

# **Dynapac Compaction Equipment**

Chemwatch: **5318-59** Version No: **4.1** Material Safety Data Sheet according to NOHSC and ADG requirements Issue Date: 23/12/2022 Print Date: 25/07/2023 L.Local.AUS.EN.E

### SECTION 1 Identification of the substance / mixture and of the company / undertaking

#### **Product Identifier**

Product name	Dynapac Paver Grease (Paver Grease)
Chemical Name	Not Applicable
Synonyms	Not Available
Chemical formula	Not Applicable
Other means of identification	Not Available

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

Relevant identified uses	Automotive and industrial grease.

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

Registered company name	Dynapac Compaction Equipment
Address	Box 504 Karlskrona SE-371 23 Sweden
Telephone	+46 455 30 60 00
Fax	+46 455 30 60 30
Website	http://www.dynapac.com
Email	info@dynapac.com

## Emergency telephone number

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Association / Organisation	CHEMWATCH EMERGENCY RESPONSE (24/7)
Emergency telephone numbers	+61 1800 951 288
Other emergency telephone numbers	+61 3 9573 3188

Once connected and if the message is not in your preferred language then please dial 01

#### **SECTION 2 Hazards identification**

#### Classification of the substance or mixture

Poisons Schedule	Not Applicable
Risk Phrases <sup>[1]</sup>	Not Applicable
Legend:	1. Classified by Chernwatch; 2. Classification drawn from HCIS; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI

#### Not Applicable

Relevant risk statements are found in section 2

Indication(s) of danger Not Applicable

#### Safety advice

Not Applicable

Other hazards

#### **SECTION 3 Composition / information on ingredients**

# Substances

See section below for composition of Mixtures

#### Mixtures

CAS No	%[weight]	Name
68411-46-1	<3	octylated diphenylamines
Not Available	0.1-0.9	bis(2-methylpentan-2-yl)dithiophosphoric acid/ amines

CAS No	%[weight]	Name
Not Available	NotSpec	polyolefins
Not Available	NotSpec	synthetic esters
Legend:	1. Classified by Chemwatch; Classification drawn from C&	<ol> <li>Classification drawn from HCIS; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 4.</li> <li>L; * EU IOELVs available</li> </ol>

# **SECTION 4 First aid measures**

Description of first aid measures	
Eye Contact	If this product comes in contact with eyes: <ul> <li>Wash out immediately with water.</li> <li>If irritation continues, seek medical attention.</li> <li>Removal of contact lenses after an eye injury should only be undertaken by skilled personnel.</li> </ul>
Skin Contact	<ul> <li>If skin or hair contact occurs:</li> <li>Flush skin and hair with running water (and soap if available).</li> <li>Seek medical attention in event of irritation.</li> <li>If failure/misuse of high pressure/hydraulic equipment results in injection of grease/oil through the skin seek urgent medical attention. Treat as surgical emergency.</li> </ul>
Inhalation	<ul> <li>If fumes, aerosols or combustion products are inhaled remove from contaminated area.</li> <li>Other measures are usually unnecessary.</li> </ul>
Ingestion	<ul> <li>Immediately give a glass of water.</li> <li>First aid is not generally required. If in doubt, contact a Poisons Information Centre or a doctor.</li> </ul>

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

Treat symptomatically.

# **SECTION 5 Firefighting measures**

#### Extinguishing media

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

Do not use water jets.

#### Special hazards arising from the substrate or mixture

Fire Incompatibility Avoid contamination with oxidising agents i.e. nitrates, oxidising acids, chlorine bleaches, pool chlorine etc. as ignition may result Advice for firefighters Alert Fire Brigade and tell them location and nature of hazard. Wear breathing apparatus plus protective gloves. Prevent, by any means available, spillage from entering drains or water courses. Use water delivered as a fine spray to control fire and cool adjacent area. Fire Fighting 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. Combustible. Slight fire hazard when exposed to heat or flame. Heating may cause expansion or decomposition leading to violent rupture of containers. On combustion, may emit toxic fumes of carbon monoxide (CO). May emit acrid smoke. Fire/Explosion Hazard Mists containing combustible materials may be explosive. Combustion products include: carbon dioxide (CO2) other pyrolysis products typical of burning organic material. May emit poisonous fumes. HAZCHEM Not Applicable

