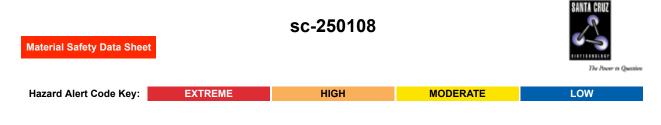
# Hexamethyleneimine



# Section 1 - CHEMICAL PRODUCT AND COMPANY IDENTIFICATION

## PRODUCT NAME

Hexamethyleneimine

# STATEMENT OF HAZARDOUS NATURE

CONSIDERED A HAZARDOUS SUBSTANCE ACCORDING TO OSHA 29 CFR 1910.1200.



## SUPPLIER

Company: Santa Cruz Biotechnology, Inc. Address: 2145 Delaware Ave Santa Cruz, CA 95060 Telephone: 800.457.3801 or 831.457.3800 Emergency Tel: CHEMWATCH: From within the US and Canada: 877-715-9305 Emergency Tel: From outside the US and Canada: +800 2436 2255 (1-800-CHEMCALL) or call +613 9573 3112

PRODUCT USE

Intermediate for pharmaceuticals, agricultural and rubber chemicals.

#### **SYNONYMS**

C6-H13-N, "11-azepine, hexahydro-", "11-azepine, hexahydro-", azacycloheptane, 1-azacyloheptane, 1-azacyloheptane, cyclohexamethyleneimine, G-0, hexahydroazepine, hexahydro-1H-azepine, homopiperidine, perhydroazepine, "piperidine, homo-"

# Section 2 - HAZARDS IDENTIFICATION

## **CANADIAN WHMIS SYMBOLS**



# EMERGENCY OVERVIEW RISK

Toxic if swallowed. Causes burns. Risk of serious damage to eyes. Harmful by inhalation and in contact with skin. Flammable.

# POTENTIAL HEALTH EFFECTS

## ACUTE HEALTH EFFECTS

#### SWALLOWED

• Toxic effects may result from the accidental ingestion of the material; animal experiments indicate that ingestion of less than 40 gram may be fatal or may produce serious damage to the health of the individual.

• The material can produce chemical burns within the oral cavity and gastrointestinal tract following ingestion.

• Amines without benzene rings when swallowed are absorbed throughout the gut. Corrosive action may cause damage throughout the gastrointestinal tract. They are removed through the liver, kidney and intestinal mucosa by enzyme breakdown.

• Exposure to the piperidines may result in increases blood pressure and heart rate, nausea, vomiting, salivation, labored breathing, muscular weakness, paralysis and convulsions. It may also excite the senses of hearing and touch.

#### EYE

• The material can produce chemical burns to the eye following direct contact. Vapors or mists may be extremely irritating.

• If applied to the eyes, this material causes severe eye damage.

• Vapors of volatile amines irritate the eyes, causing excessive secretion of tears, inflammation of the conjunctiva and slight swelling of the cornea, resulting in "halos" around lights. This effect is temporary, lasting only for a few hours. However this condition can reduce the efficiency of undertaking skilled tasks, such as driving a car. Direct eye contact with liquid volatile amines may produce eye damage, permanent for the lighter species.

#### SKIN

• Skin contact with the material may be harmful; systemic effects may resultfollowing absorption.

- The material can produce chemical burns following direct contactwith the skin.
- Open cuts, abraded or irritated skin should not be exposed to this material.

• Entry into the blood-stream, through, for example, cuts, abrasions or lesions, may produce systemic injury with harmful effects. Examine the skin prior to the use of the material and ensure that any external damage is suitably protected.

• Volatile amine vapors produce irritation and inflammation of the skin. Direct contact can cause burns. They may be absorbed through the skin and cause similar effects to swallowing, leading to death. The skin may exhibit whiteness, redness and wheals.

#### INHALED

• If inhaled, this material can irritate the throat andlungs of some persons.

- Inhalation of aerosols (mists, fumes), generated by the material during the course of normal handling, may be harmful.
- Inhalation hazard is increased at higher temperatures.

• Inhalation of quantities of liquid mist may be extremely hazardous, even lethal due to spasm, extreme irritation of larynx and bronchi, chemical pneumonitis and pulmonary edema.

• Inhalation of amine vapors may cause irritation of the mucous membrane of the nose and throat, and lung irritation with respiratory distress and cough. Swelling and inflammation of the respiratory tract is seen in serious cases; with headache, nausea, faintness and anxiety There may also be wheezing.

