Master Instruments Pty Ltd

Chemwatch: **36-8741** Version No: **4.1.1.1**

Safety Data Sheet according to WHS and ADG requirements

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

Product Identifier

Product name	Scooter Xtras XT100G, XT75G, XT55G, XT50G, XT36G, XT20G
Synonyms	Valve Regulated Lead Acid (VRLA) Batteries
Proper shipping name	BATTERIES, WET, NON-SPILLABLE, electric storage
Other means of identification	Not Available

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

Delevent identified was	Lead acid Battery. NOTE: The finished battery complies with special provision A67 and is NOT classified as dangerous goods. MSDS risk codes and the chemical hazards are related to the battery contents if shipped separately. Batteries must not be opened, heat, crush, disassemble or short circuit.
Relevant identified uses	Chemical materials are stored in sealed metal case. The toxic properties of the electrode materials are hazardous only if the materials are released by damaging the cell or if exposed to fire. The sealed battery is not hazardous in normal use.

Details of the supplier of the safety data sheet

Registered company name	Master Instruments Pty Ltd
Address	13 Sheridan Close Milperra NSW 2214 Australia
Telephone	+61 2 9519 1200
Fax	+612 9519 4604
Website	Not Available
Email	vic@master-instruments.com.au

Emergency telephone number

Association / Organisation	Master Instruments Pty Ltd	CHEMWATCH EMERGENCY RESPONSE
Emergency telephone numbers	+61 2 9186 1132 (24hrs)	+61 1800 951 288
Other emergency telephone numbers	+61 2 9186 1132 (24hrs)	+61 2 9186 1132

SECTION 2 HAZARDS IDENTIFICATION

Poisons Schedule	Not Applicable
Classification ^[1]	Metal Corrosion Category 1, Acute Toxicity (Oral) Category 4, Acute Toxicity (Inhalation) Category 4, Skin Corrosion/Irritation Category 1A, Serious Eye Damage Category 1, Carcinogenicity Category 1A, Reproductive Toxicity Category 1A, Specific target organ toxicity - repeated exposure Category 2, Chronic Aquatic Hazard Category 1
Legend:	1. Classified by Chemwatch; 2. Classification drawn from HCIS; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI

Label elements



SIGNAL WORD	DANGER
Hazard statement(s)	
H290	May be corrosive to metals.
H302	Harmful if swallowed.
H332	Harmful if inhaled.
H314	Causes severe skin burns and eye damage.
H350	May cause cancer.
H360Df	May damage the unborn child. Suspected of damaging fertility.
H373	May cause damage to organs through prolonged or repeated exposure.
H410	Very toxic to aquatic life with long lasting effects.

Precautionary statement(s) Prevention

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P201	Obtain special instructions before use.
P260	Do not breathe dust/fume/gas/mist/vapours/spray.
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.
P270	Do not eat, drink or smoke when using this product.
P273	Avoid release to the environment.

Precautionary statement(s) Response

P301+P330+P331	IF SWALLOWED: Rinse mouth. Do NOT induce vomiting.
P303+P361+P353	IF ON SKIN (or hair): Remove/Take off immediately all contaminated clothing. Rinse skin with water/shower.
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 or doctor/physician.
P321	Specific treatment (see advice on this label).
P363	Wash contaminated clothing before reuse.
P390	Absorb spillage to prevent material damage.
P391	Collect spillage.
P301+P312	IF SWALLOWED: Call a POISON CENTER or doctor/physician if you feel unwell.
P304+P340	IF INHALED: Remove victim to fresh air and keep at rest in a position comfortable for breathing.

Precautionary statement(s) Storage

Store locked up.

Precautionary statement(s) Disposal

P501 Dispos

P405

Dispose of contents/container in accordance with local regulations.

SECTION 3 COMPOSITION / INFORMATION ON INGREDIENTS

Substances

See section below for composition of Mixtures

Mixtures

CAS No	%[weight]	Name
Not Available		sealed metal container with
7439-92-1	45-60	lead
1309-60-0	15-25	lead dioxide
Not Available		electrolyte as
7664-93-9	15-20	sulfuric acid
7440-70-2	<0.06	calcium
7440-31-5	<0.6	<u>tin</u>
7440-38-2	<0.0006	arsenic
Not Available	5-10	non hazardous other materials

SECTION 4 FIRST AID MEASURES

Description of first aid measures

Eye Contact	 If this product comes in contact with the eyes: Wash out immediately with fresh 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. Seek medical attention without delay; if pain persists or recurs seek medical attention. Removal of contact lenses after an eye injury should only be undertaken by skilled personnel.
Skin Contact	 If skin or hair contact occurs: Immediately flush body and clothes with large amounts of water, using safety shower if available. Quickly remove all contaminated clothing, including footwear. Wash skin and hair with running water. Continue flushing with water until advised to stop by the Poisons Information Centre. Transport to hospital, or doctor.
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	 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. 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.

