

Chemwatch Material Safety Data Sheet Review Date: 9-Oct-2008

NA517EC

CHEMWATCH 17-8709 Version No:2.0 CD 2010/1 Page 1 of 17

Section 1 - CHEMICAL PRODUCT AND COMPANY IDENTIFICATION

PRODUCT NAME

POLY-DRILL K-ION

STATEMENT OF HAZARDOUS NATURE

Considered a hazardous substance according to the Controlled Products Regulations

SUPPLIER

Company: Poly- Drill

Address:

2192 - 8800 Venture Ave SE

Calgary

Alberta, T3S 0A2

CAN

Telephone: +1 403 259 5112 Emergency Tel: +1 403 540 7080

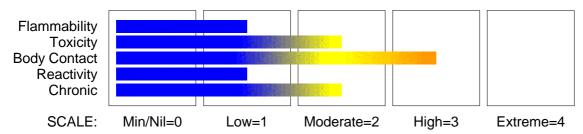
Fax: +1 403 255 7185

PRODUCT USE

Dispersant.

Section 2 - HAZARDS IDENTIFICATION

CHEMWATCH HAZARD RATINGS



CANADIAN WHMIS SYMBOLS





EMERGENCY OVERVIEW

RISK

Harmful if swallowed. Danger of cumulative effects. Risk of serious damage to eyes.

Chemwatch Material Safety Data Sheet Review Date: 9-Oct-2008 NA517EC

CHEMWATCH 17-8709 Version No:2.0 CD 2010/1 Page 2 of 17 Section 2 - HAZARDS IDENTIFICATION

Irritating to respiratory system and skin.

POTENTIAL HEALTH EFFECTS

ACUTE HEALTH EFFECTS

SWALLOWED

- Accidental ingestion of the material may be harmful; animal experiments indicate that ingestion of less than 150 gram may be fatal or may produce serious damage to the health of the individual.
- The material has NOT been classified by EC Directives or other classification systems as "harmful by ingestion". This is because of the lack of corroborating animal or human evidence. The material may still be damaging to the health of the individual, following ingestion, especially where pre-existing organ (e.g liver, kidney) damage is evident. Present definitions of harmful or toxic substances are generally based on doses producing mortality rather than those producing morbidity (disease, ill-health). Gastrointestinal tract discomfort may produce nausea and vomiting. In an occupational setting however, ingestion of insignificant quantities is not thought to be cause for concern.

EYE

- When applied to the eye(s) of animals, the material produces severe ocular lesions which are present twentyfour hours or more after instillation.
- Although the liquid is not thought to be an irritant (as classified by EC Directives), direct contact with the eye may produce transient discomfort characterised by tearing or conjunctival redness (as with windburn).

SKIN

- Evidence exists, or practical experience predicts, that the material either produces inflammation of the skin in a substantial number of individuals following direct contact, and/or produces significant inflammation when applied to the healthy intact skin of animals, for up to four hours, such inflammation being present twenty-four hours or more after the end of the exposure period. Skin irritation may also be present after prolonged or repeated exposure; this may result in a form of contact dermatitis (nonallergic). The dermatitis is often characterised by skin redness (erythema) and swelling (oedema) which may progress to blistering (vesiculation), scaling and thickening of the epidermis. At the microscopic level there may be intercellular oedema of the spongy layer of the skin (spongiosis) and intracellular oedema of the epidermis.
- Skin contact is not thought to have harmful health effects (as classified under EC Directives); the material may still produce health damage following entry through wounds, lesions or abrasions.
- Open cuts, abraded or irritated skin should not be exposed to this material.
- Entry into the blood-stream through, for example, cuts, abrasions, puncture wounds or lesions, may produce systemic injury with harmful effects. Examine the skin prior to the use of the material and ensure that any external damage is suitably protected.

INHALED

- Evidence shows, or practical experience predicts, that the material produces irritation of the respiratory system in a substantial number of individuals following inhalation.
- The material is not thought to produce adverse health effects or irritation of the respiratory tract (as classified by EC Directives using animal models). Nevertheless, good hygiene practice requires that exposure be kept to a minimum and that suitable control measures be used in an occupational setting.
- Inhalation hazard is increased at higher temperatures.
- Not normally a hazard due to non-volatile nature of product.

CHRONIC HEALTH EFFECTS

■ Long-term exposure to the product is not thought to produce chronic effects adverse to health (as classified by EC Directives using animal models); nevertheless exposure by all routes should be minimised as a matter of course.

Chemwatch Material Safety Data Sheet Review Date: 9-Oct-2008 NA517EC

CHEMWATCH 17-8709 Version No:2.0 CD 2010/1 Page 3 of 17

Section 3 - COMPOSITION / INFORMATION ON INGREDIENTS							
NAME dimethyldialkylammonium chloride/ acrylamide polymer acrylamide	CAS RN 26590-05-6 79-06-1	% 30-60					
potassium acetate corrosion inhibitors	127-08-2	10-30					
water	7732-18-5	10-30					

Section 4 - FIRST AID MEASURES

SWALLOWED

- If swallowed do NOT induce vomiting.
- If vomiting occurs, lean patient forward or place on left side (head-down position, if possible) to maintain open airway and prevent aspiration.
- Observe the patient carefully.
- Never give liquid to a person showing signs of being sleepy or with reduced awareness; i.e. becoming unconscious.
- Give water to rinse out mouth, then provide liquid slowly and as much as casualty can comfortably drink.
- · Seek medical advice.