## **CHRONIC HEALTH EFFECTS**

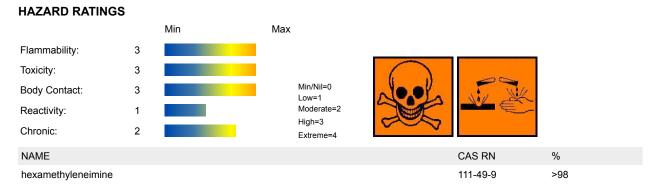
• Repeated or prolonged exposure to corrosives may result in the erosion of teeth, inflammatory and ulcerative changes in the mouth and necrosis (rarely) of the jaw. Bronchial irritation, with cough, and frequent attacks of bronchial pneumonia may ensue. Gastrointestinal disturbances may also occur. Chronic exposures may result in dermatitis and/or conjunctivitis.

There has been some concern that this material can cause cancer or mutations but there is not enough data to make an assessment.

Limited evidence suggests that repeated or long-term occupational exposure may produce cumulative health effects involving organs or biochemical systems.

Secondary amines may react with nitrites to form potentially carcinogenicN-nitrosamines.

# Section 3 - COMPOSITION / INFORMATION ON INGREDIENTS



# **Section 4 - FIRST AID MEASURES**

# SWALLOWED

• For advice, contact a Poisons Information Center 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.

#### EYE

- If this product comes in contact with the eyes:
- Immediately hold eyelids apart and flush the eye continuously with running water.
- Ensure complete irrigation of the eye by keeping eyelids apart and away from eye and moving the eyelids by occasionally lifting the upper and lower lids.
- Continue flushing until advised to stop by the Poisons Information Center or a doctor, or for at least 15 minutes.
- Transport to hospital or doctor without delay.
- Removal of contact lenses after an eye injury should only be undertaken by skilled personnel.

#### SKIN

- 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 Center.
- Transport to hospital, or doctor.

#### INHALED

- 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.

Inhalation of vapors or aerosols (mists, fumes) may cause lung edema. Corrosive substances may cause lung damage (e.g. lung edema, fluid in the lungs). As this reaction may be delayed up to 24 hours after exposure, affected individuals need complete rest (preferably in semi-recumbent posture) and must be kept under medical observation even if no symptoms are (yet) manifested. Before any such manifestation, the administration of a spray containing a dexamethasone derivative or beclomethasone derivative may be considered. This must definitely be left to a doctor or person authorized by him/her. (ICSC13719).

## NOTES TO PHYSICIAN

• For acute or short-term repeated exposures to highly alkaline materials:

- Respiratory stress is uncommon but present occasionally because of soft tissue edema.
- Unless endotracheal intubation can be accomplished under direct vision, cricothyroidotomy or tracheotomy may be necessary.
- Oxygen is given as indicated.
- The presence of shock suggests perforation and mandates an intravenous line and fluid administration.
- Damage due to alkaline corrosives occurs by liquefaction necrosis whereby the saponification of fats and solubilization of proteins allow deep penetration into the tissue.

Alkalis continue to cause damage after exposure.

INGESTION:

- Milk and water are the preferred diluents
- No more than 2 glasses of water should be given to an adult.
- Neutralizing agents should never be given since exothermic heat reaction may compound injury.
- \* Catharsis and emesis are absolutely contra-indicated.
- \* Activated charcoal does not absorb alkali.
- \* Gastric lavage should not be used.

Supportive care involves the following:

- Withhold oral feedings initially.
- If endoscopy confirms transmucosal injury start steroids only within the first 48 hours.
- Carefully evaluate the amount of tissue necrosis before assessing the need for surgical intervention.
- Patients should be instructed to seek medical attention whenever they develop difficulty in swallowing (dysphagia).

SKIN AND EYE:

- Injury should be irrigated for 20-30 minutes.
- Eye injuries require saline. [Ellenhorn & Barceloux: Medical Toxicology]

Depending on the degree of exposure, periodic medical examination is indicated. The symptoms of lung edema often do not manifest until a few hours have passed and they are aggravated by physical effort. Rest and medical observation is therefore essential. Immediate administration of an appropriate spray, by a doctor or a person authorized by him/her should be considered. (ICSC24419/24421.

|                            | Section 5 - FIRE FIGHTING MEASURES |  |
|----------------------------|------------------------------------|--|
| Vapor Pressure (mmHg):     | 7.351 @ 21.1 C                     |  |
| Upper Explosive Limit (%): | Not available                      |  |

Specific Gravity (water=1):

0.880 Not available

## **EXTINGUISHING MEDIA**

Lower Explosive Limit (%):

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

### **FIRE FIGHTING**

- •
- Alert Emergency Responders and tell them location and nature of hazard.
- May be violently or explosively reactive.
- Wear full body protective clothing with breathing apparatus.
- Prevent, by any means available, spillage from entering drains or water course.
- Consider evacuation (or protect in place).
- Fight fire from a safe distance, with adequate cover.
- If safe, switch off electrical equipment until vapor fire hazard removed.
- 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.
- Equipment should be thoroughly decontaminated after use.

