

# 6-Acetylmorphine-D3

**Novachem Pty Ltd** 

Version No: **2.4**Safety Data Sheet according to WHS and ADG requirements

Chemwatch Hazard Alert Code: 3

Issue Date: 10/09/2018 Print Date: 10/09/2018 S.GHS.AUS.EN

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

### **Product Identifier**

Product name	6-Acetylmorphine-D3
Chemical Name	acetonitrile
Synonyms	A-006
Proper shipping name	ACETONITRILE
Chemical formula	C2H3N
Other means of identification	Not Available
CAS number	75-05-8*

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

Relevant identified uses Laboratory Research Material

### Details of the supplier of the safety data sheet

Registered company name	Novachem Pty Ltd			
Address	Crissane Road, Heidelberg West Victoria 3081 Australia			
Telephone	+61384151255			
Fax	+61386250088			
Website	www.novachem.com.au			
Email	novachem@novachem.com.au			

# Emergency telephone number

Association / Organisation	Victorian Poisons Information Centre
Emergency telephone numbers	13 11 26
Other emergency telephone numbers	Not Available

# **SECTION 2 HAZARDS IDENTIFICATION**

# Classification of the substance or mixture

Poisons Schedule	Not Applicable
Classification <sup>[1]</sup>	Flammable Liquid Category 2, Acute Toxicity (Oral) Category 4, Acute Toxicity (Dermal) Category 4, Acute Toxicity (Inhalation) Category 4, Eye Irritation Category 2A
Legend:	1. Classified by Chemwatch; 2. Classification drawn from HSIS; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI

# Label elements

Hazard pictogram(s)





SIGNAL WORD DA

DANGER

### Hazard statement(s)

H225	Highly flammable liquid and vapour.		
H302	Harmful if swallowed.		
H312	Harmful in contact with skin.		
H332	Harmful if inhaled.		

Chemwatch: 9-463824 Version No: 2.4

# Page 2 of 10 6-Acetylmorphine-D3

Issue Date: **10/09/2018**Print Date: **10/09/2018** 

H319 Causes serious eye irritation.

### Precautionary statement(s) Prevention

P210	Keep away from heat/sparks/open flames/hot surfaces No smoking.		
P233	eep container tightly closed.		
P271	Use only outdoors or in a well-ventilated area.		
P240	Ground/bond container and receiving equipment.		

### Precautionary statement(s) Response

P363	Wash contaminated clothing before reuse.			
P370+P378	In case of fire: Use water spray/fog for extinction.			
P305+P351+P338	IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.			
P337+P313	If eye irritation persists: Get medical advice/attention.			

### Precautionary statement(s) Storage

P403+P235 Store in a well-ventilated place. Keep cool.

### Precautionary statement(s) Disposal

P501

Dispose of contents/container in accordance with local regulations.

### **SECTION 3 COMPOSITION / INFORMATION ON INGREDIENTS**

### Substances

CAS No	%[weight]	Name
75-05-8	99.99	acetonitrile
2784-73-8	0.01	6-acetylmorphine

### **Mixtures**

See section above for composition of Substances

### **SECTION 4 FIRST AID MEASURES**

## Description of first aid measures

3	
Eye Contact	If this product comes in contact with the eyes:  • Wash out immediately with fresh running water.  • Ensure complete irrigation of the eye by keeping eyelids apart and away from eye and moving the eyelids by occasionally lifting the upper and lower lids.  • Seek medical attention without delay; if pain persists or recurs seek medical attention.  • Removal of contact lenses after an eye injury should only be undertaken by skilled personnel.
Skin Contact	If skin 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.
Inhalation	<ul> <li>If fumes or combustion products are inhaled remove from contaminated area.</li> <li>Lay patient down. Keep warm and rested.</li> <li>Prostheses such as false teeth, which may block airway, should be removed, where possible, prior to initiating first aid procedures.</li> <li>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.</li> <li>Transport to hospital, or doctor.</li> </ul>
Ingestion	<ul> <li>IF SWALLOWED, REFER FOR MEDICAL ATTENTION, WHERE POSSIBLE, WITHOUT DELAY.</li> <li>For advice, contact a Poisons Information Centre or a doctor.</li> <li>Urgent hospital treatment is likely to be needed.</li> <li>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.</li> <li>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 SDS should be provided. Further action will be the responsibility of the medical specialist.</li> <li>If medical attention is not available on the worksite or surroundings send the patient to a hospital together with a copy of the SDS.</li> <li>Where medical attention is not immediately available or where the patient is more than 15 minutes from a hospital or unless instructed otherwise:</li> <li>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.</li> </ul>

### Indication of any immediate medical attention and special treatment needed

Treat symptomatically for a narcotic analgesic.

A vigorous program of symptomatic and supportive therapy has saved many victims of poisoning. The single most important element in therapy is the correction of anoxia by all available means: the maintenance of a patent airway, the administration of oxygen, the use of artificial respiration, and the injection of specific narcotic antagonists such as nalorphine, levallorphan or naloxone promptly antagonises the respiratory depression, coma and hypotension from overdoses of morphine, codeine, all semi-synthetics and almost all synthetic narcotics.

GOSSELIN et al: Clinical Toxicology of Commercial Products.

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

### 6-Acetylmorphine-D3

In fully conscious patients, remove swallowed poison by thorough gastric layage and emesis. The chances of removing a significant amount of the drug are better if treatment is started within the first two hours. If the patient is unconscious or depressed, emesis is contraindicated and the dangers of gastric lavage are not justified.

