • Wear safety footwear or safety gumboots, e.g. Rubber • When handling corrosive liquids, wear trousers or overalls outside of boots, to avoid spills entering boots. The selection of suitable gloves does not only depend on the material, but also on further marks of quality which vary from manufacturer to manufacturer. Where the chemical is a preparation of several substances, the resistance of the glove material can not be calculated in advance and has therefore to be checked prior to the application. The exact break through time for substances has to be obtained from the manufacturer of the protective gloves and.has to be observed when making a final choice. Suitability and durability of glove type is dependent on usage. Important factors in the selection of gloves include: t frequency and duration of contact, chemical resistance of glove material, glove thickness and devertity Select gloves tested to a relevant standard (e.g. Europe EN 374, US F739, AS/NZS 2161.1 or national equivalent). • When prolonged or frequently repeated contact may occur, a glove with a protection class of 5 or higher (breakthrough time greater than 240 minutes according to EN 374, AS/NZS 2161.10.1 or national equivalent) is recommended. • When only brief contact is expected, a glove with a protection class of 3 or higher (breakthrough time greater than 240 minutes according to EN 374, AS/NZS 2161.10.1 or national equivalent) is recommended. • When only brief contact is expected, a glove with a protection class of 3 or higher (breakthrough time gr		 Full face shield may be required for supplementary but never for primary protection of eyes. Contact lenses may pose a special hazard; soft contact lenses may absorb and concentrate irritants wearing of lenses or restrictions on use, should be created for each workplace or task. This should adsorption for the class of chemicals in use and an account of injury experience. Medical and first-removal and suitable equipment should be readily available. In the event of chemical exposure, because of the event of the class of the equipment should be readily available. 	l include a review of lens absorption and aid personnel should be trained in their
Hands/feet protection• Wear safety footwear or safety gumboots, e.g. Rubber • When handling corrosive liquids, wear trousers or overalls outside of boots, to avoid spills entering boots. The selection of suitable gloves does not only depend on the material, but also on further marks of quality which vary from manufacturer to manufacturer. Where the chemical is a preparation of several substances, the resistance of the glove material can not be calculated in advance and has therefore to be checked prior to the application. The exact break through time for substances has to be obtained from the manufacturer of the protective gloves and has to be observed when making a final choice. Suitability and durability of glove type is dependent on usage. Important factors in the selection of gloves include: 	Skin protection	See Hand protection below	
Other protection Overalls. PVC Apron. PVC protective suit may be required if exposure severe. Eyewash unit. 	Hands/feet protection	 Wear safety footwear or safety gumboots, e.g. Rubber When handling corrosive liquids, wear trousers or overalls outside of boots, to avoid spills entering I The selection of suitable gloves does not only depend on the material, but also on further marks of qua manufacturer. Where the chemical is a preparation of several substances, the resistance of the glove n and has therefore to be checked prior to the application. The exact break through time for substances has to be obtained from the manufacturer of the protective making a final choice. Suitability and durability of glove type is dependent on usage. Important factors in the selection of glove frequency and duration of contact, chemical resistance of glove material, glove thickness and dexterity Select gloves tested to a relevant standard (e.g. Europe EN 374, US F739, AS/NZS 2161.1 or national When prolonged or frequently repeated contact may occur, a glove with a protection class of 5 or hi minutes according to EN 374, AS/NZS 2161.10.1 or national equivalent) is recommended. When only brief contact is expected, a glove with a protection class of 3 or higher (breakthrough ti 	ality which vary from manufacturer to material can not be calculated in advance re gloves and has to be observed when es include: equivalent). igher (breakthrough time greater than 240
Other protection • PVC Apron. • PVC protective suit may be required if exposure severe. • Eyewash unit.	Body protection	See Other protection below	
	Other protection	 PVC Apron. PVC protective suit may be required if exposure severe. Eyewash unit. 	
Thermal hazards Not Available	Thermal hazards	Not Available	

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 *computergenerated* selection:

Auto Klene Acid Klene

Material	CPI
##sulfuric	acid
NATURAL RUBBER	А
NATURAL+NEOPRENE	A
NEOPRENE	A
NEOPRENE/NATURAL	A
PE	В
PVC	В
SARANEX-23	В
BUTYL/NEOPRENE	С
NAT+NEOPR+NITRILE	С
NITRILE	С
VITON/NEOPRENE	С

A: Best Selection

B: Satisfactory; may degrade after 4 hours continuous immersion

Appearance Clear green acidic liquid; mixes with water.