## GENERAL FIRE HAZARDS/HAZARDOUS COMBUSTIBLE PRODUCTS

- •
- Liquid and vapor are highly flammable.
- Severe fire hazard when exposed to heat, flame and/or oxidizers.
- Vapor may travel a considerable distance to source of ignition.
- Heating may cause expansion or decomposition leading to violent rupture of containers.
- On combustion, may emit toxic fumes of carbon monoxide (CO).
- Combustion products include: carbon dioxide (CO2), nitrogen oxides (NOx), other pyrolysis products typical of burning organic material.

#### FIRE INCOMPATIBILITY

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

## PERSONAL PROTECTION

Glasses: Full face- shield. Gloves: Respirator: Type AK-P Filter of sufficient capacity

## Section 6 - ACCIDENTAL RELEASE MEASURES

#### MINOR SPILLS

- ,
- Remove all ignition sources.
- Clean up all spills immediately.
- Avoid breathing vapors and contact with skin and eyes.
- Control personal contact by using protective equipment.
- Contain and absorb small quantities with vermiculite or other absorbent material.
- Wipe up.
- Collect residues in a flammable waste container.
- Drains for storage or use areas should have retention basins for pH adjustments and dilution of spills before discharge or disposal of material.
- Check regularly for spills and leaks.

MAJOR SPILLS

· Chemical Class: amines, alkyl

For release onto land: recommended sorbents listed in order of priority.

| SORBENT TYPE                          | RANK | APPLICATION | COLLECTION | LIMITATIONS    |
|---------------------------------------|------|-------------|------------|----------------|
| LAND SPILL - SMALL                    |      |             |            |                |
| cross-linked polymer -<br>particulate | 1    | shovel      | shovel     | R, W, SS       |
| cross-linked polymer -<br>pillow      | 1    | throw       | pitchfork  | R,DGC, RT      |
| sorbent clay - particulate            | 2    | shovel      | shovel     | R, I, P        |
| wood fiber - pillow                   | 3    | throw       | pitchfork  | R, P, DGC, RT, |

| treated wood fibre - pillow                  | 3 | throw  | pitchfork  | DGC, RT         |
|--|---|--------|------------|-----------------|
| foamed glass - pillow<br>LAND SPILL - MEDIUM | 4 | throw  | pitchfork  | R, P, DGC, RT   |
| cross-linked polymer<br>-particulate         | 1 | blower | skiploader | R, W, SS        |
| cross-linked polymer -<br>pillow             | 2 | throw  | skiploader | R, DGC, RT      |
| sorbent clay - particulate                   | 3 | blower | skiploader | R, I, P         |
| polypropylene - particulate                  | 3 | blower | skiploader | W, SS, DGC      |
| expanded mineral -<br>particulate            | 4 | blower | skiploader | R, I, W, P, DGC |
| polypropylene - mat                          | 4 | throw  | skiploader | DGC, RT         |

Legend

DGC: Not effective where ground cover is dense

- R; Not reusable
- I: Not incinerable

P: Effectiveness reduced when rainy

RT:Not effective where terrain is rugged

SS: Not for use within environmentally sensitive sites

W: Effectiveness reduced when windy

Reference: Sorbents for Liquid Hazardous Substance Cleanup and Control;

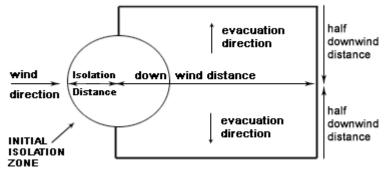
R.W Melvold et al: Pollution Technology Review No. 150: Noyes Data Corporation 1988

NOTE:

- Organic absorbents have been known to ignite when contaminated with amines in closed containers. Certain cellulosic materials used for spill cleanup such as wood chips or sawdust have shown reactivity with ethyleneamines and should be avoided.
- DO NOT touch the spill material
- Clear area of personnel and move upwind.
- Alert Emergency Responders and tell them location and nature of hazard.
- May be violently or explosively reactive.
- · Wear full body protective clothing with breathing apparatus.
- Prevent, by any means available, spillage from entering drains or water course.
- Consider evacuation.
- No smoking, naked lights or ignition sources.
- Increase ventilation.
- Stop leak if safe to do so.
- Water spray or fog may be used to disperse / absorb vapor.
- Contain spill with sand, earth or vermiculite.
- Use only spark-free shovels and explosion proof equipment.
- Collect recoverable product into labeled containers for recycling.
- Absorb remaining product with sand, earth or vermiculite.
- Collect solid residues and seal in labeled drums for disposal.
- Wash area and prevent runoff into drains.
- If contamination of drains or waterways occurs, advise emergency services.

#### **PROTECTIVE ACTIONS FOR SPILL**

#### PROTECTIVE ACTION ZONE



| From IERG (Canada/Australia) |            |
|------------------------------|------------|
| Isolation Distance           | 50 meters  |
| Downwind Protection Distance | 300 meters |

#### FOOTNOTES

1 PROTECTIVE ACTION ZONE is defined as the area in which people are at risk of harmful exposure. This zone assumes that random changes in wind direction confines the vapour plume to an area within 30 degrees on either side of the predominant wind direction, resulting in a crosswind protective action distance equal to the downwind protective action distance.

2 PROTECTIVE ACTIONS should be initiated to the extent possible, beginning with those closest to the spill and working away from the site in the downwind direction. Within the protective action zone a level of vapour concentration may exist resulting in nearly all unprotected persons becoming incapacitated and unable to take protective action and/or incurring serious or irreversible health effects.

3 INITIAL ISOLATION ZONE is determined as an area, including upwind of the incident, within which a high probability of localised wind reversal may expose nearly all persons without appropriate protection to life-threatening concentrations of the material.

4 SMALL SPILLS involve a leaking package of 200 litres (55 US gallons) or less, such as a drum (jerrican or box with inner containers). Larger packages leaking less than 200 litres and compressed gas leaking from a small cylinder are also considered "small spills". LARGE SPILLS involve many small leaking packages or a leaking package of greater than 200 litres, such as a cargo tank, portable tank or a "one-tonne" compressed gas cylinder.

5 Guide 132 is taken from the US DOT emergency response guide book.

6 IERG information is derived from CANUTEC - Transport Canada.

## ACUTE EXPOSURE GUIDELINE LEVELS (AEGL) (in ppm)

AEGL 1: The airborne concentration of a substance above which it is predicted that the general population, including susceptible individuals, could experience notable discomfort, irritation, or certain asymptomatic nonsensory effects. However, the effects are not disabling and are transient and

reversible upon cessation of exposure.

AEGL 2: The airborne concentration of a substance above which it is predicted that the general population, including susceptible individuals, could experience irreversible or other serious, long-lasting adverse health effects or an impaired ability to escape.

AEGL 3: The airborne concentration of a substance above which it is predicted that the general population, including susceptible individuals, could experience life-threatening health effects or death.

# Section 7 - HANDLING AND STORAGE

## **PROCEDURE FOR HANDLING**

- Containers, even those that have been emptied, may contain explosive vapors.
- Do NOT cut, drill, grind, weld or perform similar operations on or near containers.
- DO NOT allow clothing wet with material to stay in contact with skin
- 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, heat or ignition sources.
- When handling, DO NOT eat, drink or smoke.
- Vapor may ignite on pumping or pouring due to static electricity.
- DO NOT use plastic buckets.
- Earth and secure metal containers when dispensing or pouring product.
- Use spark-free tools when handling.
- Avoid contact with incompatible materials.
- Keep containers securely sealed.
- 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**

#### · Glass container.

Packing as supplied by manufacturer. Plastic containers may only be used if approved for flammable liquid. Check that containers are clearly labeled and free from leaks.

- For low viscosity materials (i): Drums and jerricans must be of the non-removable head type. (ii): Where a can is to be used as an inner package, the can must have a screwed enclosure.
- For materials with a viscosity of at least 2680 cSt. (23 deg. C)
- For manufactured product having a viscosity of at least 250 cSt. (23 deg. C)
- Manufactured product that requires stirring before use and having a viscosity of at least 20 cSt (23 deg. C) (i): Removable head packaging; (ii): Cans with friction closures and (iii): low pressure tubes and cartridges may be used.
- Where combination packages are used, and the inner packages are of glass, there must be sufficient inert cushioning material in contact with inner and outer packages
- In addition, where inner packagings are glass and contain liquids of packing group I there must be sufficient inert absorbent to absorb any spillage, unless the outer packaging is a close fitting molded plastic box and the substances are not incompatible with the plastic.