EYE

- 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

- If skin contact occurs:
- Immediately remove all contaminated clothing, including footwear.
- Flush skin and hair with running water (and soap if available).
- Seek medical attention in event of irritation.

INHALED

- If fumes or combustion products are inhaled remove from contaminated area.
- Other measures are usually unnecessary.

NOTES TO PHYSICIAN

■ Treat symptomatically.

Section 5 - FIRE FIGHTING MEASURES

Vapour Pressure (kPa): Not Available Upper Explosive Limit (%): Not Available Specific Gravity (water=1): Not Available Lower Explosive Limit (%): Not Available

EXTINGUISHING MEDIA

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

Chemwatch Material Safety Data Sheet Review Date: 9-Oct-2008 NA517EC

CHEMWATCH 17-8709 Version No:2.0 CD 2010/1 Page 4 of 17 Section 5 - FIRE FIGHTING MEASURES

FIRE FIGHTING

- Alert Fire Brigade and tell them location and nature of hazard.
- Wear breathing apparatus plus protective gloves.
- Prevent, by any means available, spillage from entering drains or water course.
- Use water delivered as a fine spray to control fire and cool adjacent area.
- Avoid spraying water onto liquid pools.
- 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.

GENERAL FIRE HAZARDS/HAZARDOUS COMBUSTIBLE PRODUCTS

- Combustible.
- Slight fire hazard when exposed to heat or flame.
- Heating may cause expansion or decomposition leading to violent rupture of containers.
- On combustion, may emit toxic fumes of carbon monoxide (CO).
- May emit acrid smoke.
- Mists containing combustible materials may be explosive.

Combustion products include: carbon dioxide (CO2), hydrogen chloride, phosgene, nitrogen oxides (NOx), other pyrolysis products typical of burning organic material.

FIRE INCOMPATIBILITY

 Avoid contamination with oxidising agents i.e. nitrates, oxidising acids, chlorine bleaches, pool chlorine etc. as ignition may result.

PERSONAL PROTECTION

Glasses: Gloves: Respirator: Chemical goggles. When handling larger quantities: Particulate

Section 6 - ACCIDENTAL RELEASE MEASURES

MINOR SPILLS

- Remove all ignition sources.
- · Clean up all spills immediately.
- Avoid breathing vapours and contact with skin and eyes.
- Control personal contact 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.

MAJOR SPILLS

- Moderate hazard.
- Clear area of personnel and move upwind.
- Alert Fire Brigade and tell them location and nature of hazard.
- Wear breathing apparatus plus protective gloves.
- Prevent, by any means available, spillage from entering drains or water course.
- No smoking, naked lights or ignition sources.
- Increase ventilation.
- Stop leak if safe to do so.
- Contain spill with sand, earth or vermiculite.
- Collect recoverable product into labelled containers for recycling.
- Absorb remaining product with sand, earth or vermiculite.
- Collect solid residues and seal in labelled drums for disposal.
- Wash area and prevent runoff into drains.
- If contamination of drains or waterways occurs, advise emergency services.

Chemwatch Material Safety Data Sheet Review Date: 9-Oct-2008 NA517EC

CHEMWATCH 17-8709 Version No:2.0 CD 2010/1 Page 5 of 17

Section 7 - HANDLING AND STORAGE

PROCEDURE FOR HANDLING

- Avoid all personal contact, including inhalation.
- Wear protective clothing when risk of exposure occurs.
- Use in a well-ventilated area.
- Prevent concentration in hollows and sumps.
- DO NOT enter confined spaces until atmosphere has been checked.
- Avoid smoking, naked lights or ignition sources.
- Avoid contact with incompatible materials.
- 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 soap and water after handling.
- Work clothes should be laundered separately.
- Use good occupational work practice.
- Observe manufacturer's storing and handling recommendations.
- Atmosphere should be regularly checked against established exposure standards to ensure safe working conditions.

RECOMMENDED STORAGE METHODS

- Metal can or drum
- Packaging as recommended by manufacturer.
- Check all containers are clearly labelled and free from leaks.

STORAGE REQUIREMENTS

- Store in original containers.
- Keep containers securely sealed.
- No smoking, naked lights or ignition sources.
- 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 storing and handling recommendations.

SAFE STORAGE WITH OTHER CLASSIFIED CHEMICALS















May be stored together

O: May be stored together with specific preventions

Must not be stored together

Section 8 - EXPOSURE CONTROLS / PERSONAL PROTECTION

EXPOSURE CONTROLS

US OSHA Permissible Exposure Levels (PELs)

Ζ Material

STEL TWA TWA STEL Peak ppm mg/m³ ppm mg/m³ ppm

Peak Max mg/m³ excursion

ppm

Max mg/m³

Max excursion excursion F/CC duration

(mins)