For acute or short term repeated exposures to strong acids:

- Airway problems may arise from laryngeal edema and inhalation exposure. Treat with 100% oxygen initially.
- Respiratory distress may require cricothyroidotomy if endotracheal intubation is contraindicated by excessive swelling
- > Intravenous lines should be established immediately in all cases where there is evidence of circulatory compromise.
- Strong acids produce a coagulation necrosis characterised by formation of a coagulum (eschar) as a result of the dessicating action of the acid on proteins in specific tissues.
- INGESTION:
- Immediate dilution (milk or water) within 30 minutes post ingestion is recommended.
- DO NOT attempt to neutralise the acid since exothermic reaction may extend the corrosive injury.
- Be careful to avoid further vomit since re-exposure of the mucosa to the acid is harmful. Limit fluids to one or two glasses in an adult.
- Charcoal has no place in acid management.
- Some authors suggest the use of lavage within 1 hour of ingestion.

SKIN:

+ Skin lesions require copious saline irrigation. Treat chemical burns as thermal burns with non-adherent gauze and wrapping.

Deep second-degree burns may benefit from topical silver sulfadiazine.

EYE:

Eye injuries require retraction of the eyelids to ensure thorough irrigation of the conjuctival cul-de-sacs. Irrigation should last at least 20-30 minutes. DO NOT use neutralising agents or any other additives. Several litres of saline are required.

Cycloplegic drops, (1% cyclopentolate for short-term use or 5% homatropine for longer term use) antibiotic drops, vasoconstrictive agents or artificial tears may be indicated dependent on the severity of the injury.

• Steroid eye drops should only be administered with the approval of a consulting ophthalmologist).

[Ellenhorn and Barceloux: Medical Toxicology]

- · Gastric acids solubilise lead and its salts and lead absorption occurs in the small bowel.
- Particles of less than 1 um diameter are substantially absorbed by the alveoli following inhalation.
- Lead is distributed to the red blood cells and has a half-life of 35 days. It is subsequently redistributed to soft tissue & bone-stores or eliminated. The kidney accounts for 75% of daily lead loss; integumentary and alimentary losses account for the remainder.
- Neurasthenic symptoms are the most common symptoms of intoxication. Lead toxicity produces a classic motor neuropathy. Acute encephalopathy appears infrequently in adults. Diazepam is the best drug for seizures.
- Whole-blood lead is the best measure of recent exposure; free erythrocyte protoporphyrin (FEP) provides the best screening for chronic exposure. Obvious clinical symptoms occur in adults when whole-blood lead exceeds 80 ug/dL.

British Anti-Lewisite is an effective antidote and enhances faecal and urinary excretion of lead. The onset of action of BAL is about 30 minutes and most of the chelated metal complex is excreted in 4-6 hours, primarily in the bile. Adverse reaction appears in up to 50% of patients given BAL in doses exceeding 5 mg/kg. CaNa2EDTA has also been used alone or in concert with BAL as an antidote. D-penicillamine is the usual oral agent for mobilisation of bone lead; its use in the treatment of lead poisoning remains investigational. 2,3-dimercapto-1-propanesulfonic acid (DMPS) and dimercaptosuccinic acid (DMSA) are water soluble analogues of BAL and their effectiveness is undergoing review. As a rule, stop BAL if lead decreases below 50 ug/dL; stop CaNa2EDTA if blood lead decreases below 40 ug/dL or urinary lead drops below 2 mg/24hrs.

[Ellenhorn & Barceloux: Medical Toxicology]

BIOLOGICAL EXPOSURE INDEX - BEI

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

Determinant 1. Lead in blood	Index 30 ua/100 ml	Sampling Time Not Critical	Comments
2. Lead in urine	150 ug/gm creatinine	Not Critical	В
3. Zinc protoporphyrin in blood	250 ug/100 ml erythrocytes OR 100 ug/100 ml blood	After 1 month exposure	В

B: Background levels occur in specimens collected from subjects NOT exposed.

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

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Fire Incompatibility	None known.				
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 suppected to be hot. If safe to do so, remove containers from path of fire. Equipment should be thoroughly decontaminated after use. 				

