Brenntag Canada Inc.



MATERIAL SAFETY DATA SHEET

POTASSIUM AMYL XANTHATE, SOLID

00060600

1. CHEMICAL PRODUCT AND COMPANY IDENTIFICATION

Brenntag Canada Inc. 43 Jutland Rd. Toronto, ON M8Z 2G6 (416) 259-8231

Website: http://www.brenntag.ca

Index: HCI0065/09B Effective Date: 2009 June 17 Date of Revision: 2009 June 17

WHMIS#:

EMERGENCY TELEPHONE NUMBERS (FOR EMERGENCIES INVOLVING CHEMICAL SPILLS OR RELEASE)

Toronto, ON (416) 226-6117 Montreal, QC (514) 861-1211 Winnipeg, MB (204) 943-8827 Edmonton, AB (780) 424-1754 Calgary, AB (403) 263-8660 Vancouver, BC (604) 685-5036

PRODUCT IDENTIFICATION

Product Name: Potassium Amyl Xanthate, Solid.

Chemical Name: Dithiocarbonic Acid, Amyl Ester, Potassium Salt.

Synonyms: Potassium Amyl Xanthate; KAX 51; Potassium Pentyl Xanthate; Potassium Pentyl Xanthogenate.

Chemical Family: Salts of carbonic acid dithio esters.

Molecular Formula: C6H11OS2. K.
Product Use: Flotation agent.

WHMIS Classification / Symbol:

B-6: Reactive Flammable Material D-1B: Toxic (acute effects)
D-2B: Toxic (skin and eye irritant)







READ THE ENTIRE MSDS FOR THE COMPLETE HAZARD EVALUATION OF THIS PRODUCT.

2. COMPOSITION, INFORMATION ON INGREDIENTS (Not Intended As Specifications)

Ingredient	CAS#	ACGIH TLV	% Concentration
Potassium Amyl Xanthate	2720-73-2		60 - 100
Potassium Hydroxide	1310-58-3	_	1 - 5
Isoamyl alcohol	123-51-3	100 ppm	1 - 5

Decomposition Product: Carbon disulfide 75-15-0 10 ppm (Skin)

Skin Notation: Contact with skin, eyes and mucous membranes can contribute to the overall exposure and may invalidate the TLV. Consider measures to prevent absorption by these routes.

3. HAZARDS IDENTIFICATION

EMERGENCY OVERVIEW:

May be fatal if swallowed. Harmful if inhaled. Causes skin and eye irritation. Dust is irritating to respiratory tract. See "Other Health Effects" Section. Heating of solid xanthate or aging or heating of solutions will cause formation of Carbon Bisulfide. Upon exposure of solid xanthates to moisture and/or heat, decomposition results and spontaneous combustion can occur. Contact of solid xanthate with moist air has resulted in ignition. (4) Emits a flammable gas upon contact with water or water vapour. Can decompose at high temperatures forming toxic gases. Powdered material may form explosive dustair mixtures. Contents may develop pressure on prolonged exposure to heat.

Potassium Amyl Xanthate, Solid WHMIS Number: 00060600

Date of Revision:

Page 2 of 8

POTENTIAL HEALTH EFFECTS

Inhalation: Excessive contact with powder may cause drying of mucous membranes of nose and throat due to

absorption of moisture and oils. Product may cause severe irritation of the nose, throat and respiratory

Brenntag Canada Inc.

2009 June 17

tract. Repeated and/or prolonged exposures may cause productive cough, running nose,

bronchopneumonia, pulmonary oedema (fluid build-up in lungs), and reduction of pulmonary function. Irritation of mucous membranes and respiratory tract is possible following exposure to the decomposition

product. (3) See "Other Health Effects" Section.

Skin Contact: Brief contact with the dust causes irritation. Greater exposure causes severe burns. In the presence of

moisture (perspiration, humidity, tears), the dust dissolves to form a corrosive solution which may cause burns. (3) Potassium Amyl Xanthate may cause symptoms of skin irritation such as reddening, swelling,

rash, scaling, or blistering. May cause defatting, drying and cracking of the skin.

Skin Absorption: May be absorbed through intact skin. See Section 11, "Other Studies Relevant to Material".