## SECTION 6 Accidental release measures

#### Personal precautions, protective equipment and emergency procedures

Minor Spills	<ul> <li>Clean up all spills immediately.</li> <li>Avoid contact with skin and eyes.</li> <li>Wear impervious gloves and safety goggles.</li> <li>Trowel up/scrape up.</li> <li>Place spilled material in clean, dry, sealed container.</li> <li>Flush spill area with water.</li> <li>Slippery when spilt.</li> </ul>
Major Spills	<ul> <li>Minor hazard.</li> <li>Clear area of personnel.</li> <li>Alert Fire Brigade and tell them location and nature of hazard.</li> <li>Control personal contact with the substance, by using protective equipment as required.</li> <li>Prevent spillage from entering drains or water ways.</li> <li>Contain spill with sand, earth or verniculite.</li> </ul>

<ul> <li>Collect recoverable product into labelled containers for recycling.</li> <li>Absorb remaining product with sand, earth or vermiculite and place in appropriate containers for disposal.</li> <li>Wash area and prevent runoff into drains or waterways.</li> <li>If contamination of drains or waterways occurs, advise emergency services.</li> <li>Slippery when spilt.</li> </ul>
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Personal Protective Equipment advice is contained in Section 8 of the SDS.

# **SECTION 7 Handling and storage**

Precautions for safe handling	
Safe handling	<ul> <li>Avoid all personal contact, including inhalation.</li> <li>Wear protective clothing when risk of exposure occurs.</li> <li>Use in a well-ventilated area.</li> <li>Prevent concentration in hollows and sumps.</li> <li>DO NOT enter confined spaces until atmosphere has been checked.</li> <li>DO NOT allow material to contact humans, exposed food or food utensils.</li> <li>Avoid contact with incompatible materials.</li> <li>When handling, DO NOT eat, drink or smoke.</li> <li>Keep containers securely sealed when not in use.</li> <li>Avoid physical damage to containers.</li> <li>Always wash hands with soap and water after handling.</li> <li>Work clothes should be laundered separately. Launder contaminated clothing before re-use.</li> <li>Use good occupational work practice.</li> <li>Observe manufacturer's storage and handling recommendations contained within this SDS.</li> <li>Atmosphere should be regularly checked against established exposure standards to ensure safe working conditions are maintained.</li> </ul>
Other information	<ul> <li>Store in original containers.</li> <li>Keep containers securely sealed.</li> <li>No smoking, naked lights or ignition sources.</li> <li>Store in a cool, dry, well-ventilated area.</li> <li>Store away from incompatible materials and foodstuff containers.</li> <li>Protect containers against physical damage and check regularly for leaks.</li> <li>Observe manufacturer's storage and handling recommendations contained within this SDS.</li> </ul>

## Conditions for safe storage, including any incompatibilities

Suitable container	<ul> <li>Metal can or drum</li> <li>Packaging as recommended by manufacturer.</li> <li>Check all containers are clearly labelled and free from leaks.</li> </ul>
Storage incompatibility	Avoid reaction with oxidising agents

# **SECTION 8 Exposure controls / personal protection**

## **Control parameters**

# Occupational Exposure Limits (OEL)

INGREDIENT DATA

#### Not Available

Emergency Limits

Ingredient	TEEL-1	TEEL-2		TEEL-3
Dynapac Paver Grease (Paver Grease)	Not Available	Not Available		Not Available
Ingredient	Original IDLH		Revised IDLH	
octylated diphenylamines	Not Available		Not Available	
bis(2-methylpentan- 2-yl)dithiophosphoric acid/ amines	Not Available		Not Available	
Occupational Exposure Banding	9			
Ingredient	Occupational Exposure Band Rating		Occupational Expos	ure Band Limit
octylated diphenylamines	D		> 0.01 to ≤ 0.1 mg/m <sup>3</sup>	
bis(2-methylpentan- 2-yl)dithiophosphoric acid/ amines	E		≤ 0.01 mg/m³	
Notes:	Occupational exposure banding is a process of adverse health outcomes associated with expo range of exposure concentrations that are expe	sure. The output of this p	rocess is an occupational	