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 containers. May emit corrosive, poisonous fumes. May emit acrid smoke. Decomposition may produce toxic fumes of: sulfur oxides (SOx)
HAZCHEM	2R

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	 Clean up all spills immediately. Secure load if safe to do so. Bundle/collect recoverable product. Collect remaining material in containers with covers for disposal.
Major Spills	 Clean up all spills immediately. Wear protective clothing, safety glasses, dust mask, gloves. Secure load if safe to do so. Bundle/collect recoverable product. Use dry clean up procedures and avoid generating dust. Vacuum up (consider explosion-proof machines designed to be grounded during storage and use). Water may be used to prevent dusting. Collect remaining material in containers with covers for disposal. Flush spill area with water.

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

SECTION 7 HANDLING AND STORAGE

Precautions for safe handling

Safe handling	 Avoid all personal contact, including inhalation. Wear protective clothing when risk of exposure occurs. Use in a well-ventilated area. Avoid contact with moisture. Avoid contact with noisture. Avoid contact with noisture. When handling, DO NOT eat, drink or smoke. Keep containers securely sealed when not in use. Avoid physical damage to containers. Always wash hands with scap and water after handling. Work clothes should be laundered separately. Launder contaminated clothing before re-use. Use good occupational work practice. Observe manufacturer's storage and handling recommendations contained within this SDS. Atmosphere should be regularly checked against established exposure standards to ensure safe working conditions are maintained. Avoid shorting circuit or sparks near battery. Avoid prolonged over-charging. Use only approved charging methods. Do not charge in gas tight containers.
Other information	 Store away from incompatible materials. Keep dry. Store under cover. Protect containers against physical damage. Observe manufacturer's storage and handling recommendations contained within this SDS.

Conditions for safe storage, including any incompatibilities

Suitable container	 Check regularly for spills and leaks Packaging as recommended by manufacturer.
Storage incompatibility	 Reacts with mild steel, galvanised steel / zinc producing hydrogen gas which may form an explosive mixture with air. Avoid any contamination of this material as it is very reactive and any contamination is potentially hazardous Avoid storage with reducing agents. Store away from reactive materials.

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	lead	Lead, inorganic dusts & fumes (as Pb)	0.05 mg/m3	Not Available	Not Available	Not Available

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Australia Exposure Standards	lead dioxide	Lead, inorganic dusts & fumes (as Pb)	0.05 mg/m3	Not Available	Not Available	Not Available			
Australia Exposure Standards	sulfuric acid	Sulphuric acid	1 mg/m3	3 mg/m3	Not Available	Not Available			
Australia Exposure Standards	tin	Tin, metal	2 mg/m3	Not Available	Not Available	Not Available	Not Available		
Australia Exposure Standards	arsenic	Arsenic & soluble compounds (as As)	0.05 mg/m3	Not Available	Not Available	(g) Some compounds in these groups are classified as carcinogenia or as sensitisers. Check individual classification details on the safety data sheet for information on classification.			
EMERGENCY LIMITS									
Ingredient	Material nam	e	TEEL-1			TEEL-2	TEEL-3		
lead	Lead		0.15 mg/	'm3		120 mg/m3	700 mg/m3		
lead dioxide	Lead dioxide		0.17 mg/	'm3		140 mg/m3	810 mg/m3		
sulfuric acid	Sulfuric acid		Not Avai	Not Available		Not Available	Not Available		
tin	Tin		6 mg/m3			67 mg/m3 400 mg/m3			

Ingredient	Original IDLH	Revised IDLH
lead	Not Available	Not Available
lead dioxide	100 mg/m3	Not Available
sulfuric acid	15 mg/m3	Not Available
calcium	Not Available	Not Available
tin	Not Available	Not Available
arsenic	Not Available	Not Available

MATERIAL DATA

Exposure controls

Appropriate engineering controls	Provide adequate ventilation in warehouse or closed storage areas.		
Personal protection			
Eye and face protection	 None under normal operating conditions. OTHERWISE: Safety glasses with unperforated side shields may be used where continuous eye protection is desirable, as in laboratories; spectacles are not sufficient where complete eye protection is needed such as when handling bulk-quantities, where there is a danger of splashing, or if the material may be under pressure. Chemical goggles.whenever there is a danger of the material coming in contact with the eyes; goggles must be properly fitted. Full face shield (20 cm, 8 in minimum) may be required for supplementary but never for primary protection of eyes; these afford face protection. Alternatively a gas mask may replace splash goggles and face shields. Contact lenses may pose a special hazard; soft contact lenses may absorb and concentrate irritants. 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 injury experience. Medical and first-aid personnel 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. Lens should be removed at the first signs of eye redness or irritation - lens should be removed in a clean environment only after workers have washed hands thoroughly. [CDC NIOSH Current Intelligence Bulletin 59], [AS/NZS 1336 or national equivalent] 		
Skin protection	See Hand protection below		
Hands/feet protection	None under normal operating conditions. OTHERWISE: • Elbow length PVC gloves Wear general protective gloves, eg. light weight rubber gloves.		
Body protection	See Other protection below		
Other protection	None under normal operating conditions. OTHERWISE: • Overalls. • PVC Apron. • PVC protective suit may be required if exposure severe. • Eyewash unit. • Ensure there is ready access to a safety shower.		