Eye Contact: This product may cause irritation, redness and possible damage due to abrasiveness. Brief contact with

the dust causes irritation. Greater exposure causes severe burns. In the presence of moisture (perspiration, humidity, tears), the dust dissolves to form a corrosive solution which may cause burns.

(3) Irritation of the eyes is possible following exposure to the decomposition product. (3)

Ingestion: Ingestion is not a likely route of exposure. This product causes irritation, a burning sensation of the

mouth and throat and abdominal pain.

Other Health Effects: Effects (irritancy) on the skin and eyes may be delayed, and damage may occur without the sensation or onset of pain. Strict adherence to first aid measures following any exposure is essential.

May cause cardiovascular effects, liver damage, peripheral nervous system (PNS) effects or central nervous system (CNS) depression. CNS depression is characterized by headache, dizziness, drowsiness, nausea, vomiting and incoordination. Severe overexposures may lead to coma and possible death due to respiratory failure. Peripheral Neuropathy is a progressive disorder of the nervous system characterized by sensory and motor abnormalities, muscle spasms, weakness and pain in the arms and legs, numbness and tingling of the fingers and toes and paralysis. Liver damage is characterized by the loss of appetite, jaundice (yellowish skin colour), and occasional pain in the upper left-hand side of the abdomen.

Potassium Amyl Xanthate: Symptoms of potassium poisoning may occur. These include slow heartbeat, accelerated breathing, muscle weakness and, in severe cases, paralysis.

Vapours of the decomposition products of Xanthates (Carbon Bisulphide) can cause severe disturbances of mood and behaviour, including excitation, anger and violent dreams. High concentrations of vapours can cause death. (4)

Carbon Bisulphide: Contact with moisture in the body by inhalation may yield sodium hydroxide (corrosive) and 2-mercaptobenzothiazole, an irritant. (4) Contact with acids will liberate carbon disulphide. (3) Exposure to carbon disulphide (500 to 1000 ppm) may cause severe mood and personality disturbances, including excitability, confusion, irritability, uncontrollable anger, bizarre dreams, insomnia, psychosis and suicide. Exposure to carbon disulphide at 4800 ppm for thirty minutes results in coma and may be fatal. Carbon disulphide is readily absorbed through intact skin. Chronic exposure to carbon disulphide produces central and peripheral nervous system, cardiovascular, gastrointestinal, kidney, endocrine and eye disorders. (4)

Potassium Hydroxide: Exposure to very low doses, even for a short period of time, has produced extensive damage to the esophagus, stomach and intestine extending into surrounding tissues, as well as hyperexcitability followed by apathy and weakness. In some cases, death has resulted from hemorrhage, adhesions or perforation. Following esophageal damage, strictures have frequently developed in surviving animals. (4)

4. FIRST AID MEASURES

FIRST AID PROCEDURES

Skin Contact:

Inhalation: Move victim to fresh air. Give artificial respiration ONLY if breathing has stopped. Give cardiopulmonary

resuscitation (CPR) if there is no breathing AND no pulse. Obtain medical advice IMMEDIATELY.

Prompt removal of the material from the skin is essential. Remove all contaminated clothing and

immediately wash the exposed areas with copious amounts of soap and water for a minimum of 30 minutes or up to 60 minutes for critical body areas. Immerse the exposed part immediately in ice water to relieve pain and to prevent swelling and blistering. Place cold packs, ice or wet cloths on the burned

area if immersion is not possible. Cover the exposed part with a clean, preferably sterile, lint-free dressing. Obtain medical attention IMMEDIATELY and monitor breathing and treat for shock for severe

exposure.

Eye Contact: Immediately flush eyes with running water for a minimum of 20 minutes. Hold eyelids open during

flushing. If irritation persists, repeat flushing. Obtain medical attention IMMEDIATELY.

WHMIS Number: 00060600 Date of Revision: 2009 June 17

Page 3 of 8

Ingestion: Do not attempt to give anything by mouth to an unconscious person. If victim is alert and not convulsing,

rinse mouth out and give 1/2 to 1 glass of water to dilute material. IMMEDIATELY contact local Poison Control Centre. Vomiting should only be induced under the direction of a physician or a poison control centre. If spontaneous vomiting occurs, have victim lean forward with head down to avoid breathing in of vomitus, rinse mouth and administer more water. IMMEDIATELY transport victim to an emergency

facility.