DREISBACH AND ROBERTSON: Handbook of Poisoning, Appleton & Lange

- For cyanide intoxication (and for certain nitriles which produce cyanide ion) Signs symptoms of acute cyanide poisoning reflect cellular hypoxia and are often non-specific.
  - Cyanosis may be a late finding.
  - A bradycardic, hypertensive and tachypneic patient suggests poisoning especially if CNS and cardiovascular depression subsequently occurs.
  - Immediate attention should be directed towards assisted ventilation, administration of 100% oxygen, insertion of intravenous lines and institution of cardiac monitoring.
  - Obtain an arterial blood gas immediately and correct any severe metabolic acidosis (pH below 7.15).
- Mildly symptomatic patients generally require supportive care alone. Nitrites should not be given indiscriminately in all cases of moderate to severe poisoning, they should be given in conjunction with thiosulfate. As a temporizing measure supply amyl nitrite perles (0.2ml inhaled 30 seconds every minute) until intravenous lines for sodium nitrite are established. 10 ml of a 3% solution is administered over 4 minutes to produce 20% methaemoglobin in adults. Follow directly with 50 ml of 25% sodium thiosulfate, at the same rate, IV. If symptoms reappear or persist within 1/2-1 hour, repeat nitrite and thiosulfate at 50% of initial dose. As the mode of action involves the metabolic conversion of the thiosulfate to thiocyanate, renal failure may enhance thiocyanate toxicity.
- ▶ Methylene blue is not an antidote. [Ellenhorn and Barceloux: Medical Toxicology]

If amyl nitrite intervention is employed then Medical Treatment Kits should contain the following:

- One box containing one dozen amyl nitrite ampoules
- Two sterile ampoules of sodium nitrite solution (10 mL of a 3% solution in each)
- ▶ Two sterile ampoules of sodium thiosulfate solution (50 mL of a 25% solution in each)
- One 10 mL sterile syringe. One 50 mL sterile syringe. Two sterile intravenous needles. One tourniquet.
- One dozen gauze pads.
- Latex gloves
- ► A "Biohazard" bag for disposal of bloody/contaminated equipment.
- · A set of cyanide instructions on first aid and medical treatment.
- Notes on the use of amyl nitrite:-
- AN is highly volatile and flammable do not smoke or use around a source of ignition.
- If treating patient in a windy or draughty area provide some shelter or protection (shirt, wall, drum, cupped hand etc.) to prevent amyl nitrite vapour from being blown away. Keep ampoule upwind from the nose, the objective is to get amyl nitrite into the patients lungs
- Rescuers should avoid AN inhalation to avoid becoming dizzy and losing competence.
- Lay the patient down. Since AN dilates blood vessels and lowers blood pressure, lying down will help keep patient conscious.
- DO NOT overuse excessive use might put the patient into shock. Experience at DuPont plants has not shown any serious after-effects from treatment with amyl nitrite.

### ADDITIONAL NOTES:

Major medical treatment procedures may vary e.g. US (FDA method as recommended by DuPont) uses amyl nitrite as a methaemoglobin generator, followed by treatment with sodium nitrite and then sodium thiosulfate

MODES OF ACTION: Amyl nitrite (AN) reacts with haemoglobin (HB) to form about 5% methaemoglobin (MHB). Sodium nitrite (NaNO2) reacts with haemoglobin to form approximately 20-30% methaemoglobin, Methaemoglobin attracts cvanide ions (CN) from tissue and binds with them to become cvanmethaemoglobin (CNMHB). Sodium thiosulfate (Na2S2O3) converts cyanmethaemoglobin to thiocyanate (HSCN) which is excreted by the kidneys. i.e. AN + HB = MHB NaNO2 + HB = MHB CN + MHB = CNMHB Na2S2O3 + CNMHB + O2 = HSCN

- ▶ The administration of the antidote salts is intravenous in normal saline, Ringers lactate or other available IV fluid.
- European practice may use 4-dimethylaminophenol (DMAP) as a methaemoglobin generator. Also hydroxycobalamin (Vitamin B12a) is used. Hydroxycobalamin works by reacting with cyanide to form cyanocobalamin (Vitamin B12) which is excreted in the urine.
- ▶ European and Australian NOHSC (ASCC) propose dicobalt edetate (Kelocyanor) as antidote. This acts by chelating cyanide to form stable cobalticyanide, which is excreted in the urine. In all cases hyperbaric therapy may increase the efficiency of a cyanide antidote kit.

## **SECTION 5 FIREFIGHTING MEASURES**

Fire Incompatibility

# **Extinguishing media**

- Dry chemical powder.
- BCF (where regulations permit).
- Carbon dioxide.

### Special hazards arising from the substrate or mixture

Advice for firefighters					
Fire Fighting	<ul> <li>Alert Fire Brigade and tell them location and nature of hazard.</li> <li>May be violently or explosively reactive.</li> <li>Wear breathing apparatus plus protective gloves in the event of a fire.</li> <li>Prevent, by any means available, spillage from entering drains or water course.</li> </ul>				
Fire/Explosion Hazard	Liquid and vapour are highly flammable.  Severe fire hazard when exposed to heat, flame and/or oxidisers.  Vapour may travel a considerable distance to source of ignition.  Heating may cause expansion or decomposition leading to violent rupture of containers.  Combustion products include:				

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

carbon dioxide (CO2)

nitrogen oxides (NOx)

other pyrolysis products typical of burning organic material.

HAZCHEM

### **SECTION 6 ACCIDENTAL RELEASE MEASURES**

# Personal precautions, protective equipment and emergency procedures

See section 8

## **Environmental precautions**

See section 12

Chemwatch: 9-463824 Page 4 of 10 Issue Date: 10/09/2018 Version No: 2.4 Print Date: 10/09/2018

### 6-Acetylmorphine-D3

Methods and material for containment and cleaning up

## Minor Spills

Major Spills

- ▶ Remove all ignition sources.
- Clean up all spills immediately.
- Avoid breathing vapours and contact with skin and eyes.
- Control personal contact with the substance, by using protective equipment.

### ▶ DO NOT touch the spill materia

### For alkyl nitriles:

# For residue:

- ► Add alkaline hypochlorite solution to spill to produce cyanate.
- Neutralise liquid, and absorb with sawdust
- Collect solid residues and seal in drums for disposal.
- Wash spill area with large quantities of water.
- Clear area of personnel and move upwind.
- Alert Fire Brigade and tell them location and nature of hazard.
- May be violently or explosively reactive.
- Wear breathing apparatus plus protective gloves.