## STORAGE REQUIREMENTS

- •
- Store in original containers in approved flame-proof area.
- No smoking, naked lights, heat or ignition sources.
- DO NOT store in pits, depressions, basements or areas where vapors may be trapped.
- Keep containers securely sealed.
- Store away from incompatible materials in a cool, dry well ventilated area.
- Protect containers against physical damage and check regularly for leaks.

• Observe manufacturer's storing and handling recommendations.

## SAFE STORAGE WITH OTHER CLASSIFIED CHEMICALS



X: Must not be stored together

O: May be stored together with specific preventions

+: May be stored together

# Section 8 - EXPOSURE CONTROLS / PERSONAL PROTECTION

## **EXPOSURE CONTROLS**

The following materials had no OELs on our records

• hexamethyleneimine: CAS:111-49-9

## **MATERIAL DATA**

HEXAMETHYLENEIMINE:

• No exposure limits set by NOHSC or ACGIH.

Exposure limits with "skin" notation indicate that vapor and liquid may be absorbed through intact skin. Absorption by skin may readily exceed vapor inhalation exposure. Symptoms for skin absorption are the same as for inhalation. Contact with eyes and mucous membranes may also contribute to overall exposure and may also invalidate the exposure standard.

CEL TWA: 0.5 ppm (cf .DuPont OEL) OEL-STEL (Russia): 0.5 mg/m3 (skin)

## PERSONAL PROTECTION





# Consult your EHS staff for recommendations

EYE

- •
- Chemical goggles.
- Full face shield.
- Contact lenses pose a special hazard; soft lenses may absorb irritants and all lenses concentrate them.

#### HANDS/FEET

• Wear chemical protective gloves, eg. PVC.

Wear safety footwear or safety gumboots, eg. Rubber.

- When handling corrosive liquids, wear trousers or overalls outside of boots, to avoid spills entering boots.
- Suitability and durability of glove type is dependent on usage. Important factors in the selection of gloves include: such as:
- frequency and duration of contact,
- chemical resistance of glove material,
- glove thickness and
- dexterity
- Select gloves tested to a relevant standard (e.g. Europe EN 374, US F739).
- When prolonged or frequently repeated contact may occur, a glove with a protection class of 5 or higher (breakthrough time greater than 240 minutes according to EN 374) is recommended.
- When only brief contact is expected, a glove with a protection class of 3 or higher (breakthrough time greater than 60 minutes according to EN 374) is recommended.
- Contaminated gloves should be replaced.

Gloves must only be worn on clean hands. After using gloves, hands should be washed and dried thoroughly. Application of a non-perfumed moisturiser is recommended.

• Neoprene rubber gloves

#### OTHER

- •
- Overalls.
- PVC Apron.
- PVC protective suit may be required if exposure severe.

- Eyewash unit.
- Ensure there is ready access to a safety shower.
- Some plastic personal protective equipment (PPE) (e.g. gloves, aprons, overshoes) are not recommended as they may produce static electricity.
- For large scale or continuous use wear tight-weave non-static clothing (no metallic fasteners, cuffs or pockets), non sparking safety footwear.

#### RESPIRATOR

• Where the concentration of gas/particulates in the breathing zone, approaches or exceeds the OSHA Permissible Exposure Limit (or PEL), 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.

| Protection Factor | Half-Face Respirator | Full-Face Respirator | Powered Air Respirator |  |
|-------------------|----------------------|----------------------|------------------------|--|
| 10 x PEL          | AK-1 P               | -                    | AK-PAPR-1 P            |  |
| 50 x PEL          | -                    | AK-1 P               | -                      |  |
| 100 x PEL         | -                    | AK-2 P               | AK-PAPR-2 P^           |  |
|                   |                      |                      |                        |  |

^ - Full-face

Explanation of Respirator Codes:

Class 1 low to medium absorption capacity filters.

Class 2 medium absorption capacity filters.

Class 3 high absorption capacity filters.

PAPR Powered Air Purifying Respirator (positive pressure) cartridge.

Type A for use against certain organic gases and vapors.

Type AX for use against low boiling point organic compounds (less than 65°C).