### MATERIAL DATA

## 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. Ventilation can remove or dilute an air contaminant if designed properly. The design of a
	adds and removes an in the work environment, ventilation can remove of dilute an air contaminant if designed propeny. The design of a

	General exhaust is adequate under normal operating conditi essential to obtain adequate protection. Provide adequate very workplace possess varying "escape" velocities which, in turn remove the contaminant.	entilation in warehouse or closed storage areas. Air contamin	nants generated in the
	Type of Contaminant:		Air Speed:
	solvent, vapours, degreasing etc., evaporating from tank (in still air)		0.25-0.5 m/s (50-100 f/min)
	aerosols, fumes from pouring operations, intermittent container filling, low speed conveyer transfers, welding, spray drift, plating acid fumes, pickling (released at low velocity into zone of active generation) direct spray, spray painting in shallow booths, drum filling, conveyer loading, crusher dusts, gas discharge (active generation into zone of rapid air motion)		0.5-1 m/s (100-200 f/min.)
			1-2.5 m/s (200-500 f/min)
	grinding, abrasive blasting, tumbling, high speed wheel ge very high rapid air motion).	nerated dusts (released at high initial velocity into zone of	2.5-10 m/s (500-2000 f/min.)
	Within each range the appropriate value depends on:		
	Lower end of the range	Upper end of the range	
	1: Room air currents minimal or favourable to capture	1: Disturbing room air currents	
	2: Contaminants of low toxicity or of nuisance value only	2: Contaminants of high toxicity	
	3: Intermittent, low production.	3: High production, heavy use	
	4: Large hood or large air mass in motion	4: Small hood - local control only	
		in a tank 2 meters distant from the extraction point. Other m	
Individual protection measures, such as personal	considerations, producing performance deficits within the ex factors of 10 or more when extraction systems are installed of	traction apparatus, make it essential that theoretical air veloc	
•	considerations, producing performance deficits within the ex	traction apparatus, make it essential that theoretical air veloc	
measures, such as personal	<ul> <li>considerations, producing performance deficits within the exfactors of 10 or more when extraction systems are installed of the system of 10 or more when extraction systems are installed of the system of 10 or more when extraction systems are installed of the system of 10 or more when extraction systems are installed of the system of 10 or more when extraction systems are installed of the system of 10 or more when extraction systems are installed of the system of 10 or more when extraction systems are installed of the system of 10 or more when extractions on use, should be cand adsorption for the class of chemicals in use and an their removal and suitable equipment should be readily a remove contact lens as soon as practicable. Lens should</li> </ul>	raction apparatus, make it essential that theoretical air veloc or used.	r document, describing ew of lens absorption should be trained in tion immediately and ens should be removed
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measures, such as personal protective equipment	<ul> <li>considerations, producing performance deficits within the exfactors of 10 or more when extraction systems are installed of the system of 10 or more when extraction systems are installed of the system of 10 or more when extraction systems are installed of the system of 10 or more when extraction systems are installed of the system of 10 or more when extraction systems are installed of the system of 10 or more when extraction systems are installed of the system of 10 or more when extractions of the system of 10 or more when extractions of the system of 10 or more when extractions on use, should be called a desorption for the class of chemicals in use and an their removal and suitable equipment should be readily a remove contact lens as soon as practicable. Lens should a clean environment only after workers have washed had the extraction of the system of the sys</li></ul>	raction apparatus, make it essential that theoretical air veloc or used. equivalent] lenses may absorb and concentrate irritants. A written policy reated for each workplace or task. This should include a revi account of injury experience. Medical and first-aid personnel available. In the event of chemical exposure, begin eye irriga d be removed at the first signs of eye redness or irritation - le	r document, describing ew of lens absorption should be trained in tion immediately and ens should be removed
measures, such as personal protective equipment	<ul> <li>considerations, producing performance deficits within the exfactors of 10 or more when extraction systems are installed of the system of the period of the system and the system are installed of the system of the syste</li></ul>	raction apparatus, make it essential that theoretical air veloc or used. equivalent] lenses may absorb and concentrate irritants. A written policy reated for each workplace or task. This should include a revi account of injury experience. Medical and first-aid personnel available. In the event of chemical exposure, begin eye irriga d be removed at the first signs of eye redness or irritation - le	y document, describing ew of lens absorption should be trained in tion immediately and ens should be removed