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

### **SECTION 7 HANDLING AND STORAGE**

### Precautions for safe handling

# Safe handling

- Containers, even those that have been emptied, may contain explosive vapours.
- Do NOT cut, drill, grind, weld or perform similar operations on or near containers
- 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.

### Other information

NOTE: Special security requirements may be mandated under Federal/State Regulation(s).

- ► Store in original containers
- Store in vault fitted with warning devices or detectors recommended by various Federal/State authorities. ▶ Store in vault used only for the purpose of storage of drugs of addiction.

### Conditions for safe storage, including any incompatibilities

- ▶ Packaging as recommended by manufacturer.
- Check that containers are clearly labelled.
- Tamper-proof containers

# Suitable container

- Polyethylene or polypropylene containers.
- Glass container is suitable for laboratory quantities
- For low viscosity materials (i): Drums and jerry cans 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.

### Acetonitrile

- ► forms cyanide gas on contact with steam
- reacts violently with oxidisers such as chlorine, bromine, fluorine; with chlorosulfonic acid, oleum or sulfuric acid
- is incompatible with water (especially if acid or alkaline), acids, caustics, nitrating agents, indium, nitrogen tetroxide, sulfur trioxide, iron(III) salts of perchlorate, nitrogen fluoride compounds
- attacks most rubber and plastics
- ▶ may accumulate electrical charges, causing ignition of vapours
- Contact with acids produces toxic fumes
- Nitriles may polymerise in the presence of metals and some metal compounds.
- They are incompatible with acids; mixing nitriles with strong oxidising acids can lead to extremely violent reactions
- Nitriles are generally incompatible with other oxidising agents such as peroxides and epoxides.
- The combination of bases and nitriles can produce hydrogen cyanide.
  - The covalent cyano group is endothermic and many organic nitriles are reactive under certain conditions; N-cyano derivatives are reactive or unstable.
  - The majority of endothermic compounds are thermodynamically unstable and may decompose explosively under various circumstances of initiation.
  - Many but not all endothermic compounds have been involved in decompositions, reactions and explosions and, in general, compounds with significantly positive values of standard heats of formation, may be considered suspect on stability grounds.

BRETHERICK L.: Handbook of Reactive Chemical Hazards

### WARNING.

May decompose violently or explosively on contact with other substances.

- ▶ This substance, or one of its components, is one of the relatively few compounds which are described as "endothermic" i.e. heat is absorbed into the compound, rather than released from it, during its formation.
  - ► The majority of endothermic compounds are thermodynamically unstable and may decompose explosively under various circumstances of initiation.
- Many but not all endothermic compounds have been involved in decompositions, reactions and explosions and, in general, compounds with significantly positive values of standard heats of formation, may be considered suspect on stability grounds.
- Avoid reaction with oxidising agents

# **SECTION 8 EXPOSURE CONTROLS / PERSONAL PROTECTION**

### Control parameters

# OCCUPATIONAL EXPOSURE LIMITS (OEL)

Storage incompatibility

### INGREDIENT DATA

Source	Ingredient	Material name	TWA	STEL	Peak	Notes
Australia Exposure Standards	acetonitrile	Acetonitrile	40 ppm / 67 mg/m3	101 mg/m3 / 60 ppm	Not Available	Not Available

Version No. 2.4

Page 5 of 10

6-Acetylmorphine-D3

Issue Date: **10/09/2018**Print Date: **10/09/2018** 

Ingredient	Material name	TEEL-1	TEEL-2	TEEL-3
acetonitrile	Acetonitrile Not Available		Not Available	Not Available
Ingredient	Original IDLH		Revised IDLH	
acetonitrile	500 ppm		137 ppm	
6-acetylmorphine	Not Available		Not Available	

### **Exposure controls**

# Appropriate engineering controls

Engineering controls are used to remove a hazard or place a barrier between the worker and the hazard. Well-designed engineering controls can be highly effective in protecting workers and will typically be independent of worker interactions to provide this high level of protection.

The basic types of engineering controls are: Process controls which involve changing the way a job activity or process is done to reduce the risk.

Enclosure and/or isolation of emission source which keeps a selected hazard "physically" away from the worker and ventilation that strategically "adds" and "removes" air in the work environment.

### Personal protection









### Eye and face protection

- ► 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 lenses or restrictions on use, should be created for each workplace or task.

### Skin protection

See Hand protection below

- ► Wear chemical protective gloves, e.g. PVC.
- ▶ Wear safety footwear or safety gumboots, e.g. Rubber

## Hands/feet protection

The selection of suitable gloves does not only depend on the material, but also on further marks of quality which vary from manufacturer to manufacturer. Where the chemical is a preparation of several substances, the resistance of the glove material can not be calculated in advance and has therefore to be checked prior to the application.

The exact break through time for substances has to be obtained from the manufacturer of the protective gloves and has to be observed when making a final choice.

Personal hygiene is a key element of effective hand care.

for acetonitrile:

Butyl rubber, PVAL, Teflon, Saranex, Silvershield, Viton/chlorobutyl are all highly resistant to permeation

# Body protection

See Other protection below

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

# Other protection

- ▶ Eyewash un
- 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 or conductive footwear should be considered. Conductive footwear describes a boot or shoe with a sole made from a conductive compound chemically bound to the bottom components, for permanent control to electrically ground the foot an shall dissipate static electricity from the body to reduce the possibility of ignition of volatile compounds.

### Recommended material(s)

### GLOVE SELECTION INDEX

Glove selection is based on a modified presentation of the:

"Forsberg Clothing Performance Index".

