

# Produktinformation



Forschungsprodukte & Biochemikalien
Zellkultur & Verbrauchsmaterial
Diagnostik & molekulare Diagnostik
Laborgeräte & Service

Weitere Information auf den folgenden Seiten! See the following pages for more information!



Lieferung & Zahlungsart siehe unsere Liefer- und Versandbedingungen

### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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### **Section 2 - HAZARDS IDENTIFICATION**

### **CANADIAN WHMIS SYMBOLS**



EMERGENCY OVERVIEW RISK Harmful if swallowed. Irritating to eyes, respiratory system and skin.

POTENTIAL HEALTH EFFECTS

sc-228444



Material Safety Data Sheet

Hazard Alert Code Key: EXTREME HIGH MODERATE LOW

### 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.

Poisonings rarely occur after oral administration of manganese salts because they are poorly absorbed from the gut.

■ Ingestion of 2,4-pentanedione (acetylacetone) may produce irritation of the mouth, oesophagus and stomach producing abdominal discomfort, nausea, vomiting diarrhoea, dizziness, malaise and fainting.

At sufficiently high doses the material may be neurotoxic(i.e. poisonous to the nervous system).

#### EYE

This material can cause eye irritation and damage in some persons.

• Exposure to 2,4-pentadione may produce excessive redness of the eyes and swelling of the conjunctivae; blinking and tearing may occur. Corneal damage is unlikely.

#### SKIN

• The material may cause mild but significant inflammation of the skin either following direct contact or after a delay of some time. Repeated exposure can cause contact dermatitis which is characterized by redness, swelling and blistering.

• Open cuts, abraded or irritated skin should not be exposed to this material.

■ 2,4-Pentadione may produce contact dermatitis or urticaria. Prolonged contact with 2,4-pentanedione may produce severe discomfort or pain, redness and swelling and corrosion, ulceration and development of fissures. The inflamed area may show bleeding.

• 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.

### INHALED

The material can cause respiratory irritation in some persons. The body's response to such irritation can cause further lung damage.

Inhalation of dusts, generated by the material during the course of normal handling, may be damaging to the health of the individual.

Persons with impaired respiratory function, airway diseases and conditions such as emphysema or chronic bronchitis, may incur further disability if excessive concentrations of particulate are inhaled.

• Manganese fume is toxic and produces nervous system effects characterized by tiredness. Acute poisoning is rare although acute inflammation of the lungs may occur. A chemical pneumonia may also result from frequent exposure. Inhalation of freshly formed metal oxide particles sized below 1.5 microns and generally between 0.02 to 0.05 microns may result in "metal fume fever". Symptoms may be delayed for up to 12 hours and begin with the sudden onset of thirst, and a sweet, metallic or foul taste in the mouth. Other symptoms include upper respiratory tract irritation accompanied by coughing and a dryness of the mucous membranes, lassitude and a generalized feeling of malaise. Mild to severe headache, nausea, occasional vomiting, fever or chills, exaggerated mental activity, profuse sweating, diarrhea, excessive urination and prostration may also occur. Tolerance to the fumes develops rapidly, but is quickly lost. All symptoms usually subside within 24-36 hours following removal from exposure.

At sufficiently high doses the material may be neurotoxic(i.e. poisonous to the nervous system).

■ High doses of 2,4-pentanedione produced dyspnae, severe, central nervous system depression and death in experimental animals. Similar effects were produced at lower repeated doses although some animals survive and develop a central nervous system disorder characterised by irreversible cerebellar

syndrome. Thymic necrosis and atrophy accompany the central nervous system damage. [Patty's].

Inhalation of vapours may produce narcosis.

### CHRONIC HEALTH EFFECTS

• Long-term exposure to respiratory irritants may result in disease of the airways involving difficult breathing and related systemic problems. Limited evidence suggests that repeated or long-term occupational exposure may produce cumulative health effects involving organs or biochemical systems.

There is limited evidence that, skin contact with this product is more likely to cause a sensitization reaction in some persons compared to the general population.

Long term exposure to high dust concentrations may cause changes in lung function i.e. pneumoconiosis; caused by particles less than 0.5 micron penetrating and remaining in the lung. Prime symptom is breathlessness; lung shadows show on X-ray.