Type B for use against certain inorganic gases and other acid gases and vapors.

Type E for use against sulfur dioxide and other acid gases and vapors.

Type K for use against ammonia and organic ammonia derivatives

Class P1 intended for use against mechanically generated particulates of sizes most commonly encountered in industry, e.g. asbestos, silica. Class P2 intended for use against both mechanically and thermally generated particulates, e.g. metal fume.

Class P3 intended for use against all particulates containing highly toxic materials, e.g. beryllium.

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**

• For flammable liquids and flammable gases, local exhaust ventilation or a process enclosure ventilation system may be required. Ventilation equipment should be explosion-resistant. Air contaminants generated in the workplace possess varying "escape" velocities which, in turn, determine the "capture velocities" of fresh circulating air required to effectively remove the contaminant.

| Type of Contaminant:   | Air Speed:                       |
|--|----------------------------------|
| solvent, vapors, degreasing etc., evaporating from tank (in still air).  | 0.25-0.5 m/s (50-100 f/min.)     |
| aerosols, fumes from pouring operations, intermittent container filling,<br>low speed conveyer transfers, welding, spray drift, plating acid<br>fumes, pickling (released at low velocity into zone of active<br>generation) | 0.5-1 m/s (100-200 f/min.)       |
| direct spray, spray painting in shallow booths, drum filling, conveyer<br>loading, crusher dusts, gas discharge (active generation into zone of<br>rapid air motion)   | 1-2.5 m/s (200-500 f/min.)       |
| Within each range the appropriate value depends on:  |                                  |
| Lower end of the range   | Upper end of the range           |
| 1: Room air currents minimal or favorable to capture   | 1: Disturbing room air currents  |
| 2: Contaminants of low toxicity or of nuisance value only.   | 2: Contaminants of high toxicity |
| 3: Intermittent, low production.   | 3: High production, heavy use    |
| 4: Large hood or large air mass in motion  | 4: Small hood-local control only |

Simple theory shows that air velocity falls rapidly with distance away from the opening of a simple extraction pipe. Velocity generally decreases with the square of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to distance from the contaminating source. The air velocity at the extraction fan, for example, should be a minimum of 1-2 m/s (200-400 f/min.) for extraction of solvents generated in a tank 2 meters distant from the extraction point. Other mechanical considerations, producing performance deficits within the extraction apparatus, make it essential that theoretical air velocities are multiplied by factors of 10 or more when extraction systems are installed or used.

# Section 9 - PHYSICAL AND CHEMICAL PROPERTIES

#### PHYSICAL PROPERTIES

Liquid. Mixes with water. Corrosive. Alkaline.

| State                     | Liquid            | Molecular Weight               | 99.18          |
|---------------------------|-------------------|--------------------------------|----------------|
| Melting Range (°F)        | -34.6             | Viscosity                      | Not Available  |
| Boiling Range (°F)        | 280.4 (749 mm Hg) | Solubility in water (g/L)      | Miscible       |
| Flash Point (°F)          | 64.994            | pH (1% solution)               | Not available  |
| Decomposition Temp (°F)   | Not Available     | pH (as supplied)               | Not applicable |
| Autoignition Temp (°F)    | Not available     | Vapor Pressure (mmHg)          | 7.351 @ 21.1 C |
| Upper Explosive Limit (%) | Not available     | Specific Gravity (water=1)     | 0.880          |
| Lower Explosive Limit (%) | Not available     | Relative Vapor Density (air=1) | >1             |
| Volatile Component (%vol) | Not available     | Evaporation Rate               | Not available  |

### APPEARANCE

Colourless liquid with ammonia-like odour; mixes with water.

## Section 10 - CHEMICAL STABILITY

## CONDITIONS CONTRIBUTING TO INSTABILITY

- Presence of incompatible materials.
- Product is considered stable.
- Houdde is considered stable.
- Hazardous polymerization will not occur.

### STORAGE INCOMPATIBILITY

· Hexamethyleneimine:

- reacts violently with strong oxidisers
- is incompatible with organic anhydrides, acrylates, alcohols, aldehydes, alkylene oxides, substituted allyls, cellulose nitrate, cresols, caprolactam solution, epichlorohydrin, ethylene dichloride, glycols, isocyanates, ketones, maleic anhydride, nitrates, nitromethane, phenols, vinyl acetate
- increases the explosive sensitivity if nitromethane
- reacts with nitroalkanes, forming explosive products
- attacks, aluminium, copper, lead, tin, zinc and their alloys
- Avoid strong acids.
- Avoid contact with copper, aluminium and their alloys.