TWA

Chemwatch Material Safety Data Sheet

Review Date: 9-Oct-2008 NA517EC

CHEMWATCH 17-8709 Version No:2.0 CD 2010/1 Page 6 of 17

CD 2010/1 Page 6 of 17 Section 8 - EXPOSURE CONTROLS / PERSONAL PROTECTION

Z 3	Inert or Nuisance				
	Dust: (d) Respirabl fraction	е			
Z3	Inert or Nuisance Dust: (d) Total dus	15 t			
Z1	Acrylamide	0.3			
Z3	Inert or Nuisance	5			
	Dust: (d) Respirabl	е			
Z3	fraction Inert or Nuisance	15			
23	Dust: (d) Total dus				
Sour	rce	Material	TWA ppm	TWA mg/m³ STEL ppm	STEL mg/m³ Notes
	Oregon_	dimethyldialkylammoni		10	*
	nissible Exposure	um chloride/			
Limi	ts (Z3)	acrylamide polymer (Inert or Nuisance			
		Dust: (d) Total dust)			
US (OSHA Permissible	dimethyldialkylammoni		5	
	osure Levels	um chloride/			
(PEL	₋s) - Table Z3	acrylamide polymer			
		(Inert or Nuisance			
		Dust: (d) Respirable fraction)			
US (OSHA Permissible	dimethyldialkylammoni		15	
	osure Levels	um chloride/			
(PEL	∟s) - Table Z3	acrylamide polymer			
		(Inert or Nuisance			
119 -	Hawaii Air	Dust: (d) Total dust) dimethyldialkylammoni		10	
	taminant Limits	um chloride/		10	
		acrylamide polymer			
		(Particulates not			
		other wise regulated			
HC	Hawaii Air	- Total dust)		E	
	taminant Limits	dimethyldialkylammoni um chloride/		5	
COIII	tarriirarit Lirriits	acrylamide polymer			
		(Particulates not			
		other wise regulated			
		- Respirable			
HC	Oregon	fraction) dimethyldialkylammoni		5	*
	nissible Exposure	um chloride/		5	
	ts (Z3)	acrylamide polymer			
	,	(Inert or Nuisance			
		Dust: (d) Respirable			
HC	Tonnogos	fraction)		E	
	Tennessee upational Exposure	dimethyldialkylammoni um chloride/		5	
	ts - Limits For	acrylamide polymer			
	Contaminants	(Particulates not			
		otherwise regulated			
		Respirable fraction)			

Chemwatch Material Safety Data Sheet Review Date: 9-Oct-2008

NA517EC

CHEMWATCH 17-8709

Version No:2.0

CD 2010/1 Page 7 of 17

CD 2010/1 Page 7 of 17 Section 8 - EXPOSURE CONTROLS / PERSONAL PROTECTION

Source	Material	TWA ppm	TWA mg/m³ STEL ppm	STEL mg/m³ Notes
US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants	dimethyldialkylammoni um chloride/ acrylamide polymer (Particulates not otherwise regulated (PNOR)(f)- Respirable fraction)		5	
US - Michigan Exposure Limits for Air Contaminants	dimethyldialkylammoni um chloride/ acrylamide polymer (Particulates not otherwise regulated, Respirable dust)		5	
Canada - Alberta Occupational Exposure Limits	acrylamide (Acrylamide)		0.03	
Canada - British Columbia Occupational Exposure Limits	acrylamide (Acrylamide, Inhalable Revised 2005)		0.03 (V)	Skin; 2A
Canada - Ontario Occupational Exposure Limits	acrylamide (Acrylamide, inhalable, vapour and aerosol)		0.03	Skin
US OSHA Permissible Exposure Levels (PELs) - Table Z1	acrylamide (Acrylamide)		0.3	
US ACGIH Threshold Limit Values (TLV)	acrylamide (Acrylamide)		0.03	TLV Basis CNS
US NIOSH Recommend Exposure Limits (RELs)	e d crylamide (Acrylamide)		0.03	impairmen
US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	acrylamide (Acrylamide)		0.03	
US - Vermont Permissible Exposure Limits Table Z- 1- A Transitional Limits for Air Contaminants	acrylamide (Acrylamide)		0.3	
US - Vermont Permissible Exposure Limits Table Z- 1- A Final Rule Limits for Air Contaminants	acrylamide (Acrylamide)		0.03	
US - Minnesota Permissible Exposure Limits (PELs)	acrylamide (Acrylamide)		0.03	
US - California Permissible Exposure Limits for Chemical Contaminants	acrylamide (Acrylamide)		0.03	

Chemwatch Material Safety Data Sheet Review Date: 9-Oct-2008

NA517EC

CHEMWATCH 17-8709
Version No:2.0
CD 2010/1 Page 8 of 17
Section 8 - EXPOSURE CONTROLS / PERSONAL PROTECTION