The effect(s) of the following substance(s) are taken into account in the *computer-generated* selection:

6-Acetylmorphine-D3

Material	СРІ
BUTYL	А
BUTYL/NEOPRENE	Α
CPE	А
PE/EVAL/PE	А
PVA	А
SARANEX-23	А
NEOPRENE	В
TEFLON	В
NATURAL RUBBER	С
NATURAL+NEOPRENE	С
NITRILE	С
VITON/NEOPRENE	С

<sup>\*</sup> CPI - Chemwatch Performance Index

A: Best Selection

B: Satisfactory; may degrade after 4 hours continuous immersion

C: Poor to Dangerous Choice for other than short term immersion

**NOTE**: As a series of factors will influence the actual performance of the glove, a final selection must be based on detailed observation. -

\* Where the glove is to be used on a short term, casual or infrequent basis, factors such as

### Respiratory protection

Cartridge respirators should never be used for emergency ingress or in areas of unknown vapour concentrations or oxygen content. The wearer must be warned to leave the contaminated area immediately on detecting any odours through the respirator. The odour may indicate that the mask is not functioning properly, that the vapour concentration is too high, or that the mask is not properly fitted. Because of these limitations, only restricted use of cartridge respirators is considered appropriate.

Issue Date: **10/09/2018**Print Date: **10/09/2018** 

"feel" or convenience (e.g. disposability), may dictate a choice of gloves which might otherwise be unsuitable following long-term or frequent use. A qualified practitioner should be consulted.

# **SECTION 9 PHYSICAL AND CHEMICAL PROPERTIES**

### Information on basic physical and chemical properties

Appearance	Not Available			
Physical state	Liquid	Relative density (Water = 1)	0.8	
Odour	Not Available	t Available Partition coefficient n-octanol / water		
Odour threshold	Not Available	Auto-ignition temperature (°C)	524.0	
pH (as supplied)	Not Applicable	Decomposition temperature	Not Available	
Melting point / freezing point (°C)	-45	Viscosity (cSt)	Not Available	
Initial boiling point and boiling range (°C)	81.1	Molecular weight (g/mol)	41.05 Pure	
Flash point (°C)	5.5 (OC)	Taste	Not Available	
Evaporation rate	5.79 BuAc=1 BuAC = 1	Explosive properties	Not Available	
Flammability	HIGHLY FLAMMABLE.	Oxidising properties	Not Available	
Upper Explosive Limit (%)	16.0	Surface Tension (dyn/cm or mN/m)	Not Available	
Lower Explosive Limit (%)	4.4	Volatile Component (%vol)	100	
Vapour pressure (kPa)	13.3 @ 27 deg.C	Gas group	Not Available	
Solubility in water (g/L)	Miscible	pH as a solution (1%)	Not Available	
Vapour density (Air = 1)	1.4	VOC g/L	792.8	

### **SECTION 10 STABILITY AND REACTIVITY**

Reactivity	See section 7		
Chemical stability	<ul> <li>Presence of elevated temperatures.</li> <li>Unstable in the presence of incompatible materials.</li> <li>Product is considered stable.</li> <li>Hazardous polymerisation will not occur.</li> </ul>		
Possibility of hazardous reactions	See section 7		
Conditions to avoid	See section 7		
Incompatible materials	See section 7		
Hazardous decomposition products	See section 5		

# **SECTION 11 TOXICOLOGICAL INFORMATION**

# Information on toxicological effects

Inhaled	Inhalation of vapours or aerosols (mists, fumes), generated by the material during the course of normal handling, may be harmful.  The material is not thought to produce respiratory irritation (as classified by EC Directives using animal models). Nevertheless inhalation of vapours, fumes or aerosols, especially for prolonged periods, may produce respiratory discomfort and occasionally, distress.  The smell of acetonitrile does not give enough warning of exposure. The gas is highly toxic, and inhaling it can cause loss of consciousness.
Ingestion	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.  Nitrile poisoning exhibits similar symptoms to poisoning due to hydrogen cyanide. The substances irritate the eyes and skin, and are absorbed quickly and completely through the skin.  Morphine and other analgesics cause nausea, vomiting, constipation, drowsiness and confusion. Urination can be difficult, and the bowel and bile ducts can spasm.  Cyanide poisoning can cause increased saliva output, nausea without vomiting, anxiety, confusion, vertigo, dizziness, stiffness of the lower jaw, convulsions, spasm, paralysis, coma and irregular heartbeat, and stimulation of breathing followed by failure. Often the skin becomes cyanosed (blue-grey), and this is often delayed.
Skin Contact	Skin contact with the material may be harmful; systemic effects may result following absorption.  The material is not thought to be a skin irritant (as classified by EC Directives using animal models). Temporary discomfort, however, may result from prolonged dermal exposures.  Contact dermatitis has been reported with morphine and other narcotic analgesics.  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.
Eye	There is evidence that material may produce eye irritation in some persons and produce eye damage 24 hours or more after instillation. Severe inflammation may be expected with pain.
Chronic	Long-term exposure to the product is not thought to produce chronic effects adverse to the health (as classified by EC Directives using animal models); nevertheless exposure by all routes should be minimised as a matter of course.  Chronic exposure to cyanides and certain nitriles may result in interference to iodine uptake by thyroid gland and its consequent enlargement. This occurs following metabolic conversion of the cyanide moiety to thiocyanate.  Chronic morphine poisoning or addiction causes pin-point pupils, rapid mood changes and poor social adaptation. As dependence and tolerance occurs, there is an overwhelming need to continue taking the drug or similar drugs and to increase the dose.