Note to Physicians: Treat symptomatically.

Medical conditions that may be aggravated by exposure to this product include neurological and cardiovascular disorders, diseases of the skin, eyes or respiratory tract, preexisting liver and kidney

disorders.

5. FIRE-FIGHTING MEASURES

		Flammability Limits in Air (%):	
Flashpoint (°C)	Autolgnition Temperature (°C)	LEL	UEL
-30. (Carbon Disulphide)	90.1 (Carbon Disulphide)	1.25. (Carbon Disulphide)	50. (Carbon Disulphide)
Flammability Class (WHMIS):	B-6: Reactive Flammable Material		
Hazardous Combustion Products:	Thermal decomposition products are carbonyl sulphide, Amyl Alcohols, ox		
Unusual Fire or Explosion Hazards:	This product may be capable of form of dust to reduce explosion potential. slippery. Heating of solid xanthate or Bisulfide. Upon exposure of solid xa spontaneous combustion can occur. Vapours from this product are heaving heaters, electric motors) some distart causing an explosion and fire. Enforce	Spilled material may aging or heating of so nthates to moisture ar Contact of solid xantler than air, and may "the away, and then "flate".	cause floors and contact surfaces to plutions will cause formation of Carbond/or heat, decomposition results and nate with moist air has resulted in igrand are solved in igrand to a source of ignition (eg. pilotash back" to the point of product discourse.
Sensitivity to Mechanical Impact:	Not expected to be sensitive to mechanical impact.		
Rate of Burning:	Not available.		
Explosive Power:	Not available.		
Sensitivity to Static Discharge:		e to static discharge if	Bisulphide gas has evolved, then Ca vapours are present between the lov ild-up is possible when significant qu
EXTINGUISHING MEDIA			
Fire Extinguishing Media:	Use carbon dioxide or dry chemical r fog. Cool containers with flooding qu moisture may cause the decompositi Carbon Bisulphide vapours. (3)	antities of water until v	vell after the fire is out. Exposure to h
FIRE FIGHTING INSTRUCTIONS			
Instructions to the Fire Fighters:	Use water spray to cool fire-exposed ignition is possible. Clean up immedi watercourses. Avoid accumulation at	ately to eliminate slipp	oing hazard. Do not allow to enter se
Fire Fighting Protective Equipment:	Use self-contained breathing appara	tus and protective clot	hing.

6. ACCIDENTAL RELEASE MEASURES

Information in this section is for responding to spills, leaks or releases in order to prevent or minimize the adverse effects on persons, property and the environment. There may be specific reporting requirements associated with spills, leaks or releases, which change from region to region.

WHMIS Number: 00060600 Date of Revision: 2009 June 17

Page 4 of 8

Containment and Clean-Up Procedures:

In all cases of leak or spill contact vendor at Emergency Number shown on the front page of this MSDS. Avoid accumulation and dispersion of dust to reduce explosion potential. Wear respirator, protective clothing and gloves. Spilled material may cause floors and contact surfaces to become slippery. Any recovered product can be used for the usual purpose, depending on the extent and kind of contamination. Where a package (drum or bag) is damaged and / or leaking, repair it, or place it into an over-pack drum immediately so as to avoid or minimize material loss and contamination of surrounding environment. Replace damaged containers immediately to avoid loss of material and contamination of surrounding atmosphere. Avoid dry sweeping. Do not use compressed air to clean surfaces. Vacuuming or wet sweeping is preferred. Return all material possible to container for proper disposal. Do not flush with water as aqueous solutions or powders that become wet render surfaces extremely slippery. Eliminate all sources of ignition. Collect product for recovery or disposal. For release to land, or storm water runoff, contain discharge by constructing dykes or applying inert absorbent; for release to water, utilize damming and/or water diversion to minimize the spread of contamination. Ventilate enclosed spaces. Notify applicable government authority if release is reportable or could adversely affect the environment.

7. HANDLING AND STORAGE

HANDLING

Handling Practices: Avoid accumulation and dispersion of dust to reduce explosion potential. Ground and bond equipment

and containers to prevent a static charge buildup. Use spark-resistant tools. Use normal "good" industrial hygiene and housekeeping practices. Clean up immediately to eliminate slipping hazard.