Source	Material	TWA ppm	TWA mg/m³	STEL ppm	STEL mg/m	³ Notes
US - Idaho - Limits	acrylamide		0.3			
for Air Contaminants Canada - Quebec Permissible Exposure Values for Airborne	(Acrylamide) acrylamide (Acrylamide)		0.03			
Contaminants (English) US - Hawaii Air	acrylamide		0.03			
Contaminant Limits US - Alaska Limits	(Acrylamide) acrylamide		0.03			
for Air Contaminants Canada - Yukon Permissible Concentrations for	(Acrylamide) acrylamide (Acrylamide - Skin)	-	0.3	-	0.6	
Airborne Contaminant Substances US - Washington	acrylamide		0.03		0.09	
Permissible exposure limits of air contaminants Canada - Saskatchewan	(Acrylamide)		0.03		0.09	T20, Skin
Occupational Health and Safety Regulations -	(Acrylamide (inhalable fraction++ and vapour))		0.00		0.00	120, OKIII
Contamination Limits Canada - Nova Scotia Occupational Exposure Limits	acrylamide (Acrylamide)		0.03			TLV Basis CNS impairmer
Canada - Prince Edward Island Occupational Exposure	acrylamide (Acrylamide)		0.03			TLV Basis CNS impairmer
Limits US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air	acrylamide (Acrylamide)		0.3			
Contaminants US - Michigan Exposure Limits for Air Contaminants	acrylamide (Acrylamide)		0.03			
US - Oregon Permissible Exposure	acrylamide (Acrylamide)		0.3			
Limits (Z1) Canada - Northwest Territories Occupational Exposure	acrylamide (Acrylamide - Skin)		0.3		0.6	
Limits (English) US OSHA Permissible Exposure Levels (PELs) - Table Z3	potassium acetate (Inert or Nuisance Dust: (d) Respirable		5			
US OSHA Permissible Exposure Levels (PELs) - Table Z3	fraction) potassium acetate (Inert or Nuisance Dust: (d) Total dust)		15			

Chemwatch Material Safety Data Sheet Review Date: 9-Oct-2008

NA517EC

CHEMWATCH 17-8709 Version No:2.0 CD 2010/1 Page 9 of 17

Section 8 - EXPOSURE CONTROLS / PERSONAL PROTECTION

Source	Material	TWA ppm	TWA mg/m³ STEL ppm	STEL mg/m³ Notes
				<u> </u>
US - Oregon Permissible Exposure Limits (Z3)	potassium acetate (Inert or Nuisance Dust: (d) Respirable fraction)		5	*
US - Oregon Permissible Exposure Limits (Z3)	potassium acetate (Inert or Nuisance Dust: (d) Total dust)		10	*

The following materials had no OELs on our records

• water: CAS:7732- 18- 5

EMERGENCY EXPOSURE LIMITS

Material Revised IDLH Value (mg/m3) Revised IDLH Value (ppm)

acrylamide 60

MATERIAL DATA

POLY-DRILL K-ION:

Not available

DIMETHYLDIALKYLAMMONIUM CHLORIDE/ ACRYLAMIDE POLYMER:

■ It is the goal of the ACGIH (and other Agencies) to recommend TLVs (or their equivalent) for all substances for which there is evidence of health effects at airborne concentrations encountered in the workplace.

At this time no TLV has been established, even though this material may produce adverse health effects (as evidenced in animal experiments or clinical experience). Airborne concentrations must be maintained as low as is practically possible and occupational exposure must be kept to a minimum.

NOTE: The ACGIH occupational exposure standard for Particles Not Otherwise Specified (P.N.O.S) does NOT apply.

Sensory irritants are chemicals that produce temporary and undesirable side-effects on the eyes, nose or throat. Historically occupational exposure standards for these irritants have been based on observation of workers' responses to various airborne concentrations. Present day expectations require that nearly every individual should be protected against even minor sensory irritation and exposure standards are established using uncertainty factors or safety factors of 5 to 10 or more. On occasion animal no-observable-effect-levels (NOEL) are used to determine these limits where human results are unavailable. An additional approach, typically used by the TLV committee (USA) in determining respiratory standards for this group of chemicals, has been to assign ceiling values (TLV C) to rapidly acting irritants and to assign short-term exposure limits (TLV STELs) when the weight of evidence from irritation, bioaccumulation and other endpoints combine to warrant such a limit. In contrast the MAK Commission (Germany) uses a five-category system based on intensive odour, local irritation, and elimination half-life. However this system is being replaced to be consistent with the European Union (EU) Scientific Committee for Occupational Exposure Limits (SCOEL); this is more closely allied to that of the USA.

OSHA (USA) concluded that exposure to sensory irritants can:

- cause inflammation
- cause increased susceptibility to other irritants and infectious agents
- lead to permanent injury or dysfunction
- permit greater absorption of hazardous substances and
- acclimate the worker to the irritant warning properties of these substances thus increasing the risk of overexposure.

ACRYLAMIDE:

■ For acrylamide:

Toxic effects on the central nervous system are generally reversible if exposure is terminated although prolonged exposure may significantly increase the time of recovery. These effects include muscle weakness, ataxia, incoordination, tremours and hallucination. Based on feeding studies in sensitive animals it is

Chemwatch Material Safety Data Sheet Review Date: 9-Oct-2008 NA517EC

CHEMWATCH 17-8709 Version No:2.0 CD 2010/1 Page 10 of 17

Section 8 - EXPOSURE CONTROLS / PERSONAL PROTECTION

recommended that no more than 0.05 mg/kg/day be absorbed by workmen and is consistent with data indicating a no effect level between 0.014 and 0.1 mg/kg/day. A TLV-TWA of 0.3 mg/m3 has been calculated assuming a respiratory exchange of 10 m3/day but has been recalculated based on the carcinogenic response of rats.

POTASSIUM ACETATE:

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- lead to permanent injury or dysfunction
- permit greater absorption of hazardous substances and
- acclimate the worker to the irritant warning properties of these substances thus increasing the risk of overexposure.