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:

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Respiratory protection

Type E-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.

Material	CPI	Required M
NATURAL RUBBER	А	Protection

quired Minimum Half-Face Full-Face Powered Air otection Factor Respirator Respirator Respirator

NATURAL+NEOPRENE	A
NEOPRENE	A
NEOPRENE/NATURAL	A
NITRILE	А
PE	A
PVC	A
SARANEX-23	А

* 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.

up to 10 x ES	E-AUS P2	-	E-PAPR-AUS / Class 1 P2
up to 50 x ES	-	E-AUS / Class 1 P2	-
up to 100 x ES	-	E-2 P2	E-PAPR-2 P2 ^

^ - Full-face

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

- Respirators may be necessary when engineering and administrative controls do not adequately prevent exposures.
- The decision to use respiratory protection should be based on professional judgment that takes into account toxicity information, exposure measurement data, and frequency and likelihood of the worker's exposure ensure users are not subject to high thermal loads which may result in heat stress or distress due to personal protective equipment (powered, positive flow, full face apparatus may be an option).
- Published occupational exposure limits, where they exist, will assist in determining the adequacy of the selected respiratory protection. These may be government mandated or vendor recommended.
- Certified respirators will be useful for protecting workers from inhalation of particulates when properly selected and fit tested as part of a complete respiratory protection program.
- Use approved positive flow mask if significant quantities of dust becomes airborne.
- Try to avoid creating dust conditions.

SECTION 9 PHYSICAL AND CHEMICAL PROPERTIES

Information on basic physical and chemical properties

Appearance Solid shape article consisting of an opaque plastic case with two lead terminals; Electrolyte is clear corrosive liquid absorbed in glass mat material with little pungent odor; miscible with water.

Physical state	Manufactured	Relative density (Water = 1)	1.170-1.335 (electrolyte)
Odour	Not Available	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	357 (polypropylene)
pH (as supplied)	Not Available	Decomposition temperature	Not Available
Melting point / freezing point (°C)	Not Available	Viscosity (cSt)	Not Available
Initial boiling point and boiling range (°C)	110-112 (electrolyte)	Molecular weight (g/mol)	Not Applicable
Flash point (°C)	Not Available	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Not Available	Oxidising properties	Not Available
Upper Explosive Limit (%)	74.2 (H2 gas)	Surface Tension (dyn/cm or mN/m)	Not Applicable
Lower Explosive Limit (%)	4.1 (H2 gas)	Volatile Component (%vol)	Not Available
Vapour pressure (kPa)	3 (electrolyte)	Gas group	Not Available
Solubility in water	Miscible	pH as a solution (1%)	Not Available
Vapour density (Air = 1)	3.4 (electrolyte)	VOC g/L	Not Available

SECTION 10 STABILITY AND REACTIVITY

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

Inhaled

Information on toxicological effects

Evidence shows, or practical experience predicts, that the material produces irritation of the respiratory system, in a substantial number of individuals, following inhalation. In contrast to most organs, the lung is able to respond to a chemical insult by first removing or neutralising the irritant and then repairing the damage. The repair process, which initially evolved to protect mammalian lungs from foreign matter and antigens, may however, produce further lung damage resulting in the impairment of gas exchange, the primary function of the lungs. Respiratory tract irritation often results in an inflammatory response involving the recruitment and activation of many cell types, mainly derived from the vascular system.