Enforce NO SMOKING rules in area of use.

Ventilation Requirements: See Section 8, "Engineering Controls".

Other Precautions: Use only with adequate ventilation and avoid breathing dusts (aerosols, vapours or mists). Avoid

contact with eyes, skin or clothing. Wash thoroughly with soap and water after handling. Wash contaminated clothing thoroughly before re-use. Do not use cutting or welding torches on empty drums that contained this material/product. Absorption via contact with skin, eyes and mucous membranes can

contribute to the overall exposure. Consider measures to prevent absorption by these routes.

STORAGE

Storage Temperature (°C): See below.

Ventilation Requirements: Ventilation should be explosion proof.

Storage Requirements: Store solid Xanthates under cool, dark, dry conditions. Liquid products must be kept cool and used as

quickly as possible. (3) Store in a cool, well-ventilated area. Keep away from heat, sparks and flames. Keep containers closed. Do not expose sealed containers to temperatures above 40° C. Avoid moisture

contamination. Prolonged storage may result in lumping or caking.

Special Materials to be Used for

Packaging or Containers:

Materials of construction for storing the product include: carbon steel. Copper and its alloys should not be used in equipment for storage, handling or transportation. Attacks some types of rubber, plastics and

coatings. Confirm suitability of any material before using.

8. EXPOSURE CONTROLS / PERSONAL PROTECTION

Recommendations listed in this section indicate the type of equipment, which will provide protection against overexposure to this product. Conditions of use, adequacy of engineering or other control measures, and actual exposures will dictate the need for specific protective devices at your workplace.

ENGINEERING CONTROLS

Engineering Controls: Local exhaust ventilation required. Ventilation should be explosion proof. Make up air should be

supplied to balance air that is removed by local or general exhaust ventilation. Avoid accumulation and dispersion of dust to reduce explosion potential. Ventilate low lying areas such as sumps or pits where

dense dust may collect. Enforce NO SMOKING rules.

PERSONAL PROTECTIVE EQUIPMENT (PPE)

Eye Protection: Use chemical safety goggles when there is potential for eye contact. Use full face-shield and chemical

safety goggles when there is potential for contact.

Skin Protection: Gloves and protective clothing made from neoprene, PVC, polyethylene, rubber or plastic should be

impervious under conditions of use. Attacks some types of rubber, plastics and coatings. Prior to use,

user should confirm impermeability. Discard contaminated gloves.

Respiratory Protection: No specific guidelines available. A NIOSH/MSHA-approved air-purifying respirator equipped with dust,

mist, fume cartridges for concentrations up to 2 mg/m³ Potassium Hydroxide. An air-supplied respirator if

concentrations are higher or unknown.

WHMIS Number: 00060600 Date of Revision: 2009 June 17

Page 5 of 8

Other Personal Protective

Equipment:

Avoid accumulation and dispersion of dust to reduce explosion potential. Wear an impermeable apron and boots. Locate safety shower and eyewash station close to chemical handling area. Take all precautions to avoid personal contact. Clothing and footwear that is fire retardant and dissipates static electrical charges should be worn when handling flammable materials. Natural fibers (cotton, wool, leather and linen) should be selected in favour of synthetic materials (rayon, nylon and polyester).

Skin Notation: Contact with skin, eyes and mucous membranes can contribute to the overall exposure and may invalidate the TLV. Consider measures to prevent absorption by these routes.

EXPOSURE GUIDELINES

SUBSTANCE	ACGIH TLV	OSHA	OSHA PEL		NIOSH REL	
	(STEL)	(TWA)	(STEL)	(TWA)	(STEL)	
Potassium Hydroxide	2 mg/m³ (Ceiling)				2 mg/m³ (Ceiling)	
Isoamyl alcohol	125 ppm	100 ppm		100 ppm	125 ppm	
Decomposition Product: Carbon disulfide	_	20 ppm (Skin)	30 ppm (Skin)	1 ppm (Skin)	3 ppm (Skin)	

9. PHYSICAL AND CHEMICAL PROPERTIES (Not intended as Specifications)

Physical State: Solid.