#### **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.
- Cartridge performance is affected by humidity. Cartridges should be changed after 2 hr of continuous use unless it is determined that the humidity is less than 75%, in which case, cartridges can be used for 4 hr. Used cartridges should be discarded daily, regardless of the length of time used

# **SECTION 9** Physical and chemical properties

#### Information on basic physical and chemical properties

Appearance	Light brown paste with slight hydrocarbon odour; does not mix with water.		
Physical state	Non Slump Paste	Relative density (Water = 1)	1.0 @15C
Odour	Not Available	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	>320
pH (as supplied)	Not Applicable	Decomposition temperature (°C)	Not Available

Initial boiling point and boiling range (°C)Not AvailableMolecular weight (g/mol)Not ApplicableFlash point (°C)Not AvailableTasteNot AvailableEvaporation rateNot AvailableExplosive propertiesNot AvailableFlammabilityNot AvailableOxidising propertiesNot AvailableUpper Explosive Limit (%)1Surface Tension (dyn/cm or mN/m)Not AvailableLower Explosive Limit (%)1Volatile Component (%vol)NegligibleVapour pressure (kPa)<0.0005 @20CGas groupNot AvailableSolubility in wateImmisciblepH as a solution (1%)Not Applicable	Melting point / freezing point (°C)	250 (pour pt.)	Viscosity (cSt)	Not Available
Evaporation rateNot AvailableExplosive propertiesNot AvailableFlammabilityNot AvailableOxidising propertiesNot AvailableUpper Explosive Limit (%)10Surface Tension (dyn/cm or mN/m)Not AvailableLower Explosive Limit (%)1Volatile Component (%vol)NegligibleVapour pressure (kPa)<0.0005 @20CGas groupNot AvailableSolubility in waterImmisciblePH as a solution (1%)Not Applicable	•. •	Not Available	Molecular weight (g/mol)	Not Applicable
FlammabilityNot AvailableOxidising propertiesNot AvailableUpper Explosive Limit (%)10Surface Tension (dyn/cm or mN/m)Not AvailableLower Explosive Limit (%)1Volatile Component (%vol)NegligibleVapour pressure (kPa)<0.0005 @20CGas groupNot AvailableSolubility in waterImmisciblePH as a solution (1%)Not Applicable	Flash point (°C)	Not Available	Taste	Not Available
Upper Explosive Limit (%)       10       Surface Tension (dyn/cm or mN/m)       Not Available         Lower Explosive Limit (%)       1       Volatile Component (%vol)       Negligible         Vapour pressure (kPa)       <0.0005 @20C       Gas group       Not Available         Solubility in water       Immiscible       pH as a solution (1%)       Not Applicable	Evaporation rate	Not Available	Explosive properties	Not Available
Upper Explosive Limit (%)     10     Not Available       Lower Explosive Limit (%)     1     Volatile Component (%vol)     Negligible       Vapour pressure (kPa)     <0.0005 @20C     Gas group     Not Available       Solubility in water     Immiscible     pH as a solution (1%)     Not Applicable	Flammability	Not Available	Oxidising properties	Not Available
Vapour pressure (kPa)     <0.0005 @20C     Gas group     Not Available       Solubility in water     Immiscible     pH as a solution (1%)     Not Applicable	Upper Explosive Limit (%)	10		Not Available
Solubility in water         Immiscible         pH as a solution (1%)         Not Applicable	Lower Explosive Limit (%)	1	Volatile Component (%vol)	Negligible
	Vapour pressure (kPa)	<0.0005 @20C	Gas group	Not Available
	Solubility in water	Immiscible	pH as a solution (1%)	Not Applicable
VOC g/L   Not Available	Vapour density (Air = 1)	>1	VOC g/L	Not Available

# **SECTION 10 Stability and reactivity**

Reactivity	See section 7
Chemical stability	<ul> <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**