Chemwatch: **9-463824**Version No: **2.4** 

Page **7** of **10** 

6-Acetylmorphine-D3

Issue Date: **10/09/2018**Print Date: **10/09/2018** 

G-Acetylmorphine-D3  G-Acetylmorphine-D3  G-Acetylmorphine-D3  G-Acetylmorphine-D3  G-Acetylmorphine-D3  G-Acetylmorphine  Acetonitrile  G-Acetylmorphine  G-Acetylmorphine  G-Acetylmorphine  Acetonitrile  G-Acetylmorphine  G-Acetylmorphine-D3 & Acetholicania of the skin.  Manning: Abuse can lead to habituation. Subject to Federal and State Regulations. Narcotic Substance, Schedule I (UN).  Absorption of acetonitrile occurs after oral, skin, or inhalation exposure. The liquid or vapour is irritating to the skin, eyes, and airways. At high end carbod control of the skin.  Acetonitrile  Acute Toxicity  General (rabbit) LD50: 980 mg/kg <sup>[2]</sup> Inhalation (rat) LD50: -2000 mg/kg <sup>[1]</sup> Inhalation (rat) LD50					
Inhalation (rat) LC50: 17080.4889 mg/4 h <sup>[1]</sup> Oral (rat) LD50: <2000 mg/kg <sup>[1]</sup> TOXICITY  Dermal (rabbit) LD50: 980 mg/kg <sup>[2]</sup> Inhalation (rat) LC50: 17080.4889 mg/4 h <sup>[1]</sup> Oral (rat) LD50: 980 mg/kg <sup>[2]</sup> Eye (rabbit)-20 mg (open)-SEVERE  Inhalation (rat) LC50: 17080.4889 mg/4 h <sup>[1]</sup> Skin (rabbit)-500 mg (open)-mild  TOXICITY  IRRITATION  Not Available  TOXICITY  Not Available  I. Value obtained from Europe ECHA Registered Substances - Acute toxicity 2.* Value obtained from manufacturer's SDS. Unless otherwise spe data extracted from RTECS - Register of Toxic Effect of chemical Substances  ACETONITRILE  The material may produce severe irritation to the eye causing pronounced inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis. The material may cause skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of scaling and thickening of the skin. No significant acute toxicological data identified in literature search.  WARNING: Abuse can lead to habituation. Subject to Federal and State Regulations. Narcotic Substance, Schedule I (UN).  Absorption of acetonitrile occurs after oral, skin, or inhalation exposure. The liquid or vapour is irritating to the skin, eyes, and airways. At high encodess, death can occur quickly from respiratory failure. Lower doses cause typical symptoms of cyanide poisoning such as salivation, nausea, von anxiety, confusion, rapid and difficult breathing, rapid pulse, unconsciousness, and convulsions.		TOXICITY			IRRITATION
Inhalation (rat) LCS0: 17080.4889 mg/l4 h <sup>11</sup>     Oral (rat) LD50: 2000 mg/kg <sup>11</sup>		Dermal (rabbit) LD50: 980 mg/kg <sup>[2]</sup>	Dermal (rabbit) LD50: 980 mg/kg <sup>[2]</sup>		
acetonitrile    TOXICITY	6-Acetylmorphine-D3	Inhalation (rat) LC50: 17080.4889 mg/l4 h <sup>[1]</sup>			
Dermal (rabbit) LD50: 980 mg/kg <sup>[2]</sup> Dermal (rabbit) LD50: 17080.4889 mg/kd h <sup>[1]</sup> Skin (rabbit):20 mg (open)-SEVERE  Inhalation (rat) LD50: 17080.4889 mg/kd h <sup>[1]</sup> Skin (rabbit):500 mg (open)-mild  Final (rabbit):500 mg (open)-mild  TOXICITY  Not Available  TOXICITY  Not Available  1. Value obtained from Europe ECHA Registered Substances - Acute toxicity 2.* Value obtained from manufacturer's SDS. Unless otherwise spedata extracted from RTECS - Register of Toxic Effect of chemical Substances  ACETONITRILE  The material may produce severe irritation to the eye causing pronounced inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis. The material may cause skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of scaling and thickening of the skin.  No significant acute toxicological data identified in literature search.  WARNING: Abuse can lead to habituation. Subject to Federal and State Regulations. Narcotic Substance, Schedule I (UN).  Absorption of acetonitrile occurs after oral, skin, or inhalation exposure. The liquid or vapour is irritating to the skin, eyes, and airways. At high end doses, death can occur quickly from respiratory failure. Lower doses cause typical symptoms of cyanide poisoning such as salivation, nausea, von anxiety, confusion, rapid and difficult breathing, rapid pulse, unconsciousness, and convulsions.		Oral (rat) LD50: <2000 mg/kg <sup>[1]</sup>			
Dermal (rabbit) LD50: 980 mg/kg <sup>[2]</sup> Dermal (rabbit) LD50: 980 mg/kg <sup>[2]</sup> Inhalation (rat) LC50: 17080.4889 mg/4 h <sup>[1]</sup> Skin (rabbit):500 mg (open)-SEVERE  Skin (rabbit):500 mg (open)-mild  Foxicity  Not Available  TOXICITY  Not Available  Invalue obtained from Europe ECHA Registered Substances - Acute toxicity 2.* Value obtained from manufacturer's SDS. Unless otherwise spedata extracted from RTECS - Register of Toxic Effect of chemical Substances  ACETONITRILE  The material may produce severe irritation to the eye causing pronounced inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis. The material may cause skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of scaling and thickening of the skin.  No significant acute toxicological data identified in literature search.  WARNING: Abuse can lead to habituation. Subject to Federal and State Regulations. Narcotic Substance, Schedule I (UN).  Absorption of acetonitrile occurs after oral, skin, or inhalation exposure. The liquid or vapour is irritating to the skin, eyes, and airways. At high end doses, death can occur quickly from respiratory failure. Lower doses cause typical symptoms of cyanide poisoning such as salivation, nausea, von anxiety, confusion, rapid and difficult breathing, rapid pulse, unconsciousness, and convulsions.					
Inhalation (rat) LC50: 17080.4889 mg/4 h <sup>[1]</sup> Skin (rabbit):500 mg (open)-mild  For a cetylmorphine  TOXICITY Not Available  TOXICITY Not Available  Toxic Effect of chemical Substances - Acute toxicity 2: Value obtained from manufacturer's SDS. Unless otherwise spendata extracted from RTECS - Register of Toxic Effect of chemical Substances  ACETONITRILE  The material may produce severe irritation to the eye causing pronounced inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis. The material may cause skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of scaling and thickening of the skin.  No significant acute toxicological data identified in literature search.  WARNING: Abuse can lead to habituation. Subject to Federal and State Regulations. Narcotic Substance, Schedule I (UN).  Absorption of acetonitrile occurs after oral, skin, or inhalation exposure. The liquid or vapour is irritating to the skin, eyes, and airways. At high end doses, death can occur quickly from respiratory failure. Lower doses cause typical symptoms of cyanide poisoning such as salivation, nausea, von anxiety, confusion, rapid and difficult breathing, rapid pulse, unconsciousness, and convulsions.		TOXICITY	IF	RRITATION	
Inhalation (rat) LC50: 47080.4889 mg/l4 h <sup>[1]</sup> Oral (rat) LD50: <2000 mg/kg <sup>[1]</sup> FOXICITY  Not Available  1. Value obtained from Europe ECHA Registered Substances - Acute toxicity 2.* Value obtained from manufacturer's SDS. Unless otherwise spedata extracted from RTECS - Register of Toxic Effect of chemical Substances  ACETONITRILE  The material may produce severe irritation to the eye causing pronounced inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis. The material may cause skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of scaling and thickening of the skin.  No significant acute toxicological data identified in literature search.  WARNING: Abuse can lead to habituation. Subject to Federal and State Regulations. Narcotic Substance, Schedule I (UN).  Absorption of acetonitrile occurs after oral, skin, or inhalation exposure. The liquid or vapour is irritating to the skin, eyes, and airways. At high end doses, death can occur quickly from respiratory failure. Lower doses cause typical symptoms of cyanide poisoning such as salivation, nausea, von anxiety, confusion, rapid and difficult breathing, rapid pulse, unconsciousness, and convulsions.		Dermal (rabbit) LD50: 980 mg/kg <sup>[2]</sup>	E	eye (rabbit):20 mg (open)-	SEVERE
