

Product no. 71A/7110-01A Product name CHLORPYRIFOS 480 g/I EC Cheminova A/S P.O. Box 9 DK-7620 Lemvig Denmark Phone (+45) 96 90 96 90 Fax (+45) 96 90 96 91 www.cheminova.com CVR-No. DK 12 76 00 43

December 2012 Supersedes December 2010

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SAFETY DATA SHEET

CHLORPYRIFOS 480 g/I EC

Revision: Sections containing a revision or new information are marked with a .

♣ SECTION 1: IDENTIFICATION OF THE SUBSTANCE/MIXTURE AND OF THE COMPANY/UNDERTAKING

1.1.	Product identifier	CHLORPYRIFOS 480 g/I EC Contains: chlorpyrifos and solvent naphtha (petroleum), heavy aromatic
	Trade names	Chlorpyrifos 4E, Cyren, Nufos
1.2.	Relevant identified uses of the substance or mixture and uses advised against	Can be used as insecticide only.
1.3.	Details of the supplier of the safety data sheet	CHEMINOVA A/S P.O. Box 9 DK-7620 Lemvig Denmark sds@cheminova.dk
1.4.	Emergency telephone number	(+45) 97 83 53 53 (24 h; for emergencies only)
♣ SEC	TION 2: HAZARDS IDENTIFICATION	
2.1.	Classification of the substance or mixture	See section 16 for full text of R-phrases and hazard statements.
	DPD classification of the product according to Dir. 1999/45/EC as amended	Carc3;R40 Xn;R20/22 Xi;R38 R65 N;R50/53
	CLP classification of the product according to Reg. 1272/2008 as amended	Acute oral toxicity: Category 3 (H301) Acute inhalation toxicity: Category 4 (H332) Eye irritation: Category 2 (H319) Carcinogenicity: Category 2 (H351) Aspiration toxicity: Category 1 (H304) Hazards to the aquatic environment: Acute Category 1 (H400) Chronic Category 1 (H410)
	WHO classification Guidelines to Classification 2009	Class II: Moderately hazardous
	Health hazards	The product is harmful by inhalation and by ingestion. It has irritating properties.
		Chlorpyrifos is a dangerous poison (cholinesterase inhibitor). It rapidly enters the body on contact with all skin surfaces and eyes. Exposed persons must receive prompt medical treatment.



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Repeated exposures to cholinesterase inhibitors such as **chlorpyrifos** may, without warning, cause increased susceptibility to doses of any cholinesterase inhibitor.

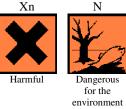
The product contains max. 5% of **naphthalene**, which is a suspected carcinogen.

Environmental hazards The product is very toxic to aquatic organisms.

2.2. Label elements

Product no.

According to Dir. 1999/45/EC as amended Hazard symbolsXr



Contains chlorpyrifos and solvent naphtha (petroleum), heavy aromatic

R-phrases	Timited evidence of a consistence is offered
R40	Limited evidence of a carcinogenic effect.
R20/22	Harmful by inhalation and if swallowed.
R38	Irritating to skin.
R65	Harmful: may cause lung damage if swallowed.
R50/53	Very toxic to aquatic organisms, may cause long-term adverse
	effects in the aquatic environment.
S-phrases	
S24	Avoid contact with skin.
\$36/37	Wear suitable protective clothing and gloves.
S46	If swallowed, seek medical advice immediately and show this container or label.
S60	This material and its container must be disposed of as hazardous waste.
S61	Avoid release to the environment. Refer to special instructions/safety data sheets.
Other mention	To avoid risks to man and the environment, comply with the instructions of use.

Additional phrases for final use of the product for plant protection

ruanional philases joi jinai use	of the product for plant protection
S2	. Keep out of the reach of children.
S13	. Keep away from food, drink and animal feedingstuffs.
S23	. Do not breathe spray.
S29	. Do not empty into drains.
SP1	. Do not contaminate water with the product or its container (Do not
	clean application equipment near surface water/Avoid
	contamination via drains from farmyards and roads).