Manganese is an essential trace element. Chronic exposure to low levels of manganese can include a mask-like facial expression, spastic gait, tremors, slurred speech, disordered muscle tone, fatigue, anorexia, loss of strength and energy, apathy and poor concentration.

Repeated overexposure to 200 ppm 2,4-pentanedione vapor may result in inflammation of the nasal mucosa. Higher concentrations may produce central nervous system effects, and immune system and bone marrow deficits. Recurrent exposure to high concentrations of the 2,4-pentadione vapor (~650 ppm) produces lethal degenerative lesions in the central nervous system and thymus. Exposure in pregnancy can damage to fetus.

In use the substance may decompose to produce 2,4-pentanedione a powerful neurotoxin.

### Section 3 - COMPOSITION / INFORMATION ON INGREDIENTS

### sc-228444



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Hazard Alert Code Ke	у:	EXTREME		HIGH	MODERATE			LOW	
HAZARD RATINGS									
		Min	Max						
Flammability:	1								
Toxicity:	2								
Body Contact:	2			Min/Nil=0 Low=1					
Reactivity:	1			Moderate=2					
Chronic:	2			High=3 Extreme=4					
NAME						CAS RN		%	
manganese(III) acetylac	etonate	9				14284-89	-0	>98	
NOTE: In use may evolv	/e								
2,4-pentanedione						123-54-6			

### **Section 4 - FIRST AID MEASURES**

### SWALLOWED

- I
- IF SWALLOWED, REFER FOR MEDICAL ATTENTION, WHERE POSSIBLE, WITHOUT DELAY.
- Where Medical attention is not immediately available or where the patient is more than 15 minutes from a hospital or unless instructed otherwise:
- For advice, contact a Poisons Information Center or a doctor.
- Urgent hospital treatment is likely to be needed.
- If conscious, give water to drink.
- INDUCE vomiting with fingers down the back of the throat, ONLY IF CONSCIOUS. Lean patient forward or place on left side (head-down position, if possible) to maintain open airway and prevent aspiration.

NOTE: Wear a protective glove when inducing vomiting by mechanical means.

- In the mean time, qualified first-aid personnel should treat the patient following observation and employing supportive measures as indicated by the patient's condition.
- If the services of a medical officer or medical doctor are readily available, the patient should be placed in his/her care and a copy of the MSDS should be provided. Further action will be the responsibility of the medical specialist.
- If medical attention is not available on the worksite or surroundings send the patient to a hospital together with a copy of the MSDS.

#### 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.
- 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.
- Lay patient down. Keep warm and rested.
- Prostheses such as false teeth, which may block airway, should be removed, where possible, prior to initiating first aid procedures.
- Apply artificial respiration if not breathing, preferably with a demand valve resuscitator, bag-valve mask device, or pocket mask as trained. Perform CPR if necessary.
- Transport to hospital, or doctor, without delay.



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Hazard Alert Code Key: EXTREME HIGH MODERATE LOW	Hazard Alert Code Key:	EXTREME	HIGH	MODERATE	LOW
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### NOTES TO PHYSICIAN

### Treat symptomatically.

Both dermal and oral toxicity of manganese salts is low because of limited solubility of manganese. No known permanent pulmonary sequelae develop after acute manganese exposure. Treatment is supportive. [Ellenhorn and Barceloux: Medical Toxicology] In clinical trials with miners exposed to manganese-containing dusts, L-dopa relieved extrapyramidal symptoms of both hypokinetic and dystonic patients. For short periods of time symptoms could also be controlled with scopolamine and amphetamine. BAL and calcium EDTA prove ineffective. [Gosselin et al: Clinical Toxicology of Commercial Products.].