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 Respiratory protection

Type BE-P 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 Protection Factor	Half-Face Respirator	Full-Face Respirator	Powered Air Respirator
up to 10 x ES	BE-AUS P2	-	BE-PAPR-AUS / Class 1 P2
up to 50 x ES	-	BE-AUS / Class 1 P2	-
up to 100 x ES	-	BE-2 P2	BE-PAPR-2 P2 ^

^ - 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)

Page 7 of 12 Version No: 3.1.1.1 Issue Date: 27/01/2015 Print Date: 23/03/2016

Information on basic physical and chemical properties

Divisional state			
Physical state	Liquid	Relative density (Water = 1)	1.0
Odour	Not Available	Partition coefficient n- octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	Not Available
pH (as supplied)	<1.0	Decomposition temperature	Not Available
Melting point / freezing point (°C)	~0	Viscosity (cSt)	Not Available
Initial boiling point and boiling range (°C)	100	Molecular weight (g/mol)	Not Applicable
Flash point (°C)	Not Applicable	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Not Applicable	Oxidising properties	Not Available
Upper Explosive Limit (%)	Not Applicable	Surface Tension (dyn/cm or mN/m)	Not Available
Lower Explosive Limit (%)	Not Applicable	Volatile Component (%vol)	Not Available
Vapour pressure (kPa)	Not Available	Gas group	Not Available
Solubility in water (g/L)	Miscible	pH as a solution (1%)	Not Available
Vapour density (Air = 1)	Not Available	VOC g/L	Not Available

SECTION 10 STABILITY AND REACTIVITY

Reactivity	See section 7
Chemical stability	 Contact with alkaline material liberates heat 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

Information on toxicological effects

internation on textoologica	
Inhalec	Inhalation of vapours or aerosols (mists, fumes), generated by the material during the course of normal handling, may produce toxic effects. Corrosive acids can cause irritation of the respiratory tract, with coughing, choking and mucous membrane damage. There may be dizziness, headache, nausea and weakness.
Ingestion	Toxic effects may result from the accidental ingestion of the material; animal experiments indicate that ingestion of less than 40 gram may be fatal or may produce serious damage to the health of the individual. The material can produce chemical burns within the oral cavity and gastrointestinal tract following ingestion. Ingestion of acidic corrosives may produce burns around and in the mouth, the throat and oesophagus. Immediate pain and difficulties in swallowing and speaking may also be evident.