<u>According to EU Reg. 1272/2008 as amended</u>		
Product identifier	Chlorpyrifos 480 g/l EC	
	Contains chlorpyrifos and solvent naphtha (petroleum), heavy	
	aromatic	



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Hazard pictograms (GHS06, GHS08, GHS09)



May be fatal if swallowed and enters airways.

Very toxic to aquatic life with long lasting effects.

Repeated exposure may cause skin dryness and cracking.

clean application equipment near surface water/Avoid

To avoid risks to human health and the environment, comply with

Do not contaminate water with the product or its container (Do not

Signal word

Hazard statements		
H301		
H332		
H319		
H351		
H304		
H410		

Supplementary hazard statements EUH066 EUH401

Supplementary phrase for final use of the product for plant protection: SP1

		contamination via drains from farmyards and roads).
	Precautionary statements	
	P261	Avoid breathing vapours.
	P273	Avoid release to the environment.
	P280	Wear eye/face protection.
	P305+P351+P338	IF IN EYES: Rinse cautiously with water for several minutes.
		Remove contact lenses, if present and easy to do. Continue rinsing.
	P310	Immediately call a POISON CENTER or doctor/physician.
	P501	Dispose of contents/container as hazardous waste.
2.3.	Other hazards	None of the ingredients in the product meets the criteria for being PBT or vPvB.

the instructions of use.

Danger

Toxic if swallowed. Harmful if inhaled.

Causes serious eye irritation.

Suspected of causing cancer.

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

3.1.	Substances	The product is a mixture, not a substance.
3.2.	Mixtures	See section 16 for full text of R-phrases and hazard statements.
	<u>Active ingredient</u> Chlorpyrifos CAS name	Content: 45% by weight Phosphorothioic acid, O,O-diethyl O-(3,5,6-trichloro-2-pyridinyl) ester
	CAS no IUPAC name ISO name/EU name EC no. (EINECS no.) EU index no DSD classification of the ingredient CLP classification of the ingredient	2921-88-2 O,O-Diethyl O-3,5,6-trichloro-2-pyridyl phosphorothioate Chlorpyrifos 220-864-4 015-084-00-4 T;R25 N;R50/53 Acute oral toxicity: Category 3 (H301) Hazards to the aquatic environment: Acute Category 1 (H400) Chronic Category 1 (H410)



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Structural formula CI _CI N $C_2H_5O_{$ C₂H₅O ĊI

Reportable ingredients

	Content (% w/w)	CAS no.	EC no. (EINECS no.)	DSD classification	CLP classification
Solvent naphtha (petroleum), heavy aromatic Reg. no. 01- 2119464588-24	51	64742-94-5	265-198-5	Carc3;R40 R65 R66 R67 N;R51/53 Harmful, dangerous for the environment	Carc. 2 (H351) STOT SE 3 (H336) Asp. Tox. 1 (H304) Aquatic Chronic 2 (H411)
Naphthalene	5	91-20-3	202-049-5	Carc3;R40 Xn;R22 N;R50/53 Harmful, dangerous for the environment	Carc. 2 (H351) Acute Tox. 4 (H302) Aquatic Acute 1 (H400) Aquatic Chronic 1 (H410)
Calcium dodecylbenzene sulphonate	1.5	26264-06-2	247-557-8	Xi;R38-41 N;R51/53 Irritant, dangerous for the environment	Skin Irrit. 2 (H315) Eye Dam. 1 (H318) Aquatic Chronic 2 (H411)
2-Ethylhexan-1-ol	1	104-76-7	203-234-3	Xi;R36 Irritant	Eye Irrit. 2 (H319)

SECTION 4: FIRST AID MEASURES

4.1.	Description of first aid measures	If exposure has occurred, do not wait for symptoms to develop, but immediately start the procedures described below.
	Inhalation	If exposure occurs, immediately remove from it. Light cases: Keep person under surveillance. Get medical attention immediately if symptoms develop. Serious cases: Get medical attention immediately or call for an ambulance.
		If breathing has stopped, immediately start artificial respiration and maintain until a physician takes charge of the exposed person.
	Skin contact	Immediately flush with much water while removing contaminated clothing and footwear. Wash with water and soap. See physician immediately if symptoms develop.
	Eye contact	Immediately rinse eyes with much water or eyewash solution, occasionally opening eyelids, until no evidence of chemical remains. Remove contact lenses after a few minutes and rinse again. See physician immediately.