Section 5 - FIRE FIGHTING MEASURES					
Vapour Pressure (mmHG):	Not available				
Upper Explosive Limit (%):	Not available				
Specific Gravity (water=1):	Not available				
Lower Explosive Limit (%):	Not available				

### **EXTINGUISHING MEDIA**

- Foam.
- ٠ 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.
- 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.
- 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**

- Combustible solid which burns but propagates flame with difficulty.
- Avoid generating dust, particularly clouds of dust in a confined or unventilated space as dusts may form an explosive mixture with air, and • any source of ignition, i.e. flame or spark, will cause fire or explosion. Dust clouds generated by the fine grinding of the solid are a particular hazard; accumulations of fine dust may burn rapidly and fiercely if ignited.
- Dry dust can be charged electrostatically by turbulence, pneumatic transport, pouring, in exhaust ducts and during transport.
- Build-up of electrostatic charge may be prevented by bonding and grounding.
- Powder handling equipment such as dust collectors, dryers and mills may require additional protection measures such as explosion venting.

Combustion products include: carbon monoxide (CO), carbon dioxide (CO2), other pyrolysis products typical of burning organic material. May emit poisonous fumes.

#### May emit corrosive fumes.

#### FIRE INCOMPATIBILITY

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

### PERSONAL PROTECTION

Glasses: Chemical goggles. Gloves: Respirator: Type A-P Filter of sufficient capacity

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LOW Hazard Alert Code Key: EXTREME HIGH MODERATE Section 6 - ACCIDENTAL RELEASE MEASURES MINOR SPILLS • Remove all ignition sources. Clean up all spills immediately. Avoid contact with skin and eyes. Control personal contact by using protective equipment. Use dry clean up procedures and avoid generating dust. Place in a suitable, labelled container for waste disposal. MAJOR SPILLS Moderate hazard. CAUTION: Advise personnel in area. • Alert Emergency Responders and tell them location and nature of hazard. Control personal contact by wearing protective clothing. • Prevent, by any means available, spillage from entering drains or water courses. Recover product wherever possible. IF DRY: Use dry clean up procedures and avoid generating dust. Collect residues and place in sealed plastic bags or other containers for disposal. IF WET: Vacuum/shovel up and place in labelled containers for disposal. ALWAYS: Wash area down with large amounts of water and prevent runoff into drains. If contamination of drains or waterways occurs, advise emergency services. 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

- Avoid all personal contact, including inhalation.
- Wear protective clothing when risk of exposure occurs.
- Use in a well-ventilated area.

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- Prevent concentration in hollows and sumps.
- DO NOT enter confined spaces until atmosphere has been checked.
- DO NOT allow material to contact humans, exposed food or food utensils.
- 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.
- Launder contaminated clothing before re-use.
- 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 are maintained.
 Empty containers may contain residual dust which has the potential to accumulate following settling. Such dusts may explode in the presence

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Hazard Alert Code Key:	EXTREME	HIGH	MODERATE	LOW
of an appropriate ignition sour • Do NOT cut, drill, grind or v				

• In addition ensure such activity is not performed near full, partially empty or empty containers without appropriate workplace safety authorisation or permit.

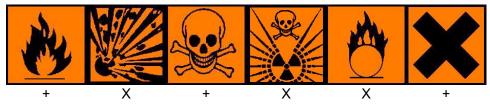
### **RECOMMENDED STORAGE METHODS**

- Glass container.
- Polyethylene or polypropylene container.
- Check all containers are clearly labelled and free from leaks.

### STORAGE REQUIREMENTS

- Store in original containers.
- Keep containers securely sealed.
- Store in a cool, dry, well-ventilated area.
- Store away from incompatible materials and foodstuff containers.
- Protect containers against physical damage and check regularly for leaks.
- Observe manufacturer's 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**

Source	Material	TWA ppm	TWA mg/m³	STEL ppm	STEL mg/m³	Peak ppm	Peak mg/m³	TWA F/CC	Notes
US - California Permissible Exposure Limits for Chemical Contaminants	manganese(III) acetylacetonate (Manganese and compounds, as Mn)		0.2						
US - Idaho - Limits for Air Contaminants	manganese(III) acetylacetonate (Manganese compounds (as Mn))						5		
US NIOSH Recommended Exposure Limits (RELs)	manganese(III) acetylacetonate (Manganese compounds and fume (as Mn))		1		3				
US - Minnesota Permissible Exposure Limits (PELs)	manganese(III) acetylacetonate (Manganese compounds (as Mn))						5		
US - Minnesota Permissible Exposure Limits (PELs)	manganese(III) acetylacetonate (Manganese fume (as Mn))		1		3				
US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air Contaminants	manganese(III) acetylacetonate (Manganese fume (as Mn))		(C)5						