6-acetylmorphine  Legend:  1. Value obtained from Europe ECHA Registered Substances - Acute toxicity 2.* Value obtained from manufacturer's SDS. Unless otherwise spedata extracted from RTECS - Register of Toxic Effect of chemical Substances  The material may produce severe irritation to the eye causing pronounced inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis. The material may cause skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of scaling and thickening of the skin.  6-ACETYLMORPHINE  6-ACETYLMORPHINE  MARNING: Abuse can lead to habituation. Subject to Federal and State Regulations. Narcotic Substance, Schedule I (UN).  Absorption of acetonitrile occurs after oral, skin, or inhalation exposure. The liquid or vapour is irritating to the skin, eyes, and airways. At high end doses, death can occur quickly from respiratory failure. Lower doses cause typical symptoms of cyanide poisoning such as salivation, nausea, von anxiety, confusion, rapid and difficult breathing, rapid pulse, unconsciousness, and convulsions.	acetonitrile	Inhalation (rat) LC50: 17080.4889 mg/l4 h <sup>[1]</sup>	S	Skin (rabbit):500 mg (oper	n)-mild
Not Available   Not Available   Not Available		Oral (rat) LD50: <2000 mg/kg <sup>[1]</sup>			
Not Available   Not Available   Not Available					
Legend:  1. Value obtained from Europe ECHA Registered Substances - Acute toxicity 2.* Value obtained from manufacturer's SDS. Unless otherwise speciate extracted from RTECS - Register of Toxic Effect of chemical Substances  ACETONITRILE  The material may produce severe irritation to the eye causing pronounced inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis. The material may cause skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of scaling and thickening of the skin.  No significant acute toxicological data identified in literature search.  WARNING: Abuse can lead to habituation. Subject to Federal and State Regulations. Narcotic Substance, Schedule I (UN).  Absorption of acetonitrile occurs after oral, skin, or inhalation exposure. The liquid or vapour is irritating to the skin, eyes, and airways. At high end doses, death can occur quickly from respiratory failure. Lower doses cause typical symptoms of cyanide poisoning such as salivation, nausea, von anxiety, confusion, rapid and difficult breathing, rapid pulse, unconsciousness, and convulsions.  Carcinogenicity		TOXICITY	IRRITAT	ΓΙΟΝ	
ACETONITRILE  The material may produce severe irritation to the eye causing pronounced inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis. The material may cause skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of scaling and thickening of the skin.  No significant acute toxicological data identified in literature search.  WARNING: Abuse can lead to habituation. Subject to Federal and State Regulations. Narcotic Substance, Schedule I (UN).  Absorption of acetonitrile occurs after oral, skin, or inhalation exposure. The liquid or vapour is irritating to the skin, eyes, and airways. At high end doses, death can occur quickly from respiratory failure. Lower doses cause typical symptoms of cyanide poisoning such as salivation, nausea, von anxiety, confusion, rapid and difficult breathing, rapid pulse, unconsciousness, and convulsions.	6-acetylmorphine	Not Available	Not Avai	Not Available	
ACETONITRILE  The material may produce severe irritation to the eye causing pronounced inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis. The material may cause skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of scaling and thickening of the skin.  No significant acute toxicological data identified in literature search.  WARNING: Abuse can lead to habituation. Subject to Federal and State Regulations. Narcotic Substance, Schedule I (UN).  Absorption of acetonitrile occurs after oral, skin, or inhalation exposure. The liquid or vapour is irritating to the skin, eyes, and airways. At high end doses, death can occur quickly from respiratory failure. Lower doses cause typical symptoms of cyanide poisoning such as salivation, nausea, von anxiety, confusion, rapid and difficult breathing, rapid pulse, unconsciousness, and convulsions.					
ACETONITRILE  conjunctivitis. The material may cause skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of scaling and thickening of the skin.  No significant acute toxicological data identified in literature search.  WARNING: Abuse can lead to habituation. Subject to Federal and State Regulations. Narcotic Substance, Schedule I (UN).  6-Acetylmorphine-D3 & ACETONITRILE  ACETONITRILE  ACETONITRILE  ACUATE Toxicity  Carcinogenicity  Carcinogenicity	Legend:	Value obtained from Europe ECHA Registered Substances - Acute toxicity 2.* Value obtained from manufacturer's SDS. Unless otherwise specified data extracted from RTECS - Register of Toxic Effect of chemical Substances			
ACETONITRILE  conjunctivitis. The material may cause skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of scaling and thickening of the skin.  No significant acute toxicological data identified in literature search.  WARNING: Abuse can lead to habituation. Subject to Federal and State Regulations. Narcotic Substance, Schedule I (UN).  6-Acetylmorphine-D3 & ACETONITRILE  ACETONITRILE  ACETONITRILE  ACUATE Toxicity  Carcinogenicity  Carcinogenicity					
6-ACETYLMORPHINE  WARNING: Abuse can lead to habituation. Subject to Federal and State Regulations. Narcotic Substance, Schedule I (UN).  6-Acetylmorphine-D3 & ACETONITRILE  ACETONITRILE  Acute Toxicity  WARNING: Abuse can lead to habituation. Subject to Federal and State Regulations. Narcotic Substance, Schedule I (UN).  Absorption of acetonitrile occurs after oral, skin, or inhalation exposure. The liquid or vapour is irritating to the skin, eyes, and airways. At high end doses, death can occur quickly from respiratory failure. Lower doses cause typical symptoms of cyanide poisoning such as salivation, nausea, von anxiety, confusion, rapid and difficult breathing, rapid pulse, unconsciousness, and convulsions.	ACETONITRILE	conjunctivitis.  The material may cause skin irritation after prolonged or re			
WARNING: Abuse can lead to habituation. Subject to Federal and State Regulations. Narcotic Substance, Schedule I (UN).  6-Acetylmorphine-D3 & ACETONITRILE  ACETONITRILE  Acute Toxicity  WARNING: Abuse can lead to habituation. Subject to Federal and State Regulations. Narcotic Substance, Schedule I (UN).  Absorption of acetonitrile occurs after oral, skin, or inhalation exposure. The liquid or vapour is irritating to the skin, eyes, and airways. At high end doses, death can occur quickly from respiratory failure. Lower doses cause typical symptoms of cyanide poisoning such as salivation, nausea, von anxiety, confusion, rapid and difficult breathing, rapid pulse, unconsciousness, and convulsions.		No significant acute toxicological data identified in literatu	re search.		
ACETONITRILE  doses, death can occur quickly from respiratory failure. Lower doses cause typical symptoms of cyanide poisoning such as salivation, nausea, von anxiety, confusion, rapid and difficult breathing, rapid pulse, unconsciousness, and convulsions.  Acute Toxicity  Carcinogenicity	6-ACETYLMORPHINE	WARNING: Abuse can lead to habituation. Subject to Fe-	deral and State Regulations.	Narcotic Substance, Sche	edule I (UN).
		doses, death can occur quickly from respiratory failure. Lov	wer doses cause typical symp	otoms of cyanide poisoning	
Skin Irritation/Corrosion			<b>0</b> '		
	Acute Toxicity	•	Carcinog	genicity 🚫	
Serious Eye Damage/Irritation STOT - Single Exposure		· ·			
Respiratory or Skin sensitisation STOT - Repeated Exposure	Skin Irritation/Corrosion	0	Reprodu	uctivity 🛇	
Mutagenicity Aspiration Hazard	Skin Irritation/Corrosion Serious Eye Damage/Irritation Respiratory or Skin	<ul><li>○</li><li>✓</li></ul>	Reprodu STOT - Single Ex	uctivity O	