Avoid reaction with oxidizing agents.

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

## Section 11 - TOXICOLOGICAL INFORMATION

hexamethyleneimine

#### **TOXICITY AND IRRITATION**

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

| TOXICITY                    | IRRITATION   |
|-----------------------------|--------------|
| Oral (rat) LD50: 410 mg/kg  | Nil Reported |
| Oral (rat) LD50: 32 mg/kg * |              |

····· (····) \_\_ ····g····g

Inhalation (mouse) LC50: 10800 mg/m³/2h

#### • For hexamethyleneimine:

Acute toxicity: Reported values for acute oral toxicity in rats with hexamethyleneimine ranged from highly toxic to slightly toxic. However, only 1 study specified the purity of the sample tested, which was 98.18%. In this study, the acute oral toxicity in rats was 1,000 mg/kg (slightly toxic). Hexamethyleneimine was moderately toxic via inhalation with an ALC in rats of 2.45 mg/L. Organs showing possible test substance related effects included the lungs, trachea, and eyes; however, histopathologic effects were difficult to interpret in the absence of a concurrent control group. Hexamethyleneimine had a dermal MLD (Minimum Lethal Dose) of 1,260-2,000 mg/kg when tested in rabbits, was corrosive to the skin and eye, and produced sensitisation reactions in 40% of mice tested using the mouse ear swelling test (MEST).

Hexamethyleneimine did not produce mortality, clinical signs of toxicity (other than some temporary discomfort at dosing), or evidence of treatment related pathological changes when administered orally to rats at a dose of 90 mg/kg, 5 times/week for 2 weeks. When administered as a single 10 mg/kg dose, hexamethyleneimine produced no effects upon plasma or interstitial fluid concentrations. At 10 mg/kg daily for 7 days, no morphological changes in testes, or abnormal changes in epididymal sperm morphology in male rats were observed.

Repeat dose, reproductive, developmental toxicity: Except for mucosal thickening of glandular/forestomach tissue observed at necropsy (without correlation to microscopic tissue damage), no other treatment related effects were observed up to the highest dosage tested (50 mg/kg/day hexamethyleneimine (by gavage) during a Combined Repeated Dose Toxicity and Reproductive/Developmental Toxicity Screen study (OECD 422). Additionally, no effects were observed on gonadal tissue or on reproductive parameters including mating, parturition or lactation in parental animals. No evidence of developmental or fetal effects were seen in offspring even at the highest dosage (50 mg/kg/day) tested

Genotoxicity: Hexamethyleneimine was negative in Salmonella typhimurium and Escherichia coli when tested in the in vitro bacterial reverse mutation assay. No clastogenic effects were observed when hexamethyleneimine was tested in an in vitro chromosomal aberration study

(OECD 473) in human lymphocytes, with and without incorporation of a mammalian metabolic activation system.

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.

## Section 12 - ECOLOGICAL INFORMATION

1.31

2 17

Refer to data for ingredients, which follows: HEXAMETHYLENEIMINE:

• BOD5:

· COD:

• For hexamethyleneimine:

Environmental fate:

If released to air, a vapor pressure of 8.09 mm Hg at 25 C indicates hexamethyleneimine will exist solely in the vapour phase in the ambient atmosphere. Vapour-phase hexamethyleneimine will be degraded in the atmosphere by reaction with photochemically-produced hydroxyl radicals; the half-life for this reaction in air is estimated to be 4.3 hours. If released into water, hexamethyleneimine is expected to have limited adsorption to suspended solids and sediment in water based upon the estimated Koc of 20. A pKa of 11.07 indicates hexamethyleneimine will exist almost entirely in the protonated form in aqueous environments, and is not expected to volatilise from water surfaces. The maximum reported biodegradability for hexamethyleneimine was 1.19% over 28 days. In addition, hexamethyleneimine was toxic to the microbial inoculum at a test concentration of 2 mg active substance/L. An estimated BCF of 3.9 suggests the potential for bioconcentration in aquatic organisms is low . Fugacity model prediction indicates that hexamethyleneimine will partition mainly to the soil and water, with virtually none going to the air or sediment.

Ecotoxicity:

Aquatic toxicity studies conducted with hexamethyleneimine on fish, invertebrates, and algae according to current OECD TG (203, 202, and 201, respectively) indicate low to moderate acute aquatic toxicity. LC50/EC50 values were determined to be >100 mg/L for rainbow trout and Daphnia and 88 mg/L (growth index) for green algae.