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Hazard Alert Code Key:	XTREME	HIGH	1		MODER	ATE		LOW
US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants	manganese(III) ac (Manganese comp Mn))						5	
US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants	manganese(III) ac (Manganese fume			1	:	3		
US - Idaho - Limits for Air Contaminants	manganese(III) ac (Manganese fume	•					5	
US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	manganese(III) ac (Manganese com Mn))						5	
US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air Contaminants	manganese(III) ac (Manganese com Mn))			(C)5				
US - Hawaii Air Contaminant Limits	manganese(III) ac (Manganese fume	•		1	;	3		
US - Alaska Limits for Air Contaminants	manganese(III) ac (Manganese Com Mn))						5	
US - Michigan Exposure Limits for Air Contaminants	manganese(III) ac (Manganese, Con Mn))						5	
Canada - Yukon Permissible Concentrations for Airborne Contaminant Substances	manganese(III) ac (Manganese and Mn))		-	5				
US - Washington Permissible exposure limits of air contaminants	manganese(III) ac (Manganese and Mn))						5	
US - Hawaii Air Contaminant Limits	manganese(III) ac (Manganese comp Mn))	•					5	
Canada - Quebec Permissible Exposure Values for Airborne Contaminants (English)	manganese(III) ac (Manganese (as M compounds)			5				
US OSHA Permissible Exposure Levels (PELs) - Table Z1	manganese(III) ac (Manganese com Mn))	•					5	
US OSHA Permissible Exposure Levels (PELs) - Table Z1	manganese(III) ac (Manganese fume						5	
US - Oregon Permissible Exposure Limits (Z1)	manganese(III) ac (Manganese Com Mn))						5	
Canada - Northwest Territories Occupational Exposure Limits (English)	manganese(III) ac (Manganese & co Mn))						5	
Canada - Northwest Territories Occupational Exposure Limits (English)	manganese(III) ac (Manganese fume	•		1	:	3		
Canada - Alberta Occupational Exposure Limits	2,4-pentanedione fuels, as total hyd vapour)			200				

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The Power to Question

Hazard Alert Code Key:	EXTREME	HIGH	I	MODERATE	LOW	
Canada - Ontario Occupational Exposure Limits	2,4-pentanedione total hydrocarbor aerosol)	•	100			Skin
Canada - Saskatchewan Occupational Health and Safety Regulations - Contamination Limits	2,4-pentanedione total hydrocarbor	•	100	150		Skin
Canada - Alberta Occupational Exposure Limits	2,4-pentanedione total hydrocarbor	( ,	100			
Canada - British Columbia Occupational Exposure Limits	2,4-pentanedione total hydrocarbor	· · · ·	100 (V)			Skin
EMERGENCY EXPOSURE LIMITS						
Material	Revis	ed IDLH Value (mg/m3)		Revised IDLH	Value (ppm)	
manganese(III) acetylacetonate	500					

### MATERIAL DATA

2,4-PENTANEDIONE:

MANGANESE(III) ACETYLACETONATE:

■ CEL TWA for 2,4-pentadione: 20 ppm, 82 mg/m3 [cf recommended OEL, Union Carbide]

The disagreeable odour associated with relatively low concentrations of 2,4-pentanedione, in the atmosphere, may produce nausea. Exposure to about 1000 ppm for 30 minutes may result in the inhalation of harmful and potentially fatal amounts of the material.

Odour Safety Factor(OSF) OSF=21 (2,4-pentanedione.

MANGANESE(III) ACETYLACETONATE:

• Ceiling values were recommended for manganese and compounds in earlier publications. As manganese is a chronic toxin a TWA is considered more appropriate. Because workers exposed to fume exhibited manganism at air-borne concentrations below those that effect workers exposed to dust a lower value has been proposed to provide an extra margin of safety. This value is still above that experienced by two workers exposed to manganese fume in the course of one study.