Inhaled	The material is not thought to produce adverse health effects or irritation of the respiratory tract (as classified by EC Directives using animal models). Nevertheless, good hygiene practice requires that exposure be kept to a minimum and that suitable control measures be used in an occupational setting.		
Ingestion	Although ingestion is not thought to produce harmful effects (as classified under EC Directives), the material may still be damaging to the health of the individual, following ingestion, especially where pre-existing organ (e.g liver, kidney) damage is evident. Present definitions of harmful or toxic substances are generally based on doses producing mortality rather than those producing morbidity (disease, ill-health). Gastrointestinal tract discomfort may produce nausea and vomiting. In an occupational setting however, ingestion of insignificant quantities is not thought to be cause for concern.		
Skin Contact	dermatitis is often characterised by skin redness (erytherr spongy layer (spongiosis) and intracellular oedema of the Open cuts, abraded or irritated skin should not be expose	d to this material prasions, puncture wounds or lesions, may produce systemic injury with harmful effec	
Eye	Although the material is not thought to be an irritant (as cl characterised by tearing or conjunctival redness (as with	lassified by EC Directives), direct contact with the eye may produce transient discomf windburn).	
Chronic		cupational exposure may produce cumulative health effects involving organs or	
	biochemical systems.		
	biochemical systems. TOXICITY	IRRITATION	
ynapac Paver Grease (Paver	· · · · · · · · · · · · · · · · · · ·	IRRITATION Not Available	
ynapac Paver Grease (Paver Grease)	ΤΟΧΙΟΙΤΥ		
•••	TOXICITY Dermal (Rabbit) LD50: >5000 mg/kg* <sup>[2]</sup>		
•••	TOXICITY           Dermal (Rabbit) LD50: >5000 mg/kg* <sup>[2]</sup> Oral (Rat) LD50: >5000 mg/kg* <sup>[2]</sup>	Not Available	
•••	TOXICITY           Dermal (Rabbit) LD50: >5000 mg/kg* <sup>[2]</sup> Oral (Rat) LD50: >5000 mg/kg* <sup>[2]</sup> TOXICITY	Not Available IRRITATION	
Grease)	TOXICITY           Dermal (Rabbit) LD50: >5000 mg/kg*[2]           Oral (Rat) LD50: >5000 mg/kg*[2]           TOXICITY           dermal (rat) LD50: >2000 mg/kg <sup>[1]</sup>	Not Available       IRRITATION       Eye (rabbit): Non Irritant	
Grease)	TOXICITY           Dermal (Rabbit) LD50: >5000 mg/kg*[2]           Oral (Rat) LD50: >5000 mg/kg*[2]           TOXICITY           dermal (rat) LD50: >2000 mg/kg <sup>[1]</sup>	Not Available         IRRITATION         Eye (rabbit): Non Irritant         Eye: adverse effect observed (irritating) <sup>[1]</sup>	
Grease)	TOXICITY           Dermal (Rabbit) LD50: >5000 mg/kg*[2]           Oral (Rat) LD50: >5000 mg/kg*[2]           TOXICITY           dermal (rat) LD50: >2000 mg/kg <sup>[1]</sup>	Not Available         IRRITATION         Eye (rabbit): Non Irritant         Eye: adverse effect observed (irritating) <sup>[1]</sup> Skin (rabbit): Non Irritant [Bay]	

OCTYLATED DIPHENYLAMINES For substituted diphenylamines: Based upon reviewed data the physicochemical and toxicological properties of the substituted diphenylamines are similar and follow a regular pattern as a result of that structural similarity.