	Auto Klene Ferris Alloy Wheel	Cleaner		
Chemwatch: 9126549	-			
Page 8 of 12	Issue Date: 27/01/2015			
Version No: 3.1.1.1	Print Date: 23/03/2016			
Skin Contact	spread beneath skin. Skin contact with acidic corrosives may result in pain and burns; these	the skin. ft tissue and erode bone. Healing is delayed and death of tissue may continue to e may be deep with distinct edges and may heal slowly with the formation of scar ons or lesions, may produce systemic injury with harmful effects. Examine the		
Eye	If applied to the eyes, this material causes severe eye damage. Direct eye contact with acid corrosives may produce pain, tears, sensitivity to light and burns. Mild burns of the epithelia generally recover rapidly and completely.			
Chronic	cough, and inflammation of lung tissue often occurs.			
	TOXICITY	IRRITATION		
Auto Klene Acid Klene				
	Not Available	Not Available		
	TOXICITY	IRRITATION		
hydrofluoric acid				
nyuronuone acia	Inhalation (rat) LC50: 1.1 mg/L/601	Eye (human): 50 mg - SEVERE		
	Inhalation (rat) LC50: 1276 ppm/ ² h			
		1		
	TOXICITY	IRRITATION		
sulfuric acid	Inhalation (guinea pig) LC50: 0.018 mg/L/8H Inhalation (mouse) LC50: 0.32 mg/L/2H	Eye (rabbit): 1.38 mg SEVERE Eye (rabbit): 5 mg/30sec SEVERE		
	Inhalation (rat) LC50: 0.51 mg/L/2hE			
	Oral (rat) LD50: 2140 mg/kg ^{2]}			
Legend:	data extracted from RTECS - Register of Toxic Effect of chemical Sub	toxicity 2.* Value obtained from manufacturer's SDS. Unless otherwise specified ostances		
HYDROFLUORIC ACID	conjunctivitis. Asthma-like symptoms may continue for months or even years after known as reactive airways dysfunction syndrome (RADS) which can for the diagnosis of RADS include the absence of preceding respiratory disease, in a nu minutes to hours of a documented exposure to the irritant. A reversibl bronchial hyperreactivity on methacholine challenge testing and the la included in the criteria for diagnosis of RADS. RADS (or asthma) follo concentration of and duration of exposure to the irritating substance. I exposure due to high concentrations of irritating substance (often part	nced inflammation. Repeated or prolonged exposure to irritants may produce exposure to the material ceases. This may be due to a non-allergenic condition occur following exposure to high levels of highly irritating compound. Key criteria on-atopic individual, with abrupt onset of persistent asthma-like symptoms within e airflow pattern, on spirometry, with the presence of moderate to severe ick of minimal lymphocytic inflammation, without eosinophilia, have also been wing an irritating inhalation is an infrequent disorder with rates related to the ndustrial bronchitis, on the other hand, is a disorder that occurs as result of iculate in nature) and is completely reversible after exposure ceases. The 'he material may produce respiratory tract irritation, and result in damage to the pour)		
SULFURIC ACID	known as reactive airways dysfunction syndrome (RADS) which can for the diagnosis of RADS include the absence of preceding respiratory disease, in a m minutes to hours of a documented exposure to the irritant. A reversibl bronchial hyperreactivity on methacholine challenge testing and the la included in the criteria for diagnosis of RADS. RADS (or asthma) follo concentration of and duration of exposure to the irritating substance. I exposure due to high concentrations of irritating substance (often part disorder is characterised by dyspnea, cough and mucus production. V by the IARC as Group 1: CARCINOGENIC TO HUMANS Occupation			
Acute Toxicity	*	Carcinogenicity		
Skin Irritation/Corrosion	Reproductivity			
Damage/Irritation	✓ STO	T - Single Exposure		

Chemwatch: 9126549

Page 9 of 12

Version No: 3.1.1.1

Issue Date: 27/01/2015

Print Date: 23/03/2016

Respiratory or Skin sensitisation	0	STOT - Repeated Exposure	0
Mutagenicity	0	Aspiration Hazard	0
		Legend: 🗙	- Data available but does not fill the criteria for classification

- Data required to make classification available - Data Not Available to make classification

SECTION 12 ECOLOGICAL INFORMATION

Toxicity

Ingredient	Endpoint	Test Duration (hr)	Species	Value	Source
hydrofluoric acid	LC50	96	Fish	51mg/L	2
hydrofluoric acid	EC50	48	Crustacea	97mg/L	2
hydrofluoric acid	EC50	96	Crustacea	10.5mg/L	2
hydrofluoric acid	NOEC	504	Crustacea	3.7mg/L	2
hydrofluoric acid	EC50	96	Algae or other aquatic plants	43mg/L	2
sulfuric acid	EC50	48	Crustacea	=42.5mg/L	1
sulfuric acid	EC50	240	Algae or other aquatic plants	2.5000mg/L	4
sulfuric acid	LC50	96	Fish	=8mg/L	1
sulfuric acid	NOEC	1560	Fish	0.025mg/L	2
sulfuric acid	EC50	72	Algae or other aquatic plants	>100mg/L	2
Legend:	V3.12 - Aquatic Toxicity Da (Japan) -		CHA Registered Substances - Ecotoxicolog x database - Aquatic Toxicity Data 5. ECET tration Data 8. Vendor Data		-

Prevent, by any means available, spillage from entering drains or water courses. DO NOT discharge into sewer or waterways.

Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
	No Data available for all ingredients	No Data available for all ingredients
Bioaccumulative potential		
Ingredient	Bioaccumulation	
	No Data available for all ingredients	

Mobility in soil

Ingredient	Mobility
	No Data available for all ingredients

SECTION 13 DISPOSAL CONSIDERATIONS

Waste treatment methods

	Auto Klene Ferris Alloy Wheel Cleaner
Chemwatch: 9126549	
Page 10 of 12	Issue Date: 27/01/2015
/ersion No: 3.1.1.1	Print Date: 23/03/2016
Product / Packaging disposal SECTION 14 TRANSPOR	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 recycled if unused, or if it has not been contaminated so as to make it unsuitable for its intended use. If it has been contaminated, it 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. Note that properties of a material may change in use, and recycling or reuse may not always be appropriate. • DO 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 sewer 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. Treat and neutralise at an approved treatment plant. Treatment should involve: Neutralisation with soda-ash or soda-lime followed by: burial in a land-fill specifically licenced to accept chemical and / or pharmaceutical wastes or Incineration in a licenced apparatus (after admixture with suitable combustible material). Decontaminate empty containers with 5% aqueous sodium hydroxide or soda ash, followed by water. Observe all label safeguards until containers are cleaned and destroyed.

SECTION 14 TRANSPORT INFORMATION

Labels Required		
	CORROSVE 8 6	
Marine Pollutant	NO	
HAZCHEM	2X	
Land transport (ADG)		
UN number	2922	
Packing group	II	
UN proper shipping name	CORROSIVE LIQUID, TOXIC, N.O.S. (contains hydrofluoric acid and sulfuric acid)	
Environmental hazard	Not Applicable	
Transport hazard class(es)	Class 8 Subrisk 6.1	
Special precautions for user	Special provisions 274 Limited quantity 1 L	
Air transport (ICAO-IATA / D	GR)	
UN number	2922	
Packing group	П	
UN proper shipping name	Corrosive liquid, toxic, n.o.s. * (contains hydrofluoric acid and sulfuric acid)	
Environmental hazard	Not Applicable	
	ICAO/IATA Class 8	
Transport hazard class(es)	ICAO / IATA Subrisk 6.1	
	ERG Code 8P	

Print Date: 23/03/2016

Auto Klene Ferris Alloy Wheel Cleaner

Version No: 3.1.1.1

Special precautions for user	Special provisions	A3A803	
	Cargo Only Packing Instructions	855	
	Cargo Only Maximum Qty / Pack	30 L	
	Passenger and Cargo Packing Instructions	851	
	Passenger and Cargo Maximum Qty / Pack	1 L	
	Passenger and Cargo Limited Quantity Packing Instructions Y840		
	Passenger and Cargo Limited Maximum Qty / Pack	0.5 L	

Sea transport (IMDG-Code / GGVSee)

UN number	2922		
Packing group	П		
UN proper shipping name	CORROSIVE LIQUID, TOXIC, N.O.S. (contains hydrofluoric acid and sulfuric acid)		
Environmental hazard	Not Applicable		
Transport hazard class(es)	IMDG Class 8 IMDG Subrisk 6.1		
Special precautions for user	EMS NumberF-A, S-BSpecial provisions274Limited Quantities1 L		

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

SECTION 15 REGULATORY INFORMATION

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

 HYDROFLUORIC ACID(7664-39-3) IS FOUND ON THE FOLLOWING REGULATORY LISTS

 Australia Exposure Standards
 Australia Inventory of Chemical Substances (AICS)

Australia Hazardous Substances Information System - Consolidated Lists		International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs
Australia Exposure Standards		International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs
SULFURIC ACID(7664-93-9) IS FOUND ON THE FOLLOWING REGULATORY LISTS		nonographo
		Australia Hazardous Substances Information System - Consolidated Lists
Australia Inventory of Chemical Substances (AICS)		International Air Transport Association (IATA) Dangerous Goods Regulations - Prohibited List
		Passenger and Cargo Aircraft
National Inventory	Status	
Australia - AICS	Y	
Canada - DSL Y		
Canada - NDSL	N (sulfuric acid; hydrofluoric acid)	
China - IECSC Y		
Europe - EINEC / ELINCS / NLP Y		
Japan - ENCS	Y	
Korea - KECI	Y	
New Zealand - NZIoC	Y	
Philippines - PICCS Y		
USA - TSCA Y		

	Auto Klene Ferris Alloy Wheel Cleaner
Version No: 3.1.1.1	Print Date: 23/03/20
Legend:	Y = All ingredients are on the inventory N = Not determined or one or more ingredients are not on the inventory and are not exempt from listing(see specific ingredients in brackets)
SECTION 16 OTH	

Other information

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

A list of reference resources used to assist the committee may be found at:

www.chemwatch.net

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.