A number of studies have shown that susceptibility to the effects of manganese at or about 1 - 5 mg/m3 (TWA) can lead to clinical manifestations of manganism or more commonly to the development of indicators of sub- clinical manganism (e.g. hand tremor, exaggerated reflexes, short-term memory deficits, poor psychomotor performance). Long term exposure at, or below the recommended ES TWA should provide protection for those individuals susceptible to neurological effects of prolonged exposure.

2,4-PENTANEDIONE:

• 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.

### PERSONAL PROTECTION



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				The Power to Question
Hazard Alert Code Key:	EXTREME	HIGH	MODERATE	LOW
Consult your EHS staff for recon	nmendations			
EYE				
<ul> <li>Safety glasses with side shie</li> <li>Chemical goggles.</li> </ul>	IOS.			
<ul> <li>Contact lenses pose a specia</li> </ul>	al hazard: soft lenses may a	absorb irritants and all len	ses concentrate them DO N	OT wear contact lenses
HANDS/FEET	a nazara, son ienses may a			
■ NOTE: The material may pro	oduce skin sensitization in	predisposed individuals	Care must be taken when	removing gloves and other
protective equipment, to avoid a				ionioring gioree and earer
Suitability and durability of glove	type is dependent on usag	e. Important factors in the	selection of gloves include:	such as:
<ul> <li>frequency and duration of co</li> </ul>				
chemical resistance of glove	material,			
<ul> <li>glove thickness and</li> <li>devtority</li> </ul>				
<ul> <li>dexterity</li> <li>Select gloves tested to a relevant</li> </ul>	at standard (e.g. Europe EN	374 LIS E730)		
<ul> <li>When prolonged or frequentl</li> </ul>		. ,	tion class of 5 or higher (bre	akthrough time greater than
240 minutes according to EN	· · ·	cul, u gioro mar u protoc		ananough anto groator anan
• When only brief contact is ex	xpected, a glove with a prot	tection class of 3 or highe	er (breakthrough time greate	r than 60 minutes according
to EN 374) is recommended.				
Contaminated gloves should				
Gloves must only be worn on cle moisturiser is recommended.	ean hands. After using glov	es, hands should be was	hed and dried thoroughly. Ap	oplication of a non-perfumed
Experience indicates that the fol	lowing polymers are suitabl	e as glove materials for p	rotection against undissolve	d dry solids where abrasive
particles are not present.		o do gio io materialo ioi p		
<ul> <li>polychloroprene</li> </ul>				
<ul> <li>nitrile rubber</li> </ul>				
<ul> <li>butyl rubber</li> </ul>				
fluorocaoutchouc				
<ul> <li>polyvinyl chloride</li> <li>Gloves should be examined for y</li> </ul>	waar and/ or degradation of	notantly		
OTHER		nistantiy.		
-				

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

### RESPIRATOR

Protection Factor	Half-Face Respirator	Full-Face Respirator	Powered Air Respirator
10 x PEL	A P1	-	A PAPR-P1
	Air-line*	-	-
50 x PEL	Air-line**	A P2 A	PAPR-P2
100 x PEL	-	A P3	-
		Air-line*	-
100+ x PEL	-	Air-line**	A PAPR-P3

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HIGH



LOW

MODERATE

### Material Safety Data Sheet

Hazard Alert Code Key:

EXTREME

<ul> <li>* - Negative pressure demand ** - Continuous flow</li> <li>Explanation of Respirator Codes:</li> <li>Class 1 low to medium absorption capacity filters.</li> <li>Class 2 medium absorption capacity filters.</li> <li>Class 3 high absorption capacity filters.</li> <li>PAPR Powered Air Purifying Respirator (positive pressure) cartridge.</li> <li>Type A for use against certain organic gases and vapors.</li> <li>Type A for use against certain inorganic gases and other acid gases and vapors.</li> <li>Type B for use against certain inorganic gases and other acid gases and vapors.</li> <li>Type K for use against sulfur dioxide and other acid gases and vapors.</li> <li>Type K for use against sulfur dioxide and other acid gases and vapors.</li> <li>Type K for use against monia and organic ammonia derivatives</li> <li>Class P2 intended for use against both mechanically generated particulates of sizes most commonly encountered in industry, e.g. asbestos, silica.</li> <li>Class P3 intended for use against all particulates containing highly toxic materials, e.g. metal fume.</li> <li>Class P3 intended for use against all particulates containing highly toxic materials, e.g. beryllium.</li> <li>The local concentration of material, quantity and conditions of use determine the type of personal protective equipment required.</li> <li>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.</li> </ul>							
ENGINEERING CONTROLS							
<ul> <li>Local exhaust ventilation is required where solids are handled as powders or crystals; even when particulates are relatively large, certain proportion will be powdered by mutual friction.</li> <li>Exhaust ventilation should be designed to prevent accumulation and recirculation of particulates in the workplace.</li> <li>If in spite of local exhaust an adverse concentration of the substance in air could occur, respiratory protection should be considered. Such protection might consist of: <ul> <li>(a): particle dust respirators, if necessary, combined with an absorption cartridge;</li> <li>(b): filter respirators with absorption cartridge or canister of the right type;</li> <li>(c): fresh-air hoods or masks</li> <li>Build-up of electrostatic charge on the dust particle, may be prevented by bonding and grounding.</li> <li>Powder handling equipment such as dust collectors, dryers and mills may require additional protection measures such as explosion venting.</li> </ul> </li> <li>Air contaminants generated in the workplace possess varying "escape" velocities which, in turn, determine the "capture velocities" of free circulating air required to efficiently remove the contaminant.</li> </ul>							
Type of Contaminant: Air Speed:							
direct spray, spray painting in shallow booths, drum filling, conveyer loading, crusher dusts, gas discharge (active generation into zone of 1-2.5 m/s (200-500 f/min.) rapid air motion)							
grinding, abrasive blasting, tumbling, high speed wheel generated dusts (released at high initial velocity into zone of very high rapid air 2.5-10 m/s (500-2000 f/min.) motion).							
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							

4: Large hood or large air mass in motion

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 4-10 m/s (800-2000 f/min) for extraction of crusher dusts generated 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.

4: Small hood-local control only

### **Section 9 - PHYSICAL AND CHEMICAL PROPERTIES**

### sc-228444



### Material Safety Data Sheet

Hazard Alert Code Key:	EXTREME	HIGH	MODERATE	LOW
PHYSICAL PROPERTIES				
Solid. Does not mix with water.				
State	Divided solid	Mole	ecular Weight	352.27
Melting Range (°F)	320 (decompos	es) Visco	osity	Not Applicable
Boiling Range (°F)	Not applicable	Solu	bility in water (g/L)	Partly miscible
Flash Point (°F)	Not Available	pH (	1% solution)	Not applicable
Decomposition Temp (°F)	320	pH (a	as supplied)	Not applicable
Autoignition Temp (°F)	Not available	Vapo	our Pressure (mmHG)	Not available
Upper Explosive Limit (%)	Not available	Spec	cific Gravity (water=1)	Not available
Lower Explosive Limit (%)	Not available	Rela	tive Vapor Density (air=1)	>1
Volatile Component (%vol)	Not available	Evap	poration Rate	Not available

### APPEARANCE

Solid; does not mix well with water. In use the substance may decompose to produce 2,4-pentanedione a powerful neurotoxin

### **Section 10 - CHEMICAL STABILITY**

### CONDITIONS CONTRIBUTING TO INSTABILITY

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

### STORAGE INCOMPATIBILITY

Avoid reaction with oxidizing agents.

- For 2,4-pentanedione:
- Segregate from halogens.
- Store away from steel, nickel, zinc, galvanized iron, tinned iron, copper and copper alloys.

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

### **Section 11 - TOXICOLOGICAL INFORMATION**

manganese(III) acetylacetonate

#### TOXICITY AND IRRITATION

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

• 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 (often particulate in nature) and is completely reversible after exposure ceases. The disorder is characterised by dyspnea, cough and mucus production.

The material may cause skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of vesicles, scaling and thickening of the skin.

No significant acute toxicological data identified in literature search.