WATER:

■ No exposure limits set by NOHSC or ACGIH.

PERSONAL PROTECTION









EYE

- · Safety glasses with side shields
- Chemical goggles.
- Contact lenses may pose a special hazard; soft contact lenses may absorb and concentrate irritants. A written policy document, describing the wearing of lens 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].

Chemwatch Material Safety Data Sheet Review Date: 9-Oct-2008 NA517EC

CHEMWATCH 17-8709 Version No:2.0 CD 2010/1 Page 11 of 17

Section 8 - EXPOSURE CONTROLS / PERSONAL PROTECTION

HANDS/FEET

■ Wear general protective gloves, eg. light weight rubber gloves.

OTHER

- No special equipment needed when handling small quantities.
- OTHERWISE:
- · Overalls.
- Barrier cream.
- Eyewash unit.

RESPIRATOR

■ Selection of the Class and Type of respirator will depend upon the level of breathing zone contaminant and the chemical nature of the contaminant. Protection Factors (defined as the ratio of contaminant outside and inside the mask) may also be important.

Breathing Zone Level	Maximum Protection	Half- face Respirator	Full- Face Respirator
ppm (volume)	Factor		
1000	10	- AUS P	-
1000	50	-	- AUS P
5000	50	Airline *	-
5000	100	-	- 2 P
10000	100	-	- 3 P
	100+		Airline**

^{* -} Continuous Flow ** - Continuous-flow or positive pressure demand.

The local concentration of material, quantity and conditions of use determine the type of personal protective equipment required.

Use appropriate NIOSH-certified respirator based on informed professional judgement. In conditions where no reasonable estimate of exposure can be made, assume the exposure is in a concentration IDLH and use NIOSH-certified full face pressure demand SCBA with a minimum service life of 30 minutes, or a combination full facepiece pressure demand SAR with auxiliary self-contained air supply. Respirators provided only for escape from IDLH atmospheres shall be NIOSH-certified for escape from the atmosphere in which they will be used.

ENGINEERING CONTROLS

■ General exhaust is adequate under normal operating conditions. If risk of overexposure exists, wear SAA approved respirator. Correct fit is essential to obtain adequate protection. Provide adequate ventilation in warehouse or closed storage areas.

Section 9 - PHYSICAL AND CHEMICAL PROPERTIES

PHYSICAL PROPERTIES

Liquid.

Mixes with water

State	Liquid	Molecular Weight	Not Applicable
Melting Range (℃)	Not Available	Viscosity	Not Available
Boiling Range (℃)	>100	Solubility in water (g/L)	Miscible
Flash Point (℃)	Not Available	pH (1% solution)	Not Available
Decomposition Temp (℃)	Not Available	pH (as supplied)	Not Available
Autoignition Temp (℃)	Not Available	Vapour Pressure (kPa)	Not Available
Upper Explosive Limit (%)	Not Available	Specific Gravity (water=1)	Not Available

Chemwatch Material Safety Data Sheet Review Date: 9-Oct-2008

NA517EC

CHEMWATCH 17-8709 Version No:2.0 CD 2010/1 Page 12 of 17

Section 9 - PHYSICAL AND CHEMICAL PROPERTIES

Not Available Relative Vapour Density Not Available Lower Explosive Limit (%)

(air=1)

Volatile Component (%vol) Not Available **Evaporation Rate** Not Available

APPEARANCE

Clear liquid; mixes with water.

Value Material

ACRYLAMIDE:

log Kow - 0.67

Section 10 - CHEMICAL STABILITY AND REACTIVITY INFORMATION

CONDITIONS CONTRIBUTING TO INSTABILITY

■ Product is considered stable and hazardous polymerisation will not occur.

STORAGE INCOMPATIBILITY

Avoid reaction with oxidising agents.

For incompatible materials - refer to Section 7 - Handling and Storage.

Section 11 - TOXICOLOGICAL INFORMATION

Poly-Drill K-Ion

TOXICITY AND IRRITATION

■ Not available. Refer to individual constituents.

DIMETHYLDIALKYLAMMONIUM CHLORIDE/ ACRYLAMIDE POLYMER:

- unless otherwise specified data extracted from RTECS Register of Toxic Effects of Chemical Substances.
- Most undiluted cationic surfactants satisfy the criteria for classification as Harmful (Xn) with R22 and as Irritant (Xi) for skin and eyes with R38 and R41.

No significant acute toxicological data identified in literature search.

POTASSIUM ACETATE:

■ unless otherwise specified data extracted from RTECS - Register of Toxic Effects of Chemical Substances.

TOXICITY **IRRITATION** Oral (rat) LD50: 3250 mg/kg Nil Reported

■ Asthma-like symptoms may continue for months or even years after exposure to the material ceases. This may be due to a non-allergenic condition known as reactive airways dysfunction syndrome (RADS) which can occur following exposure to high levels of highly irritating compound. Key criteria for the diagnosis of RADS include the absence of preceding respiratory disease, in a non-atopic individual, with abrupt onset of persistent asthma-like symptoms within minutes to hours of a documented exposure to the irritant. A reversible airflow pattern, on spirometry, with the presence of moderate to severe bronchial hyperreactivity on methacholine challenge testing and the lack of minimal lymphocytic inflammation, without eosinophilia, have also been included in the criteria for diagnosis of RADS. RADS (or asthma) following an irritating inhalation is an infrequent disorder with rates related to the concentration of and duration of exposure to the irritating substance. Industrial bronchitis, on the other hand, is a disorder that occurs as result of exposure due to high concentrations of irritating substance (often particulate in nature) and is completely reversible after exposure ceases. The disorder is characterised by dyspnea, cough and mucus production.