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	Ingestion	 Call a doctor or get medical attention immediately. Make the exposed person rinse mouth and then drink 1 or 2 glasses of water or milk. Induce vomiting only if: 1. A significant amount (more than a mouthful) has been ingested 2. Patient is fully conscious 3. Medical aid is not readily available 4. Time since ingestion is less than one hour. Let the patient induce vomiting by touching the back of the throat with a finger. If vomiting occurs, take care that vomit does not enter airways. Let the exposed person rinse mouth and drink fluids again.
4.2.	Most important symptoms and effects, both acute and delayed	The first symptom to appear may be irritation. Symptoms of cholinesterase inhibition: nausea, headache, vomiting, cramps, weakness, blurred vision, pin-point pupils, tightness in chest, laboured breathing, nervousness, sweating, watering of eyes, drooling or frothing of mouth and nose, muscle spasms and coma.
4.3.	Indication of any immediate medical attention and special treatment needed	If any sign of cholinesterase inhibition occurs, call a doctor (physician), clinic or hospital immediately. Explain that the victim has been exposed to chlorpyrifos , an organophosphorus insecticide. Describe his/her condition and the extent of exposure. Immediately remove the exposed person from the area where the product is present.
		In an industrial setting the antidote atropine sulphate should be available at the workplace.
		It may be helpful to show this safety data sheet to physician.
	Notes to physician	Chlorpyrifos is a cholinesterase inhibitor affecting the central and peripheral nervous systems producing respiratory depression.
		The product contains petroleum distillates which may pose an aspiration pneumonia hazard.
	Cholinesterase inhibition – treatment	Much information on (acetyl)cholinesterase inhibition by organophosphate insecticides and its treatment can be found on the internet.
		Decontamination procedures such as whole body washing, gastric lavage and administration of activated charcoal are often required.
		Antidote : If symptoms (see subsection 4.2.) are present, administer atropine sulphate, which often is a lifesaving antidote, in large doses, TWO to FOUR mg intravenously or intramuscularly as soon as possible. Repeat at 5 to 10 minute intervals until signs of atropinisation appear and maintain full atropinisation until all organophosphate is metabolised.
		Obidoxime chloride (Toxogonin), alternatively pralidoxime chloride (2-PAM), may be administered as an adjunct to, but not a substitute for atropine sulphate. Treatment with oxime should be maintained as long as atropine sulphate is administered.



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At first sign of pulmonary oedema the patient should be given supplementary oxygen and treated symptomatically.

Relapse can occur after initial improvement. VERY CLOSE SUPERVISION OF THE PATIENT IS INDICATED FOR AT LEAST 48 HOURS, DEPENDING ON THE SEVERITY OF POISONING.

SECTION 5: FIREFIGHTING MEASURES

5.1.	Extinguishing media	Dry chemical or carbon dioxide for small fires, water spray or foam for large fires. Avoid heavy hose streams.
5.2.	Special hazards arising from the substance or mixture	The essential breakdown products are volatile, toxic, irritant, malodorous and inflammable compounds such as nitrogen oxides, hydrogen chloride, ethyl mercaptan, diethyl sulphide, sulphur dioxide, carbon monoxide, carbon dioxide, phosphorus pentoxide and various chlorinated organic compounds.
5.3.	Advice for firefighters	Use water spray to keep fire-exposed containers cool. Approach fire from upwind to avoid hazardous vapours and toxic decomposition products. Fight fire from protected location or maximum possible distance. Dike area to prevent water runoff. Firemen should wear self-contained breathing apparatus and protective clothing.