#### SKIN

2,4-pentanedione Canada - Ontario Occupational Exposure Limits - Skin

Notes

Skin

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2,4-pentanedione US AIHA	Notes	Skin			
2,4-pentanedione Canada	Notes	Skin			
2,4-pentanedione Canada	Notation	Skin			
2,4-pentanedione Canada	- Alberta Occupational Expo	sure Limits - Skin		Substance Interaction	1

### Section 12 - ECOLOGICAL INFORMATION

Refer to data for ingredients, which follows:

2,4-PENTANEDIONE:

MANGANESE(III) ACETYLACETONATE:

DO NOT discharge into sewer or waterways.

■ for 2,4-pentanedione:

log Kow : 1.9-2.25

BOD 5 if unstated: 5.60%

Harmful to aquatic life/ birdlife. Environmental fate:

Biodegradation of 2,4-pentadione occurs in the soil. Leaching into ground water is expected to occur. Biodegradation in water is expected to proceed at a moderate rate. No significant bioaccumulation is expected to occur.

In air 2,4-pentadione undergoes photochemical degradation to hydroxyl radicals. The material is expected to be removed from air by wet deposition with a half-life between 10 and 30 days.

MANGANESE(III) ACETYLACETONATE:

For manganese and its compounds:

Environmental fate:

It has been established that while lower organisms (e.g., plankton, aquatic plants, and some fish) can significantly bioconcentrate manganese, higher organisms (including humans) tend to maintain manganese homeostasis. This indicates that the potential for biomagnification of manganese from lower trophic levels to higher ones is low.

There were two mechanisms involved in explaining the retention of manganese and other metals in the environment by soil. First, through cation exchange reactions, manganese ions and the charged surface of soil particles form manganese oxides, hydroxides, and oxyhydroxides which in turn form absorption sites for other metals. Secondly, manganese can be adsorbed to other oxides, hydroxides, and oxyhydroxides through ligand exchange reactions. When the soil solution becomes saturated, these manganese oxides, hydroxides, and oxyhydroxides can precipitate into a new mineral phase and act as a new surface to which other substances can absorb. The tendency of soluble manganese compounds to adsorb to soils and sediments depends mainly on the cation exchange capacity and the organic composition of the soil. The soil adsorption constants (the ratio of the concentration in soil to the concentration in water) for Mn(II) span five orders of magnitude, ranging from 0.2 to 10,000 mL/g, increasing as a function of the organic content and the ion exchange capacity of the soil; thus, adsorption may be highly variable. In some cases, adsorption of manganese to soils may not be a readily reversible process. At low concentrations, manganese may be "fixed" by clays and will not be released into solution readily. At higher concentrations, manganese environments resulted in the mobilization of manganese from the bottom sediments. The metals in the effluent may have been preferentially adsorbed resulting in the release of manganese. The oxidation state of manganese in soil and sediments may be altered by microbial activity; oxidation may lead to the precipitation of manganese. Bacteria and microflora can increase the mobility of manganese.

The transport and partitioning of manganese in water is controlled by the solubility of the specific chemical form present, which in turn is determined by pH, Eh (oxidation-reduction potential), and the characteristics of the available anions. The metal may exist in water in any of four oxidation states.

Manganese(II) predominates in most waters (pH 4-7) but may become oxidized at a pH >8 or 9. The principal anion associated with Mn(II) in water is usually carbonate (CO3.2), and the concentration of manganese is limited by the relatively low solubility (65 mg/L) of MnCO3. In relatively oxidized water, the solubility of Mn(II) may be controlled by manganese oxide equilibria, with manganese being converted to the Mn(II) or Mn(IV) oxidation states. In extremely reduced water, the fate of manganese tends to be controlled by formation of a poorly soluble sulfide. Manganese in water may undergo oxidation at high pH or Eh and is also subject to microbial activity. For example, Mn(II) in a lake was oxidized during the summer months, but this was inhibited by a microbial poison, indicating that the oxidation was mediated by bacteria . The microbial metabolism of manganese is presumed to be a function of pH, temperature, and other factors.