	Acidic corrosives produce respiratory tract irritation with coughing, choking and headache, nausea and weakness. In more severe exposures, pulmonary oeder Symptoms of pulmonary oedema include a tightness in the chest, dyspnoea, fro rapid pulse and moist rates. Death, due to anoxia, may occur several hours after	ma may be evident either immediately or after a latent period of 5-72 hours. thy sputum and cyanosis. Examination may reveal hypotension, a weak and
Ingestion	Accidental ingestion of the material may be harmful; animal experiments indica damage to the health of the individual. Ingestion of acidic corrosives may produce circumoral burns with a distinct dis Immediate pain and difficulties in swallowing and speaking may also be evider asphyxia. Nausea, vomiting, diarrhoea and a pronounced thirst may occur. Mo and large shreds of mucosa. Shock, with marked hypotension, weak and rapid exposure. Circulatory collapse may, if left untreated, result in renal failure. Sev fever and abdominal rigidity. Stricture of the oesophageal, gastric and pyloric se Death may be rapid and often results from asphyxia, circulatory collapse or asp severe nephritis or pneumonia. Coma and convulsions may be terminal.	colouration of the mucous membranes of the mouth, throat and oesophagus. Int. Oedema of the epiglottis may produce respiratory distress and possibly, are severe exposures may produce a vomitus containing fresh or dark blood pulse, shallow respiration and clammy skin may be symptomatic of the vere cases may show gastric and oesophageal perforation with peritonitis, sphincter may occur as within several weeks or may be delayed for years.
Skin Contact	Skin contact with acidic corrosives may result in pain and burns; these may be tissue. Open cuts, abraded or irritated skin should not be exposed to this material Entry into the blood-stream through, for example, cuts, abrasions, puncture wo the skin prior to the use of the material and ensure that any external damage is	ounds or lesions, may produce systemic injury with harmful effects. Examine
Eye	When applied to the eye(s) of animals, the material produces severe ocular le Direct eye contact with acid corrosives may produce pain, lachrymation, photo completely. Severe burns produce long-lasting and possible irreversible damay the initial contact. The comea may ultimately become deeply vascularised and	phobia and burns. Mild burns of the epithelia generally recover rapidly and ge. The appearance of the burn may not be apparent for several weeks after
Chronic	produce cerebrospinal pressure, brain damage with stupor leading to coma all include anorexia and loss of weight, constipation, apathy or irritability, occasio Advanced poisonings are characterised by intermittent vomiting, irritability, ne Severe poisonings may produce persistent vomiting, ataxia, stupor or lethargy tension, papilloedema, cranial nerve paralysis, delirium, convulsions and com and marked muscular contractions that distort the spine, limbs, hips and some Industrial exposure has been associated with irreversible kidney damage. Lead is a cumulative poison with adverse effects in pregnancy [NIOSHTIC] Lead salts have been reported to cross the placenta and induce embryo- and f deformities) in certain animal species. Organometallic lead may not produce the foetal development and postnatal mental development have also been recorde behavioural disorders and death during the first year of childhood. Paternal effe	nia may ensue. Gastrointestinal disturbances may also occur. Chronic number of interrelated factors. These include physicochemical characteristics, no to the lung); water solubility (more soluble agents are more likely to be particle size of aerosols involved in occupational exposures to acids, it is id mists containing particles with a diameter of up to a few micrometers will s epithelia, they cause dental erosion, and they produce acute effects in the be at particular risk for pulmonary effects. health effects involving organs or biochemical systems. rs involving difficult breathing and related systemic problems. inhalation, in contact with skin and if swallowed. may have toxicological significance) is likely to be caused by repeated or which produces severe lesions. Such damage may become apparent sub-acute (28 day) or chronic (two-year) toxicity tests. exposure to the material and subsequent developmental toxic effects in the the basis that results in animal studies provide sufficient evidence to cause a e of impaired fertility occurring at around the same dose levels as other toxic c effects. docrine organs and the immune system and the digestive system. The ad, neuromuscular dysfunction, possible paralysis and encephalopathy (brain muscle pain, weakness of the extensor muscles (frequently the hand and vomiting, blue line on the gums, insomnia and metallic taste. High body levels nd, in some cases, death. Early symptoms of lead poisoning ("plumbism") nal vomiting, fatigue, headache, weakness, and a metallic taste in the mouth. ervousness, myalgia of the arms and legs (often with wrist and foot drop). /, visual disturbances progressing to optic neuritis and atrophy, hyper- na. Neurological effects include mental retardation, seizures, cerebral palsy times the cranial inervated muscles (dystonia musculorum deformans).
Scooter Xtras XT100G, XT75G, XT55G, XT50G, XT36G, XT20G	TOXICITY Not Available	IRRITATION Not Available
	τοχιςιτγ	IRRITATION

X155G, X150G, X136G, X120G	Not Available	Not Available
	TOXICITY	IRRITATION
	dermal (rat) LD50: >2000 mg/kg ^[1]	Not Available
lead	Inhalation (rat) LC50: >5.05 mg/l4 h ^[1]	1 1 1 1
	Oral (rat) LD50: >2000 mg/kg ^[1]	
	TOXICITY	IRRITATION
lead dioxide	Not Available	Not Available
	TOXICITY	IRRITATION
sulfuric acid	Inhalation (guinea pig) LC50: 0.036 mg/l/8H ^[2]	Eye (rabbit): 1.38 mg SEVERE
		1