		eir common starting material Diphenylamine are
	Because of their powerful antioxidant properties, Substituted Diphenylamines, along with th regulated for use in several food-contact applications by the Food and Drug Administration sections of the Code of Federal Regulations (CFR): Heating may generate vapors which can irritate the eyes and respiratory passages. Drying may be possible from prolonged or repeated contact. Overexposure to vapors from heating respiratory tract irritation with symptoms such as, but not limited to, dizziness and flu-like sy <b>Acute toxicity</b> : As a group these materials do not produce significant acute toxicity in mam following oral administration, with LD50 values ranging from >5000 to > 34,000 mg/kg. Ove greater than the 2000 mg/kg limit dose indicating a very low order of toxicity. <b>Mammalian Toxicology - Mutagenicity</b> . Data from bacterial reverse mutation assays, in vi well as additional supporting in vitro and in vivo genetic toxicity studies indicate a low conce materials. Similarly, the data for a mixed aryl/alkyl substituted molecule also indicates a lack Potential sensitiser producing contact allergies.	as Indirect Food Additives under the following of skin and mucous membranes leading to irritation the product may cause and/or skin irritation and ymptoms mals. All show a slight to very low order of toxicity rall, the acute dermal LD50 for these materials was itro and in vivo chromosome aberration studies, as ern for mutagenicity either for aryl or alkyl substituted
	Thee rat oral LD50 is greater than 10 ml/kg bw. No mortality occurred. No signs of systemic the study, and no abnormalities were noted at necropsy. In a second study this substance s with OECD 401. The rat oral LD50is estimated at approximately 2000 mg/kg.No gross path organ toxicity is evident. The dermal route for acute toxicity is appropriate if the physicocher significant rate of absorption through the skin. The scientific literature regarding dermal toxic greater than 5 there is very limited potential for dermal absorption (e.g., 10% absorption) (A The test material has a Log Kow greater than 7.1 (small portion < 0.3) thereby demonstratir In contrast, oral absorption can be relatively fast due to contact surface areas in the GI tract tract has been regarded as the route resulting in higher bioavailability. For the test material, scenario for assessing acute exposure, and acute dermal toxicity end point satisfied by acu	shows evidence of toxicity when tested in accordance ology was found in surviving animals.No specific mical properties suggest there is potential for a city states that for those substances with a log Kow nnals of Occupatinoal Hygiene, 47(8):641-652, 2003 ng that it has very limited dermal absorption potential t resulting in a peak concentration in the body, and G dosing via oral route represents the worst case
BIS(2-METHYLPENTAN- 2-YL)DITHIOPHOSPHORIC ACID/ AMINES	sensitisation: EC3 value was determined to be 9.39%. Per the CLP guidance, substances a EC3 value is less than 2% and are to be classified as skin sensitization 1B when the EC3 v. administration of the test substance to rats by gavage in accordance with OECD Test Guide microscopic changes in the adrenal glands of the male and female rats and kidneys of the r adrenal gland changes are accompanied by an increase in adrenal weight only at the high or by an increase in hyaline droplets which is consistent with male rat species specific effect re a2-microglobulin in renal proximal tubular epithelial cells. Microscopic changes also are pre 500 mg/kg/day group and these changes were possibly treatment related. Landing foot spla high-dose females, and rectal temperature was also statistically significantly decreased in the splay was not increased and there were no statistically significant differences in mean moto parameters, this is not considered to be evidence of serious adverse effects. Therefore, the study is 150 mg/kg/day, and the no observable effect level (NOEL) is 50 mg/kg/day. * REAC identified in literature search.	alue is greater than 2%. Repeat dose toxicity: Oral eline 407 (1995) produces treatment related male rats of the 150 and 500 mg/kg/day groups. The doses level. The male kidney effects are accompanie esulting from the excessive accumulation of usent in the stomach of the male and female rats of th ay was statistically significantly decreased in mid- an he high-dose females. However, because landing for or activity data or other functional observational batte e no observable adverse effect level (NOAEL) for this