Manganese in water may be significantly bioconcentrated at lower trophic levels. A bioconcentration factor (BCF) relates the concentration of a chemical in plant and animal tissues to the concentration of the chemical in the water in which they live. The BCF of manganese was estimated as 2,500 - 6,300 for phytoplankton, 300 -5,500 for marine algae, 80 - 830 for intertidal mussels, and 35 - 930 for coastal fish. Similarly, the BCF of manganese was estimated to be 10,00 -20,000 for marine and freshwater plants, 10,000 - 40,000 for invertebrates, and 10 - 600 for fish. In general, these data indicate that lower organisms such as algae have larger BCFs than higher organisms. In order to protect consumers from the risk of manganese bioaccumulation in marine mollusks, the U.S. EPA has set a criterion for manganese at 0.1 mg/L for marine waters.

Elemental manganese and inorganic manganese compounds have negligible vapor pressures but may exist in air as suspended particulate matter derived from industrial emissions or the erosion of soils. Manganese-containing particles are mainly removed from the atmosphere by

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gravitational settling, with large particles tending to fall out faster than small particles. The half-life of airborne particles is usually on the order of days, depending on the size of the particle and atmospheric conditions. Some removal by washout mechanisms such as rain may also occur, although it is of minor significance in comparison to dry deposition. 2,4-PENTANEDIONE:						
■ log Pow (Verschueren 1983):					1.9/2.25	
Algae IC50 (72hr.) (mg/l):					2.7- 8.5	
■ BCF<100:					0.7, 0.75	
Water solubility (g/l):					166000	
■ log Kow (Prager 1995):					0.14	
Iog Pow (Verschueren 1983):					1.90/2.25	
BOD5:					5.60%	
<ul><li>BOD20:</li><li>Harmful to aquatic organisms.</li></ul>					69.6	
<b>Ecotoxicity</b> Ingredient 2,4-pentanedione	Persistence: Water LOW	Soil P	ersistence: Air	Bioaccumulation LOW	n	Mobility HIGH

### Section 13 - DISPOSAL CONSIDERATIONS

#### **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. 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.

For small quantities:

- Dissolve the material (in water or acid solution as appropriate) or convert it to a water soluble state with appropriate oxidizing agent.
- Precipitate as the sulfide, adjusting the pH to neutral to complete the precipitation.
- Filter off sulfide solids for recovery or disposal to approved land-fill.
- Destroy excess sulfide in solution with, for example, sodium hypochlorite, neutralize, and flush to sewer (subject to local regulation).
- 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.
- Dispose of by: Burial in a licensed land-fill or Incineration in a licensed apparatus (after admixture with suitable combustible material)
- Decontaminate empty containers. Observe all label safeguards until containers are cleaned and destroyed.

### **Section 14 - TRANSPORTATION INFORMATION**

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

### **Section 15 - REGULATORY INFORMATION**

manganese(III) acetylacetonate (CAS: 14284-89-0) is found on the following regulatory lists; "Canada Non-Domestic Substances List (NDSL)", "US Toxic Substances Control Act (TSCA) - Inventory"

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Regulations for ingredients

#### 2,4-pentanedione (CAS: 123-54-6) is found on the following regulatory lists;

"Canada Domestic Substances List (DSL)", "Canada Ingredient Disclosure List (SOR/88-64)", "International Council of Chemical Associations (ICCA) - High Production Volume List", "OECD Representative List of High Production Volume (HPV) Chemicals", "US - Massachusetts Oil & Hazardous Material List", "US - New Jersey Right to Know Hazardous Substances", "US DOE Temporary Emergency Exposure Limits (TEELs)", "US EPA High Production Volume Program Chemical List", "US FDA Indirect Food Additives: Adhesives and Components of Coatings - Substances for Use Only as Components of Adhesives - Adhesives", "US Toxic Substances Control Act (TSCA) - Inventory", "US TSCA Section 12(b) - List of Chemical Substances Subject to Export Notification Requirements", "US TSCA Section 5(a)(2) - Significant New Use Rules (SNURs)", "US TSCA Section 8 (a) - Preliminary Assessment Information Rules (PAIR) - Reporting List", "US TSCA Section 8 (d) - Health and Safety Data Reporting"

### **Section 16 - OTHER INFORMATION**

### LIMITED EVIDENCE

- Inhalation may produce health damage\*.
- Cumulative effects may result following exposure\*.
- Possible skin sensitizer\*.
- \* (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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