	Oral (rat) LD50: 2140 mg/kg ^[2]	Eye (rabbit): 5 mg/30sec SEVERE
	ТОХІСІТҮ	IRRITATION
calcium	dermal (rat) LD50: >2000 mg/kg ^[1]	Eye: no adverse effect observed (not irritating) ^[1]
	Oral (rat) LD50: >2000 mg/kg ^[1]	Skin: no adverse effect observed (not irritating) ^[1]
	TOXICITY	IRRITATION
tin	dermal (rat) LD50: >2000 mg/kg ^[1]	Eye: no adverse effect observed (not irritating) ^[1]
	Oral (rat) LD50: >2000 mg/kg ^[1]	Skin: no adverse effect observed (not irritating) ^[1]
	TOXICITY	IRRITATION
arsenic	Oral (rat) LD50: 763 mg/kg ^[1]	Eye: adverse effect observed (irreversible damage) ^[1]
		Skin: adverse effect observed (irritating) ^[1]
Legend:	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	

LEAD	WARNING: Lead is a cumulative poison and has the pote	ential to cause abortion and intellectual in	npairment to unborn children of pregnant workers.
	WARNING: For inhalation exposure ONLY: This substar	nce has been classified by the IARC as (Group 1: CARCINOGENIC TO HUMANS
SULFURIC ACID	Occupational exposures to strong inorganic acid mists o	f sulfuric acid:	
CALCIUM	The solid may react violently on contact with wet skin tissue, i.e. eyes, mouth, causing chemical and thermal burns. The acute effects include burns, ulceration, or tissue death, severe eye damage (corneal burns or opacification), and probable blindness. Inhalation of dust or fumes (especially from a fire involving calcium) will cause shortness of breath, nausea, headache, nose and respiratory tract irritation and in extreme, pneumonitis		
	Arsenic compounds are classified by the European Unior IARC classify arsenic in drinking water as a confirmed hu The main inorganic forms of arsenic relevant for human e (also called arsenite, As(III), or As+3). These inorganic sp tissues to form tri, and contavalent methylated methylated	uman carcinogen (IARC 1). exposures are pentavalent arsenic (also becies undergoes a series of reduction a	called arsenate, As(V), or As+5) and trivalent arsenic ind oxidative/methylation steps in human liver and other
tissues to form tri- and pentavalent methylated metabolite dimethylarsinate [DMA(V)]. Some mammalian species als ARSENIC The distinction between inorganic and organic forms is in the body and generally considered less toxic, with a relati trivalent metabolites, MA(III) and DMA(III), are significant biomonitoring or environmental occurrence data are report understanding the relevant sources of arsenic is essential exposure. WARNING: This substance has been classified by the IA Tumorigenic - Carcinogenic by RTECS criteria.		so produce trimethylated metabolites, trim portant because it is generally accepted ive rank order of $As(III) > As(V) >> MA(V)$ ly more toxic than their pentavalent cour rted as total arsenic and do not distinguis I to evaluate potential arsenic related he	nethylarsine oxide d that the organic species are excreted more quickly fro /), DMA(V) >> arsenobetaine. However, the methylated hterpart and either As(III) or As(V). In many cases, sh between the different species. In those situations, alth effects, especially those related to inorganic arsenia
SULFURIC ACID & CALCIUM	Asthma-like symptoms may continue for months or even y reactive airways dysfunction syndrome (RADS) which ca diagnosis of RADS include the absence of preceding resp within minutes to hours of a documented exposure to the bronchial hyperreactivity on methacholine challenge testin in the criteria for diagnosis of RADS. RADS (or asthma) of and duration of exposure to the irritating substance. In concentrations of irritating substance (often particulate in dyspnea, cough and mucus production.	n occur following exposure to high levels biratory disease, in a non-atopic individu- irritant. A reversible airflow pattern, on sp ig and the lack of minimal lymphocytic inf following an irritating inhalation is an inf dustrial bronchitis, on the other hand, is a	s of highly irritating compound. Key criteria for the al, with abrupt onset of persistent asthma-like symptoms birometry, with the presence of moderate to severe flammation, without eosinophilia, have also been include requent disorder with rates related to the concentration a disorder that occurs as result of exposure due to high
CALCIUM & TIN	No significant acute toxicological data identified in literati	ure search.	
Acute Toxicity	×	Carcinogenicity	✓
Skin Irritation/Corrosion	✓	Reproductivity	✓
Serious Eye Damage/Irritation	✓	STOT - Single Exposure	×
Respiratory or Skin	×	STOT - Repeated Exposure	*
sensitisation			