Appearance: Yellow to yellow-green pellets.

Odour: Strong, disagreeable sulphur odour.

Odour Threshold (ppm): 0.02 - 0.21(Carbon Disulphide)

Boiling Range (°C): Not available.

Melting/Freezing Point (°C): 255 - 280 (decomposes). (3)

Vapour Pressure (mm Hg at 20° C):

Vapour Density (Air = 1.0):

Relative Density (g/cc):

Bulk Density:

Viscosity:

Not applicable.

Not applicable.

Not applicable.

Not applicable.

Not applicable.

Not applicable.

Solubility: Soluble in water. Hygroscopic (readily absorbs water).

% Volatile by Volume: < 20. (3)

pH: 10.5 (10 % solution). (3)

Coefficient of Water/Oil Distribution: Not available.

Volatile Organic Compounds (VOC): Not applicable.

Flashpoint (°C): -30. (Carbon Disulphide)

10. STABILITY AND REACTIVITY

CHEMICAL STABILITY

Under Normal Conditions: Unstable. Solid Xanthates are stable when kept cool and dry. Exposure to heat causes decomposition.

Acids and oxidizing agents accelerate aging. In solution, Xanthates will decompose slowly even at room

temperature. (3)

Under Fire Conditions: Flammable. This product may be capable of forming flammable dust clouds in air.

Hazardous Polymerization: Will not occur.

Conditions to Avoid: High temperatures, sparks, open flames and all other sources of ignition. Avoid accumulation and

dispersion of dust to reduce explosion potential. Exposure to heat and moisture may cause the decomposition of xanthates to release flammable, explosive and poisonous Carbon Bisulphide vapours.

(3)

Materials to Avoid: Strong oxidizers. Lewis or mineral acids. Metal Salts. Copper and its alloys.. Contact with acids will

liberate Carbon Bisulphide. Avoid moisture contamination. Contact with water or moisture will liberate Carbon Bisulphide. Mixtures or reactions of alcohols with the following materials may cause explosions: barium perchlorate, chlorine, hypochlorous acid, ethylene oxide, hexamethylene diisocyanate and other isocyanates, nitrogen tetroxide, permonosulfuric acid and tri-isobutyl aluminum. (4) Attacks some types

of rubber, plastics and coatings.

Potassium Amyl Xanthate, Solid

WHMIS Number: 00060600 2009 June 17 Date of Revision:

Page 6 of 8

Decomposition or Combustion

Products:

Thermal decomposition products are toxic and may include Carbon Bisulphide, Potassium sulphide, carbonyl sulphide, Amyl Alcohols, oxides of carbon, sulphur, potassium and irritating gases.

Brenntag Canada Inc.

11. TOXICOLOGICAL INFORMATION

TOXICOLOGICAL DATA:

SUBSTANCE	LD50 (Oral, Rat)	LD50 (Dermal, Rabbit)	LC50 (Inhalation, Rat, 4h)
Potassium Amyl Xanthate	1 000 mg/kg (3)		
Potassium Hydroxide	214 - 365 mg/kg (1,3)	1 260 mg/kg (3)	
Isoamyl alcohol	1 300 mg/kg (1)	3 216 mg/kg (1)	
Decomposition Product: Carbon disulfide	1 200 mg/kg (1)		12 500 mg/m3 (1)

Mutagenicity Data:

The ingredient(s) of this product is (are) not classed as carcinogenic by ACGIH, IARC, OSHA or NTP.

Carcinogenicity Data: Reproductive Data:

This product: No adverse reproductive effects are anticipated.

Teratogenicity Data:

No adverse mutagenic effects are anticipated. No adverse teratogenic effects are anticipated.

Respiratory / Skin Sensitization

Data:

None known.

Synergistic Materials: Alcohols may interact synergistically with chlorinated solvents (example - carbon tetrachloride,

chloroform, bromotrichloromethane), dithiocarbamates (example - disulfiram), dimethylnitrosamine and

thioacetamide. (4)

Carbon Bisulphide: The toxic effects of Carbon Bisulphide, particularly on the nervous system, can be intensified by consumption of alcohol, alcoholism, treatment with disulfiram (Antibuse), and exposure to Hydrogen Sulphide. (4) In animal studies the toxicity of Carbon Bisulphide was intensified by chemicals

such as resperine and amphetamine which act on the nervous system. (4)

Other Studies Relevant to

Material:

None known.