Legend:

🗶 – Data available but does not fill the criteria for classification

Data available to make classification

O - Data Not Available to make classification

# **SECTION 12 ECOLOGICAL INFORMATION**

# Toxicity

	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
6-Acetylmorphine-D3	LC50	96	Fish	>100mg/L	4
	NOEC	24	Crustacea	0.00001mg/L	4
	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
acetonitrile	LC50	96	Fish	>100mg/L	4
	NOEC	24	Crustacea	0.00001mg/L	4
C as a to down a small in a	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
6-acetylmorphine	Not Available	Not Available	Not Available	Not Available	Not Available

Legend:

Extracted from 1. IUCLID Toxicity Data 2. Europe ECHA Registered Substances - Ecotoxicological Information - Aquatic Toxicity 3. EPIWIN Suite V3.12 (QSAR) - Aquatic Toxicity Data (Estimated) 4. US EPA, Ecotox database - Aquatic Toxicity Data 5. ECETOC Aquatic Hazard Assessment Data 6. NITE (Japan) - Bioconcentration Data 7. METI (Japan) - Bioconcentration Data 8. Vendor Data

Soil Guidelines: Dutch Criteria: free cyanide: 1 mg/kg (target)

20 mg/kg (intervention) complex cyanide (pH 5): 5 mg/kg (target)

50 mg/kg (intervention)

Air Quality Standards: no safe guidelines recommended due to carcinogenic properties.

On the basis of available evidence concerning either toxicity, persistence, potential to accumulate and or observed environmental fate and behaviour, the material may present a danger, immediate or long-term and /or delayed, to the structure and/ or functioning of natural ecosystems.

### Version No: 2.4

Issue Date: 10/09/2018
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6-Acetylmorphine-D3

Abiotic Effects: Acetonitrile is a volatile organic compound (VOC) substance, thus it is a contributor to the formation of photochemical smog in the presence of other VOCs.

Transport: Acetonitrile is primarily removed by volatilization and leaching into groundwater. It has low adsorption potential to soils. Air - Acetonitrile may persist in the troposphere and can be transported over long distances.

DO NOT discharge into sewer or waterways

### Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air	
acetonitrile	HIGH (Half-life = 360 days)	HIGH (Half-life = 541.29 days)	

### **Bioaccumulative potential**

Ingredient	Bioaccumulation
acetonitrile	LOW (BCF = 0.4)

### Mobility in soil

Ingredient	Mobility
acetonitrile	LOW (KOC = 4.5)

### **SECTION 13 DISPOSAL CONSIDERATIONS**

### Waste treatment methods

- ► Containers may still present a chemical hazard/ danger when empty.
- ► Return to supplier for reuse/ recycling if possible.