• DO NOT discharge into sewer or waterways.

#### Ecotoxicity

Ingredient hexamethyleneimine Persistence: Water/Soil Persistence: Air LOW

Bioaccumulation LOW Mobility MED

# Section 13 - DISPOSAL CONSIDERATIONS

#### **US EPA Waste Number & Descriptions**

A. General Product Information

Ignitability characteristic: use EPA hazardous waste number D001 (waste code I)

Corrosivity characteristic: use EPA hazardous waste number D002 (waste code C)

## **Disposal Instructions**

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

• Puncture containers to prevent re-use and bury at an authorized landfill. Legislation addressing waste disposal requirements may differ by country, state and/ or territory. Each user must refer to laws operating in

their area. In some areas, certain wastes must be tracked.

A Hierarchy of Controls seems to be common - the user should investigate:

- Reduction
- Reuse
- Recycling
- Disposal (if all else fails)

This material may be recycled if unused, or if it has not been contaminated so as to make it unsuitable for its intended use. If it has been contaminated, it may be possible to reclaim the product by filtration, distillation or some other means. Shelf life considerations should also be applied in making decisions of this type. Note that properties of a material may change in use, and recycling or reuse may not always be appropriate.

DO NOT allow wash water from cleaning equipment to enter drains. Collect all wash water for treatment before disposal.

Recycle wherever possible.

- Consult manufacturer for recycling options or consult Waste Management Authority for disposal if no suitable treatment or disposal facility can be identified.
- Treat and neutralize at an approved treatment plant.
- Treatment should involve: Neutralization with suitable dilute acid followed by: Burial in a licensed land-fill or Incineration in a licensed apparatus
- Decontaminate empty containers. Observe all label safeguards until containers are cleaned and destroyed.

# **Section 14 - TRANSPORTATION INFORMATION**



| DOT:   |                            |  |              |  |
|--|----------------------------|--|--------------|--|
| Symbols:   | None                       | Hazard class or Division:                      | 3            |  |
| Identification Numbers:                                    | UN2493                     | PG:  | П            |  |
| Label Codes:   | 3, 8                       | Special provisions:                            | IB2, T7, TP1 |  |
| Packaging: Exceptions:                                     | 150                        | Packaging: Non-bulk:                           | 202          |  |
| Packaging: Exceptions:                                     | 150                        | Quantity limitations: Passenger aircraft/rail: | 1 L          |  |
| Quantity Limitations: Cargo<br>aircraft only:              | 5 L                        | Vessel stowage: Location:                      | В            |  |
| Vessel stowage: Other:                                     | 40                         |  |              |  |
| Hazardous materials descriptions a<br>Hexamethyleneimine   | and proper shipping names: |  |              |  |
| Air Transport IATA:  |                            |  |              |  |
| ICAO/IATA Class:   | 3 (8)                      | ICAO/IATA Subrisk:                             | None         |  |
| UN/ID Number:  | 2493                       | Packing Group:                                 | II           |  |
| Special provisions:  | None                       |  |              |  |
| Shipping Name: HEXAMETHYLENEIMINE Maritime Transport IMDG: |                            |  |              |  |
| IMDG Class:  | 3                          | IMDG Subrisk:                                  | 8            |  |
| UN Number:   | 2493                       | Packing Group:                                 | II           |  |
| EMS Number:  | F-E,S-C                    | Special provisions:                            | None         |  |
| Limited Quantities:  | 1 L                        |  |              |  |
|  |                            |  |              |  |

Shipping Name: HEXAMETHYLENEIMINE

## Section 15 - REGULATORY INFORMATION

#### hexamethyleneimine (CAS: 111-49-9) is found on the following regulatory lists;

"Canada Domestic Substances List (DSL)","Canada Ingredient Disclosure List (SOR/88-64)","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","OECD Representative List of High Production Volume (HPV) Chemicals","US - Massachusetts Oil & Hazardous Material List","US DOE Temporary Emergency Exposure Limits (TEELs)","US EPA High Production Volume Program Chemical List","US Harmonized Tariff Schedule -Intermediate Chemicals for Dyes Appendix","US Toxic Substances Control Act (TSCA) - Inventory"

## **Section 16 - OTHER INFORMATION**

#### LIMITED EVIDENCE

- · Cumulative effects may result following exposure\*.
- · Limited evidence of a carcinogenic effect\*.

\* (limited evidence).

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• 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.

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