Chemwatch Material Safety Data Sheet Review Date: 9-Oct-2008 NA517EC

CHEMWATCH 17-8709

Version No:2.0

CD 2010/1 Page 13 of 17

Section 11 - TOXICOLOGICAL INFORMATION

WATER:

■ No significant acute toxicological data identified in literature search.

CARCINOGEN

Acrylamide (NB: Overall

evaluation upgraded from 2B to 2A with supporting evidence

from other relevant

data)

International Agency Group for Research on Cancer

(IARC) - Agents Reviewed by the IARC

Monographs

SKIN

acrylamide

acrylamide Canada - Ontario Occupational Exposure

Notes

Skin

Limits - Skin

Canada - Quebec Permissible Exposure Values

Notes

2A

Skin

1

Skin; 2A

for Airborne Contaminants - Skin (French)
acrylamide Canada - British Columbia Occupational

Canada - British Columbia Occupational Exposure Limits - Skin

Notation

Exposure Limits

acrylamide Canada - Alberta Occupational Exposure

Limits - Skin

Substance Interaction

Section 12 - ECOLOGICAL INFORMATION

Refer to data for ingredients, which follows:

ACRYLAMIDE:

POTASSIUM ACETATE:

DIMETHYLDIALKYLAMMONIUM CHLORIDE/ ACRYLAMIDE POLYMER:

■ DO NOT discharge into sewer or waterways.

DIMETHYLDIALKYLAMMONIUM CHLORIDE/ ACRYLAMIDE POLYMER:

- Toxic to aquatic organisms.
- 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.
- Although inorganic chloride ions are not normally considered toxic they can exist in effluents at acutely toxic levels (chloride >3000 mg/l). the resulting salinity can exceed the tolerances of most freshwater organisms.

Inorganic chlorine eventually finds its way into the aqueous compartment and as such is bioavailable. Incidental exposure to inorganic chloride may occur in occupational settings where chemicals management policies are improperly applied. The toxicity of chloride salts depends on the counter-ion (cation) present; that of chloride itself is unknown. Chloride toxicity has not been observed in humans except in the special case of impaired sodium chloride metabolism, e.g. in congestive heart failure. Healthy individuals can tolerate the intake of large quantities of chloride provided that there is a concomitant intake of fresh water.

Although excessive intake of drinking-water containing sodium chloride at concentrations above 2.5 g/litre has been reported to produce hypertension, this effect is believed to be related to the sodium ion concentration.

Chloride concentrations in excess of about 250 mg/litre can give rise to detectable taste in water, but the threshold depends upon the associated cations. Consumers can, however, become accustomed to concentrations in excess of 250 mg/litre. No health-based guideline value is proposed for chloride in drinking-water. In humans, 88% of chloride is extracellular and contributes to the osmotic activity of body fluids. The electrolyte balance in the body is maintained by adjusting total dietary intake and by excretion via the kidneys and gastrointestinal tract. Chloride is almost completely absorbed in normal individuals, mostly from the proximal half of the small intestine. Normal fluid loss amounts to about 1.5?2 liters/day, together with about 4 g of chloride per day. Most (90 - 95%) is excreted in the urine, with minor amounts in faeces (4-%)

Chemwatch Material Safety Data Sheet Review Date: 9-Oct-2008 NA517EC

CHEMWATCH 17-8709

Version No:2.0

CD 2010/1 Page 14 of 17

Section 12 - ECOLOGICAL INFORMATION

and sweat (2%)

Chloride increases the electrical conductivity of water and thus increases its corrosivity. In metal pipes, chloride reacts with metal ions to form soluble salts thus increasing levels of metals in drinking-water. In lead pipes, a protective oxide layer is built up, but chloride enhances galvanic corrosion. It can also increase the rate of pitting corrosion of metal pipes.

■ For organic cationics

Cationic substances, and their polymers and those polymers that are reasonably anticipated to become cationic in the natural aquatic environment (pH range 4-9) may be environmental hazards.

Exempt from this concern are those polymers to be used only in solid phase, such as ion-exchange resins, and where the FGEW (Functional Group Equivalent Weight) of cationic groups is not 5000 and above.

The numerous studies of aquatic toxicity, many of which were conducted in natural waters with and without added effluents, indicate that the source and composition of the test water dramatically affects the toxicity of the test substance. These results are consistent with the known behavior of these materials in the environment. Cationic substances in the environment instantaneously form complexes with naturally occurring negatively charged constituents in sewage, soils, sediments, and with dissolved humic substances in surface waters. This complexation behavior results in reduced bioavailability in actual environmental conditions that is not adequately represented by standard laboratory assays and/or predictions by various QSAR models. Ecotoxicity:

These chemicals, by the nature of their surfactant properties, are toxic to aquatic organisms at low concentrations. Cationic groups such as alkylsulfoniums, alkylphosphoniums and quaternary ammonium polymers are highly toxic to fish and other aquatic organisms. Similarly potentially cationic groups such as amines and isocyanates are of concern. Some cationics, however, may fall into the category of PLCs (polymers of low concern) provided they possess low charge density, and/or are not water-soluble or are not self-dispersing polycarboxylates or poly- (aromatic or aliphatic) sulfonate polymers.