### Otherwise

- ▶ If container can not be cleaned sufficiently well to ensure that residuals do not remain or if the container cannot be used to store the same product, then puncture containers, to prevent re-use, and bury at an authorised landfill.
- ▶ Where possible retain label warnings and SDS and observe all notices pertaining to the product.

Valuable substance, hold all residues for recovery. Disposal of the material must be carried out in accordance with the requirements of the relevant Federal/State Act(s) or Code(s) regulating the disposal of Drugs of Addiction.

- ► Consult manufacturer/supplier for recycling options.
- ► Decontaminate empty containers with water; incinerate plastic bags.

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:

### Product / Packaging disposal

- ▶ 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.

- ▶ DO NOT allow wash water from cleaning or process equipment to enter drains
- It may be necessary to collect all wash water for treatment before disposal.
- ► In all cases disposal to sewer may be subject to local laws and regulations and these should be considered first.
- Where in doubt contact the responsible authority.
- Recycle wherever possible.
- Consult manufacturer for recycling options or consult local or regional waste management authority for disposal if no suitable treatment or disposal facility can be identified.
- Dispose of by: burial in a land-fill specifically licensed to accept chemical and / or pharmaceutical wastes or Incineration in a licensed apparatus (after admixture with suitable combustible material).
- ▶ Decontaminate empty containers.

# **SECTION 14 TRANSPORT INFORMATION**

# Labels Required



•2YE

Marine Pollutant	L
HAZCHEM	

# Land transport (ADG)

UN number	1648			
UN proper shipping name	ACETONITRILE			
Transport hazard class(es)	Class 3 Subrisk Not Applicable			
Packing group	1			
Environmental hazard	Not Applicable			
Special precautions for user	Special provisions Not Applicable  Limited quantity 1 L			

Chemwatch: **9-463824**Version No: **2.4** 

Page **9** of **10** 

6-Acetylmorphine-D3

Issue Date: 10/09/2018 Print Date: 10/09/2018

# Air transport (ICAO-IATA / DGR)

UN number	1648			
UN proper shipping name	Acetonitrile			
Transport hazard class(es)	ICAO/IATA Class ICAO / IATA Subrisk ERG Code	3 k Not Applicable 3L		
Packing group	II .			
Environmental hazard	Not Applicable			
Special precautions for user	Special provisions  Cargo Only Packing Instructions  Cargo Only Maximum Qty / Pack  Passenger and Cargo Packing Instructions  Passenger and Cargo Maximum Qty / Pack  Passenger and Cargo Limited Quantity Packing Instructions  Passenger and Cargo Limited Maximum Qty / Pack		Not Applicable  364  60 L  353  5 L  Y341  1 L	

# Sea transport (IMDG-Code / GGVSee)

UN number	1648
UN proper shipping name	ACETONITRILE
Transport hazard class(es)	IMDG Class 3 IMDG Subrisk Not Applicable
Packing group	П
Environmental hazard	Not Applicable
Special precautions for user	EMS Number F-E , S-D Special provisions Not Applicable Limited Quantities 1 L

# Transport in bulk according to Annex II of MARPOL and the IBC code

SOURCE	PRODUCT NAME	POLLUTION CATEGORY	SHIP TYPE
	Acetonitrile Acetonitrile (Low purity grade)	Z Y	2 3

## **SECTION 15 REGULATORY INFORMATION**

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

# ACETONITRILE(75-05-8) IS FOUND ON THE FOLLOWING REGULATORY LISTS

Australia Exposure Standards
Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals
Australia Inventory of Chemical Substances (AICS)

Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Appendix E (Part 2)

Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Appendix J (Part 2)

# 6-ACETYLMORPHINE(2784-73-8) IS FOUND ON THE FOLLOWING REGULATORY LISTS

Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule 8

### **National Inventory Status**

National Inventory	Status
Australia - AICS	N (6-acetylmorphine)
Canada - DSL	N (6-acetylmorphine)
Canada - NDSL	N (acetonitrile; 6-acetylmorphine)
China - IECSC	N (6-acetylmorphine)
Europe - EINEC / ELINCS / NLP	N (6-acetylmorphine)
Japan - ENCS	N (6-acetylmorphine)
Korea - KECI	N (6-acetylmorphine)
New Zealand - NZIoC	N (6-acetylmorphine)
Philippines - PICCS	N (6-acetylmorphine)
USA - TSCA	N (6-acetylmorphine)
Legend:	Y = All ingredients are on the inventory N = Not determined or one or more ingredients are not on the inventory and are not exempt from listing(see specific ingredients in brackets)

Chemwatch: 9-463824 Page 10 of 10 Issue Date: 10/09/2018 Version No: 2.4 Print Date: 10/09/2018

# 6-Acetylmorphine-D3

### **SECTION 16 OTHER INFORMATION**

Revision Date	10/09/2018
Initial Date	05/10/2017

### Other information

Classification of the preparation and its individual components has drawn on official and authoritative sources as well as independent review by the Chemwatch Classification committee using available literature references.

The SDS is a Hazard Communication tool and should be used to assist in the Risk Assessment. Many factors determine whether the reported Hazards are Risks in the workplace or other settings. Risks may be determined by reference to Exposures Scenarios. Scale of use, frequency of use and current or available engineering controls must be considered.

### Definitions and abbreviations

 ${\sf PC-TWA: Permissible \ Concentration-Time \ Weighted \ Average}$ 

PC-STEL: Permissible Concentration-Short Term Exposure Limit

IARC: International Agency for Research on Cancer

ACGIH: American Conference of Governmental Industrial Hygienists

STEL: Short Term Exposure Limit

TEEL: Temporary Emergency Exposure Limit。

IDLH: Immediately Dangerous to Life or Health Concentrations

OSF: Odour Safety Factor

NOAEL :No Observed Adverse Effect Level LOAEL: Lowest Observed Adverse Effect Level

TLV: Threshold Limit Value LOD: Limit Of Detection OTV: Odour Threshold Value BCF: BioConcentration Factors BEI: Biological Exposure Index

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