2-YL)DITHIOPHOSPHORIC	EC3 value is less than 2% and are to be classified as skin sensitization 1B when the EC3 v administration of the test substance to rats by gavage in accordance with OECD Test Guide microscopic changes in the adrenal glands of the male and female rats and kidneys of the r adrenal gland changes are accompanied by an increase in adrenal weight only at the high by an increase in hyaline droplets which is consistent with male rat species specific effect r a2-microglobulin in renal proximal tubular epithelial cells. Microscopic changes also are pre 500 mg/kg/day group and these changes were possibly treatment related. Landing foot spla high-dose females, and rectal temperature was also statistically significantly decreased in tt splay was not increased and there were no statistically significant differences in mean moto parameters, this is not considered to be evidence of serious adverse effects. Therefore, the study is 150 mg/kg/day, and the no observable effect level (NOEL) is 50 mg/kg/day. * REAC	alue is greater than 2%. Repeat dose toxicity: Oral eline 407 (1995) produces treatment related male rats of the 150 and 500 mg/kg/day groups. The doses level. The male kidney effects are accompanie esulting from the excessive accumulation of esent in the stomach of the male and female rats of the ay was statistically significantly decreased in mid- an he high-dose females. However, because landing for or activity data or other functional observational battle en observable adverse effect level (NOAEL) for this Ch Dossier No significant acute toxicological data this product. or Quincke's oedema. The pathogenesis of contact her allergic skin reactions, e.g. contact urticaria, t simply determined by its sensitisation potential: the A weakly sensitising substance which is widely th which few individuals come into contact. From a
2-YL)DITHIOPHOSPHORIC ACID/ AMINES OCTYLATED DIPHENYLAMINES & BIS(2- METHYLPENTAN- 2-YL)DITHIOPHOSPHORIC ACID/ AMINES	EC3 value is less than 2% and are to be classified as skin sensitization 1B when the EC3 v administration of the test substance to rats by gavage in accordance with OECD Test Guide microscopic changes in the adrenal glands of the male and female rats and kidneys of the r adrenal gland changes are accompanied by an increase in adrenal weight only at the high of by an increase in hyaline droplets which is consistent with male rat species specific effect re a2-microglobulin in renal proximal tubular epithelial cells. Microscopic changes also are pre 500 mg/kg/day group and these changes were possibly treatment related. Landing foot spla high-dose females, and rectal temperature was also statistically significantly decreased in tt splay was not increased and there were no statistically significent differences in mean moto parameters, this is not considered to be evidence of serious adverse effects. Therefore, the study is 150 mg/kg/day, and the no observable effect level (NOEL) is 50 mg/kg/day. * REAC identified in literature search. The following information refers to contact allergens as a group and may not be specific to th Contact allergies quickly manifest themselves as contact eczema, more rarely as urticaria of eczema involves a cell-mediated (T lymphocytes) immune reaction of the delayed type. Oth involve antibody-mediated immune reactions. The significance of the contact allergen is not distribution of the substance and the opportunities for contact with it are equally important. <i>J</i> distributed can be a more important allergen than one with stronger sensitising potential wit clinical point of view, substances are noteworthy if they produce an allergic test reaction in re- distributed can be a more important allergen than one with stronger sensitising potential wit clinical point of view, substances are noteworthy if they produce an allergic test reaction in re-	alue is greater than 2%. Repeat dose toxicity: Oral eline 407 (1995) produces treatment related male rats of the 150 and 500 mg/kg/day groups. The doses level. The male kidney effects are accompanie esulting from the excessive accumulation of esent in the stomach of the male and female rats of th ay was statistically significantly decreased in mid- an he high-dose females. However, because landing for or activity data or other functional observational batte e no observable adverse effect level (NOAEL) for this Ch Dossier No significant acute toxicological data this product. or Quincke's oedema. The pathogenesis of contact ner allergic skin reactions, e.g. contact urticaria, t simply determined by its sensitisation potential: the A weakly sensitising substance which is widely th which few individuals come into contact. From a more than 1% of the persons tested.
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X - Data either not available or does not fill the criteria for classification Data available to make classification