The toxicity of quaternary ammonium compounds is known to be greatly reduced in the environment because of preferential binding to dissolved organics in surface water.

ACRYLAMIDE:

■ Hazardous Air Pollutant: Yes
 ■ Fish LC50 (96hr.) (mg/l): 130
 ■ Daphnia magna EC50 (48hr.) (mg/l): 105
 ■ log Kow (Prager 1995): - 0.67

■ For acrylamide:

log Kow: -0.67 Half-life (hr) air: 6.6

Half-life (hr) H2O surface water: 192-288

Henry's atm m3/mol: 3.20E-10

BOD 5: 0.97,72.8% BCF: 0.86-1.12 Environmental fate:

Terrestrial Fate: When released on soil, acrylamide would be expected to leach readily into the ground and biodegrade within a few weeks.

Aquatic Fate: Since acrylamide has been found to degrade in distilled water over a period of 1-2 months and river water in 8-12 days, it is unlikely that acrylamide would have a long residence time in natural waters. However, acclimation of the microorganisms is important. In systems in which the residence time is relatively short such as in sewage works and water treatment facilities, acrylamide may not be completely degraded, and acrylamide has been detected in the effluent from a sewage treatment plant. Adsorption to sediment and volatilization will not be appreciable.

Atmospheric Fate: If released into the atmosphere, acrylamide in the vapor phase should react with photochemically produced hydroxyl radicals (estimated half-life 6.6 hr). Due to its high solubility in water, it should be scavenged by rain and fog.

Ecotoxicity:

Fish:LC50 (24-96 h): 130-460 mg/L Bioaccumulation: not significant Anaerobic effects: some degradation Degradation Biological: significant

processes Abiotic: nophotolysis, hydrolysis, RxnOH*.

Chemwatch Material Safety Data Sheet Review Date: 9-Oct-2008 NA517EC

CHEMWATCH 17-8709
Version No:2.0
CD 2010/1 Page 15 of 17
Section 12 - ECOLOGICAL INFORMATION

POTASSIUM ACETATE:

- Acetic acid and its salts (the acetates) can be grouped together because of their close structural relationships, their natural occurrence in plants and animals, and their fundamental role in cell metabolism, particularly in the tricarboxylic acid cycle (also known as the citric acid or Kreb's cycle), which is where humans get their energy.
- Acetic acid is degraded photochemically in the atmosphere to produce hydroxyl radicals (estimated typical half-life of 22 days). Physical removal of acetates on atmospheric particulates may occur via wet or dry deposition.
- Natural water will neutralise dilute solutions of acetic acid.
- Spills of acetic acid on soil will readily biodegrade the biodegradation rate for acetic acid after 14 days under aerobic conditions is 74 days.
- In invertebrates the toxicity of acetic acid (EC50 = 50-450 mg/L, depending on test species) -under static conditions, the 48 hour EC50 value for acetic acid is 65 mg/L for aquatic invertebrates (the test media was not neutralised). When the test solutions are neutralised, to form acetates, the static 48 hour EC50 for acetic acid is 6000 mg/L. In renewal systems with aquatic invertebrates, 48 hour EC50s for acetic acid are 100 mg/L and 180 mg/L.
- Fish LC50 (96 h): 75-88 mg/L.
- Acetic acid is not expected to bioconcentrate in the aquatic system.
- Low concentrations of acetic acid are harmful to fish.
- Drinking water standards: none available.
- Soil Guidelines: none available.
- Air Quality Standards: none available.

May be harmful to fauna if not disposed of according to Section 13 and legislative requirements. [AMC]

Ecotoxicity

Ingredient	Persistence: Water/Soil	Persistence: Air	Bioaccumulation	Mobility
acrylamide	LOW		LOW	HIGH
water	LOW		LOW	HIGH

Section 13 - DISPOSAL CONSIDERATIONS

US EPA Waste Number & Descriptions

B. Component Waste Numbers

When acrylamide is present as a solid waste as a discarded commercial chemical product, off-specification species, as a container residue, or a spill residue, use EPA waste number U007 (waste code T).

Disposal Instructions

All waste must be handled in accordance with local, state and federal regulations.

- Recycle wherever possible or consult manufacturer for recycling options.
- Consult State Land Waste Authority for disposal.
- Bury or incinerate residue at an approved site.
- Recycle containers if possible, or dispose of in an authorised landfill.