SECTION 12 ECOLOGICAL INFORMATION

Toxicity

Secolar Viras VT400C VT75C	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
Scooter Xtras XT100G, XT75G, XT55G, XT50G, XT36G, XT20G	Not Available	Not Available	Not Available	Not Available	Not Available

	ENDPOINT	TEST DURATION (HR)	er	PECIES	VAL	IE	SOURCE
	LC50	96	Fi		1	-0.06756mg/L	2
	EC50	1	1		1		
lead		48		rustacea	0.029	-	2
	EC50	72		gae or other aquatic plants)5mg/L	2
	BCFD	8	Fi	-	4.324	•	4
	NOEC	672	Fi	sh	0.000	103mg/L	4
	ENDPOINT	TEST DURATION (HR)		SPECIES		VALUE	SOURCE
lead dioxide	NOEC	264		Algae or other aquatic plants		0.0091mg/L	2
	ENDPOINT	TEST DURATION (HR)		SPECIES		VALUE	SOURCE
	LC50	96	1	Fish		=8mg/L	1
sulfuric acid	EC50	48		Crustacea		=42.5mg/L	1
	EC50	72 Algae or other aquatic plants			>100mg/L	2	
	NOEC	Not Available		Crustacea		0.15mg/L	2
	ENDPOINT	TEST DURATION (HR)		SPECIES		VALUE	SOURCE
	EC50	48		Crustacea		49.1mg/L	2
calcium	EC100	48		Crustacea		75mg/L	2
	NOEC	336		Crustacea		32mg/L	2
	ENDPOINT	TEST DURATION (HR)	S	PECIES	VA	LUE	SOURCE
	LC50	96	F	Fish	>0	.0124mg/L	2
tin	EC50	48	C	Crustacea	0.0	0018mg/L	5
	EC50	72	Α	Algae or other aquatic plants 0.009-0.846		09-0.846mg/L	2
	NOEC	72	A	Algae or other aquatic plants 0.001-mg/L		01-mg/L	2
	ENDPOINT	TEST DURATION (HR)		SPECIES		VALUE	SOURCE
	LC50	96		Fish		3.38mg/L	2
arsenic	EC50	48		Crustacea		0.015mg/L	2
	EC10	48	 	Crustacea		0.006mg/L	2
-	NOEC	336		Algae or other aquatic plants		0.01mg/L	2

Legend:

Extracted from 1. IUCLID Toxicity Data 2. Europe ECHA Registered Substances - Ecotoxicological Information - Aquatic Toxicity 3. EPIWIN Suite V3.12 (QSAR) - Aquatic Toxicity Data (Estimated) 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

Very toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment.

Do NOT allow product to come in contact with surface waters or to intertidal areas below the mean high water mark. Do not contaminate water when cleaning equipment or disposing of equipment wash-waters.

Wastes resulting from use of the product must be disposed of on site or at approved waste sites.

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

Decontaminate empty containers. Observe all label safeguards until containers are cleaned and destroyed.

SECTION 14 TRANSPORT INFORMATION

Labels Required	
	B B
Marine Pollutant	
HAZCHEM	2R

Land transport (ADG)

UN number	2800		
UN proper shipping name	BATTERIES, WET, NON-SPILLABLE, electric storage		
Transport hazard class(es)	Class 8 Subrisk Not Applicable		
Packing group	Not Applicable		
Environmental hazard	Environmentally hazardous		
Special precautions for user	Special provisions 238 Limited quantity 1 L		

Air transport (ICAO-IATA / DGR)

UN number	2800			
UN proper shipping name	Batteries, wet, non-spillable electric storage			
Transport hazard class(es)	ICAO/IATA Class 8 ICAO / IATA Subrisk Not Applicable ERG Code 8L			
Packing group	Not Applicable			
Environmental hazard	Environmentally hazardous			
Special precautions for user	Special provisions Cargo Only Packing Instructions Cargo Only Maximum Qty / Pack Passenger and Cargo Packing Instructions Passenger and Cargo Maximum Qty / Pack Passenger and Cargo Limited Quantity Packing Instructions Passenger and Cargo Limited Maximum Qty / Pack		A48 A67 A164 A183 872 No Limit 872 No Limit Forbidden Forbidden	

Sea transport (IMDG-Code / GGVSee)

• •	,		
UN number	2800		
UN proper shipping name	BATTERIES, WET, NON-SPILLABLE electric storage		
Transport hazard class(es)	IMDG Class 8 IMDG Subrisk Not Applicable		
Packing group	Not Applicable		
Environmental hazard	Marine Pollutant		
Special precautions for user	EMS NumberF-A , S-BSpecial provisions238Limited Quantities1 L		