12. ECOLOGICAL INFORMATION

Ecotoxicity: Not available. May be harmful to aquatic life.

Environmental Fate: Not available. Product has an unaesthetic appearance and can be a nuisance. Can be dangerous if

allowed to enter drinking water intakes. Do not contaminate domestic or irrigation water supplies, lakes,

streams, ponds, or rivers.

13. DISPOSAL CONSIDERATIONS

Deactivating Chemicals: Not available.

Waste Disposal Methods: This information applies to the material as manufactured. Reevaluation of the product may be required

by the user at the time of disposal since the product uses, transformations, mixtures and processes may

influence waste classification. Dispose of waste material at an approved (hazardous) waste

treatment/disposal facility in accordance with applicable local, provincial and federal regulations. Do not

dispose of waste with normal garbage, or to sewer systems.

Safe Handling of Residues: See "Waste Disposal Methods".

Disposal of Packaging: Empty containers retain product residue and can be dangerous. Treat package in the same manner as

the product.

14. TRANSPORTATION INFORMATION

CANADIAN TDG ACT SHIPPING DESCRIPTION:

XANTHATES, Class 4.2, UN3342, PG III.

Label(s): Substances Liable To Spontaneous Combustion. Placard: Substances Liable To Spontaneous Combustion.

2009 June 17

Date of Revision:

WHMIS Number: 00060600

Page 7 of 8

ERAP Index: ----- Exemptions: None known.

ERAP Index: ----- Exemptions: None known. US DOT CLASSIFICATION (49CFR 172.101, 172.102):

XANTHATES, Class 4.2, UN3342, PG III.

Label(s): Spontaneously Combustible. Placard: Spontaneously Combustible.

CERCLA-RQ: Not available. Exemptions: None known.

15. REGULATORY INFORMATION

CANADA

CEPA - NSNR: All constituents of this product are included on the DSL.

CEPA - NPRI: Not included.

Controlled Products Regulations Classification (WHMIS):

B-6: Reactive Flammable MaterialD-1B: Toxic (acute effects)D-2B: Toxic (skin and eye irritant)

USA

Environmental Protection Act: All constituents of this product are included on the TSCA inventory.

OSHA HCS (29CFR 1910.1200): Flammable Solid. Toxic. Skin and Eye Irritant.

NFPA: 3 Health, 4 Fire, 0 Reactivity (6)

HMIS: Health, Fire, Reactivity (Not available.)

INTERNATIONAL

Not available.

16. OTHER INFORMATION

REFERENCES

- RTECS-Registry of Toxic Effects of Chemical Substances, Canadian Centre for Occupational Health and Safety RTECS database.
- Clayton, G.D. and Clayton, F.E., Eds., Patty's Industrial Hygiene and Toxicology, 3rd ed., Vol. IIA,B,C, John Wiley and Sons, New York, 1981.
- 3. Supplier's Material Safety Data Sheet(s).
- 4. CHEMINFO, through "CCINFOdisc", Canadian Centre for Occupational Health and Safety, Hamilton, Ontario, Canada.
- 5. Guide to Occupational Exposure Values, 2007, American Conference of Governmental Industrial Hygienists, Cincinnati, 2007.
- 6. Regulatory Affairs Group, Brenntag Canada Inc.
- The British Columbia Drug and Poison Information Centre, Poison Managements Manual, Canadian Pharmaceutical Association, Ottawa, 1981.

The information contained herein is offered only as a guide to the handling of this specific material and has been prepared in good faith by technically knowledgeable personnel. It is not intended to be all-inclusive and the manner and conditions of use and handling may involve other and additional considerations. No warranty of any kind is given or implied and Brenntag Canada Inc. will not be liable for any damages, losses, injuries or consequential damages which may result from the use of or reliance on any information contained herein. This Material Safety Data Sheet is valid for three years.

To obtain revised copies of this or other Material Safety Data Sheets, contact your nearest Brenntag Canada Regional office.

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WHMIS Number: 00060600 Date of Revision: 2009 June 17 Page 8 of 8

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