# **SECTION 12 Ecological information**

D	Endpoint	Test Duration (hr)	Species	Value	Source
Dynapac Paver Grease (Paver Grease)	Not Available	Not Available	Not Available	Not Available	Not Availabl
	Endpoint	Test Duration (hr)	Species	Value	Source
	EC50	72h	Algae or other aquatic plants	>100mg/l	2
octylated diphenylamines	EC50	48h	Crustacea	51mg/l	2
	EC50	96h	Algae or other aquatic plants	870mg/l	2
	LC50	96h	Fish	5.1mg/l	Not Availab
	EC50(ECx)	24h	Crustacea	4.2mg/l	Not Availabl
bis(2-methylpentan-	Endpoint	Test Duration (hr)	Species	Value	Source
2-yl)dithiophosphoric acid/ amines	Not Available	Not Available	Not Available	Not Available	Not Availabl
	Available Extracted from	1. IUCLID Toxicity Data 2. Europe E	Not Available CHA Registered Substances - Ecotoxicological Informatic C Aquatic Hazard Assessment Data 6. NITE (Japan) - Bic	Available on - Aquatic Toxicity 4.	US

### DO NOT discharge into sewer or waterways.

#### Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
octylated diphenylamines	HIGH	HIGH
Bioaccumulative potential		
Ingredient	Bioaccumulation	
octylated diphenylamines	LOW (BCF = 5.5)	
Mobility in soil		
Ingredient	Mobility	
octylated diphenylamines	LOW (KOC = 28640000)	

## **SECTION 13 Disposal considerations**

Waste treatment methods	
Product / Packaging disposal	<ul> <li>DO NOT allow wash water from cleaning or process equipment to enter drains.</li> <li>It may be necessary to collect all wash water for treatment before disposal.</li> <li>In all cases disposal to sewer may be subject to local laws and regulations and these should be considered first.</li> <li>Where in doubt contact the responsible authority.</li> <li>Recycle wherever possible or consult manufacturer for recycling options.</li> <li>Consult State Land Waste Authority for disposal.</li> <li>Bury or incinerate residue at an approved site.</li> <li>Recycle containers if possible, or dispose of in an authorised landfill.</li> </ul>

## **SECTION 14 Transport information**

#### Labels Required

Marine Pollutant	NO
HAZCHEM	Not Applicable

Land transport (ADG): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

Air transport (ICAO-IATA / DGR): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

Sea transport (IMDG-Code / GGVSee): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

Transport in bulk according to Annex II of MARPOL and the IBC code Not Applicable

#### Transport in bulk in accordance with MARPOL Annex V and the IMSBC Code

Product name	Group
octylated diphenylamines	Not Available
bis(2-methylpentan- 2-yl)dithiophosphoric acid/ amines	Not Available

#### Transport in bulk in accordance with the IGC Code

Product name	Ship Type
octylated diphenylamines	Not Available
bis(2-methylpentan- 2-yl)dithiophosphoric acid/ amines	Not Available

## **SECTION 15 Regulatory information**

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

# octylated diphenylamines is found on the following regulatory lists

Australian Inventory of Industrial Chemicals (AIIC)

nternational WHO List of Proposed Occupational Exposure Limit (OEL) Values for
Manufactured Nanomaterials (MNMS)

bis(2-methylpentan-2-yl)dithiophosphoric acid/ amines is found on the following reg	gulatory lists
Not Applicable	

#### **National Inventory Status**

National Inventory	Status
Australia - AIIC / Australia Non-Industrial Use	Yes
Canada - DSL	Yes

National Inventory	Status			
Canada - NDSL	No (octylated diphenylamines)			
China - IECSC	/es			
Europe - EINEC / ELINCS / NLP	Yes			
Japan - ENCS	Yes			
Korea - KECI	Yes			
New Zealand - NZIoC	Yes			
Philippines - PICCS	Yes			
USA - TSCA	Yes			
Taiwan - TCSI	Yes			
Mexico - INSQ	Yes			
Vietnam - NCI	Yes			
Russia - FBEPH	Yes			
Legend:	Yes = All CAS declared ingredients are on the inventory No = One or more of the CAS listed ingredients are not on the inventory. These ingredients may be exempt or will require registration.			

### **SECTION 16 Other information**

Revision Date	23/12/2022
Initial Date	09/08/2018

#### **SDS Version Summary**

Version	Date of Update	Sections Updated
3.1	01/11/2019	One-off system update. NOTE: This may or may not change the GHS classification
4.1	23/12/2022	Classification review due to GHS Revision change.

#### 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

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 ES: Exposure Standard 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 AIIC: Australian Inventory of Industrial Chemicals DSL: Domestic Substances List NDSL: Non-Domestic Substances List IECSC: Inventory of Existing Chemical Substance in China EINECS: European INventory of Existing Commercial chemical Substances ELINCS: European List of Notified Chemical Substances NLP: No-Longer Polymers ENCS: Existing and New Chemical Substances Inventory KECI: Korea Existing Chemicals Inventory NZIoC: New Zealand Inventory of Chemicals PICCS: Philippine Inventory of Chemicals and Chemical Substances TSCA: Toxic Substances Control Act TCSI: Taiwan Chemical Substance Inventory INSQ: Inventario Nacional de Sustancias Químicas NCI: National Chemical Inventory FBEPH: Russian Register of Potentially Hazardous Chemical and Biological Substances This document is copyright.

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