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

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

LEAD(7439-92-1) IS FOUND ON THE FOLLOWING REGULATORY LISTS Australia Exposure Standards Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Index Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule Australia Inventory of Chemical Substances (AICS) Δ IMO IBC Code Chapter 17: Summary of minimum requirements Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Appendix B (Part 3) International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs LEAD DIOXIDE(1309-60-0) IS FOUND ON THE FOLLOWING REGULATORY LISTS Australia Dangerous Goods Code (ADG Code) - Dangerous Goods List Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule 10 / Appendix C Australia Dangerous Goods Code (ADG Code) - List of Emergency Action Codes Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule Australia Exposure Standards Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule Australia Inventory of Chemical Substances (AICS) 6 Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Appendix International Agency for Research on Cancer (IARC) - Agents Classified by the IARC E (Part 2) Monographs Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Appendix International Air Transport Association (IATA) Dangerous Goods Regulations F (Part 3) International Maritime Dangerous Goods Requirements (IMDG Code) Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Index United Nations Recommendations on the Transport of Dangerous Goods Model Regulations SULFURIC ACID(7664-93-9) IS FOUND ON THE FOLLOWING REGULATORY LISTS Australia Dangerous Goods Code (ADG Code) - Dangerous Goods List **GESAMP/EHS** Composite List - GESAMP Hazard Profiles Australia Dangerous Goods Code (ADG Code) - List of Emergency Action Codes IMO IBC Code Chapter 17: Summary of minimum requirements IMO MARPOL (Annex II) - List of Noxious Liquid Substances Carried in Bulk Australia Exposure Standards Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs Australia Inventory of Chemical Substances (AICS) International Air Transport Association (IATA) Dangerous Goods Regulations Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Appendix International Air Transport Association (IATA) Dangerous Goods Regulations - Prohibited List E (Part 2) Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Appendix Passenger and Cargo Aircraft F (Part 3) International Maritime Dangerous Goods Requirements (IMDG Code) Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule United Nations Recommendations on the Transport of Dangerous Goods Model Regulations 6 CALCIUM(7440-70-2) IS FOUND ON THE FOLLOWING REGULATORY LISTS Australia Dangerous Goods Code (ADG Code) - Dangerous Goods List International Air Transport Association (IATA) Dangerous Goods Regulations Australia Dangerous Goods Code (ADG Code) - List of Emergency Action Codes International Maritime Dangerous Goods Requirements (IMDG Code) Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals United Nations Recommendations on the Transport of Dangerous Goods Model Regulation Australia Inventory of Chemical Substances (AICS) TIN(7440-31-5) IS FOUND ON THE FOLLOWING REGULATORY LISTS Australia Inventory of Chemical Substances (AICS) Australia Exposure Standards ARSENIC(7440-38-2) IS FOUND ON THE FOLLOWING REGULATORY LISTS Australia Dangerous Goods Code (ADG Code) - Dangerous Goods List Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule Australia Dangerous Goods Code (ADG Code) - List of Emergency Action Codes Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule Australia Exposure Standards Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule Australia Inventory of Chemical Substances (AICS) Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Appendix International Agency for Research on Cancer (IARC) - Agents Classified by the IARC G Monographs Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Appendix International Air Transport Association (IATA) Dangerous Goods Regulations J (Part 2) International Maritime Dangerous Goods Requirements (IMDG Code)

Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Index

United Nations Recommendations on the Transport of Dangerous Goods Model Regulation

National Inventory Status

National Inventory	Status
Australia - AICS	Yes
Canada - DSL	Yes
Canada - NDSL	No (lead; calcium; lead dioxide; arsenic; sulfuric acid; tin)
China - IECSC	Yes
Europe - EINEC / ELINCS / NLP	Yes
Japan - ENCS	No (lead; calcium; arsenic; tin)
Korea - KECI	Yes
New Zealand - NZIoC	Yes
Philippines - PICCS	Yes
USA - TSCA	Yes
Taiwan - TCSI	Yes
Mexico - INSQ	Yes
Vietnam - NCI	Yes
Russia - ARIPS	Yes

Thailand - TECI	No (lead; lead dioxide)
Legend:	Yes = All CAS declared ingredients are on the inventory No = One or more of the CAS listed ingredients are not on the inventory and are not exempt from listing(see specific ingredients in brackets)

SECTION 16 OTHER INFORMATION

Revision Date	24/01/2019
Initial Date	05/09/2013

SDS Version Summary

Version	Issue Date	Sections Updated
3.1.1.1	07/08/2015	Use
4.1.1.1	24/01/2019	Storage (suitable container)

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.

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 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 This document is copyright.

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