Section 14 - TRANSPORTATION INFORMATION

NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS: TDG, IATA, IMDG

Chemwatch Material Safety Data Sheet Review Date: 9-Oct-2008 NA517EC

CHEMWATCH 17-8709 Version No:2.0 CD 2010/1 Page 16 of 17

Section 15 - REGULATORY INFORMATION

REGULATIONS

US EPCRA Section 313 Chemical List

Ingredient CAS % de minimus

concentration

acrylamide 79- 06- 1 0.1

US CERCLA List of Hazardous Substances and Reportable Quantities

Ingredient CAS RQ

acrylamide 79- 06- 1 5000 lb (2270 kg)

Regulations for ingredients

dimethyldialkylammonium chloride/ acrylamide polymer (CAS: 26590-05-6,108464-53-5) is found on the following regulatory lists;

"Canada Domestic Substances List (DSL)"

acrylamide (CAS: 79-06-1) is found on the following regulatory lists;

"Canada - Alberta Occupational Exposure Limits", "Canada - British Columbia Occupational Exposure Limits", "Canada - Northwest Territories Occupational Exposure Limits (English)", "Canada - Nova Scotia Occupational Exposure Limits", "Canada - Ontario Occupational Exposure Limits", "Canada - Prince Edward Island Occupational Exposure Limits", "Canada - Quebec Permissible Exposure Values for Airborne Contaminants (English)", "Canada - Saskatchewan Occupational Health and Safety Regulations - Designated Chemical Substances", "Canada - Yukon Permissible Concentrations for Airborne Contaminant Substances", "Canada ARET (Accelerated Reduction / Elimination of Toxics) Substance List", "Canada Domestic Substances List (DSL)", "Canada Ingredient Disclosure List (SOR/88-64)", "Canada National Pollutant Release Inventory (NPRI)", "Canada Toxicological Index Service - Workplace Hazardous Materials Information System - WHMIS (English)", "Canada Toxicological Index Service - Workplace Hazardous Materials Information System - WHMIS (French)", "GESAMP/EHS Composite List - GESAMP Hazard Profiles", "IMO IBC Code Chapter 17: Summary of minimum requirements", "IMO MARPOL 73/78 (Annex II) - List of Noxious Liquid Substances Carried in Bulk", "International Agency for Research on Cancer (IARC) - Agents Reviewed by the IARC Monographs", "International Chemical Secretariat (ChemSec) REACH SIN* List ("Substitute It Now!) 1.0", "OECD Representative List of High Production Volume (HPV) Chemicals", "WHO Guidelines for Drinking-water"

potassium acetate (CAS: 127-08-2) is found on the following regulatory lists;

"Canada Toxicological Index Service - Workplace Hazardous Materials Information System - WHMIS (English)", "Canada Toxicological Index Service - Workplace Hazardous Materials Information System - WHMIS (French)", "CODEX General Standard for Food Additives (GSFA) - Additives Permitted for Use in Food in General, Unless Otherwise Specified, in Accordance with GMP", "International Council of Chemical Associations (ICCA) - High Production Volume List", "OECD Representative List of High Production Volume (HPV) Chemicals"

water (CAS: 7732-18-5) is found on the following regulatory lists;

"Canada Domestic Substances List (DSL)", "Canada Toxicological Index Service - Workplace Hazardous Materials Information System - WHMIS (English)", "Canada Toxicological Index Service - Workplace Hazardous Materials Information System - WHMIS (French)", "IMO IBC Code Chapter 18: List of products to which the Code does not apply", "OECD Representative List of High Production Volume (HPV) Chemicals"

No data for Poly-Drill K-Ion (CW: 17-8709)

This product has been classified in accordance with the hazard criteria of the Controlled Products Regulations and the MSDS contains all the information required by the Controlled Products Regulations.

Section 16 - OTHER INFORMATION

INGREDIENTS WITH MULTIPLE CAS NUMBERS

Ingredient Name CAS

dimethyldialkylammonium chloride/ acrylamide polymer 26590- 05- 6, 108464- 53- 5

REPRODUCTIVE HEALTH GUIDELINES

Ingredient ORG UF Endpoint CR Adeq TLV acrylamide 0.03 mg/m3 NA NA 220 Yes

■ These exposure guidelines have been derived from a screening level of risk assessment and should not be construed as unequivocally safe limits. ORGS represent an 8-hour time-weighted average unless specified otherwise.

CR = Cancer Risk/10000; UF = Uncertainty factor:

Chemwatch Material Safety Data Sheet Review Date: 9-Oct-2008 NA517EC

CHEMWATCH 17-8709 Version No:2.0 CD 2010/1 Page 17 of 17 Section 16 - OTHER INFORMATION

TLV believed to be adequate to protect reproductive health:

LOD: Limit of detection

Toxic endpoints have also been identified as:

D = Developmental; R = Reproductive; TC = Transplacental carcinogen Jankovic J., Drake F.: A Screening Method for Occupational Reproductive American Industrial Hygiene Association Journal 57: 641-649 (1996).

EXPOSURE STANDARD FOR MIXTURES

- "Worst Case" computer-aided prediction of spray/ mist or fume/ dust components and concentration:
- Composite Exposure Standard for Mixture (TWA) :100 mg/m³.
- Classification of the mixture 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/references.
- The (M)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.
- For detailed advice on Personal Protective Equipment, refer to the following Canadian Standards: CAN/CSA-Z195 Protective Footwear

Z195.1 - Guideline on Selection, Use, and Care of Protective Footwear

CAN/CSA-Z94.3 - Industrial Eye and Face Protectors

Z94.3.1 - Protective Eyewear User's Guide

CSA-Z94.4 - Selection, Use, and Care of Respirators

CAN/CSA-Z180.1 - Compressed Breathing Air and Systems.

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