***** SECTION 6: ACCIDENTAL RELEASE MEASURES

6.1.	Personal precautions, protective equipment and emergency procedures	It is recommended to have a predetermined plan for the handling of spills. Empty, sealable vessels for the collection of spills should be available.
		In case of large spill (involving 10 tons of the product or more):1. Use personal protection equipment; see section 82. Call emergency telephone no.; see section 13. Alert authorities.
		Observe all safety precautions when cleaning up spills. Use personal protection equipment. Depending on the magnitude of the spill this may mean wearing respirator, face mask or eye protection, chemical resistant clothing, gloves and boots.
		Stop the source of the spill immediately if safe to do so. Keep unprotected persons away from the spill area. Remove sources of ignition. Avoid and reduce mist formation as much as possible.
6.2.	Environmental precautions	Contain the spill to prevent any further contamination of surface, soil or water. Wash waters must be prevented from entering surface water drains. Uncontrolled discharge into water courses must be alerted to the appropriate regulatory body.
6.3.	Methods and materials for containment and cleaning up	It is recommended to consider possibilities to prevent damaging effects of spills, such as bunding or capping. See GHS (Annex 4, Section 6).
		Use non-sparking tools and equipment. If appropriate, surface



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		water drains should be covered. Minor spills on the floor or other impervious surface should be absorbed onto an absorptive material such as universal binder, hydrated lime, bentonite or other absorbent clays. Collect the contaminated absorbent in suitable containers. Clean area with soda lye and much water. Absorb wash liquid onto absorbent and transfer to suitable containers. The used containers should be properly closed and labelled.
		Spills which soak into the ground should be dug up and transferred to suitable containers.
		Spills in water should be contained as much as possible by isolation of the contaminated water. The contaminated water must be collected and removed for treatment or disposal.
6.4.	Reference to other sections	See subsection 8.2. for personal protection. See section 13 for disposal.
SE	CTION 7: HANDLING AND STOR	AGE
7.1.	Precautions for safe handling	Keep away from sources of ignition and protect from exposure to fire and heat.
		In an industrial environment it is recommended to avoid all personal contact with the product, if possible by using closed systems with remote system control. Otherwise, the material should be handled by mechanical means as much as possible. Adequate ventilation or local exhaust ventilation is required. The exhaust gases should be filtered or treated otherwise. For personal protection in this situation, see section 8.
		For its use as a pesticide, first look for precautions and personal protection measures on the officially approved label on the packaging or for other official guidance or policy in force. If these are lacking, see section 8.
		Keep all unprotected persons and children away from working area.
		Remove contaminated clothing immediately. Wash thoroughly after handling. Before removing gloves, wash them with water and soap. After work, take off all work clothes and footwear. Take a shower, using water and soap. Wear only clean clothes when leaving job. Wash protective clothing and protective equipment with water and soap after each use. Clothes that have been heavily drenched must be discarded. Do not wash and reuse them.
		Inhalation of vapours of the product can cause lowered consciousness, which increases the risks of operating machinery and driving.
		The respirator should be cleaned and filter replaced according to the accompanying instructions.
		Do not discharge to the environment. Collect all waste material and remains from cleaning equipment, etc., and dispose of as hazardous

waste. See section 13 for disposal.



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7.2.	Conditions for safe storage, including any incompatibilities	The product is stable under normal conditions of warehouse storage. Protect against sunshine for prolonged periods.
		Keep in tightly closed, labelled containers. The storage room should be constructed of incombustible material, closed, dry, ventilated and with impermeable floor, without access of unauthorised persons or children. A warning sign reading "POISON" is recommended. The room should only be used for storage of chemicals. Food, drink, feed and seed should not be present. A hand wash station should be available.
7.3.	Specific end use(s)	The product is a registered pesticide which may only be used for the applications it is registered for, in accordance with a label approved by the regulatory authorities.

♣ SECTION 8: EXPOSURE CONTROLS/PERSONAL PROTECTION

8.1. **Control parameters**

Personal	exposure	limits
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		Year	
Chlorpyrifos	ACGIH (USA) TLV	2012	TWA 0.1 mg/m ³ , measured as inhalable fraction and vapour
			Skin notation; BEI
	OSHA (USA) PEL		Not established
	EU, 2000/39/EC	2009	Not established
	as amended	2012	
	Germany, MAK		Not established; BAT
	HSE (UK) WEL	2007	8-h TWA 0.2 mg/m ³
			STEL 0.6 mg/m ³ ; 15-minute reference period Skin notation
			Skill hotation
Naphthalene	ACGIH (USA) TLV	2012	TWA 10 ppm (52 mg/m ³)
- ··· F			Ceiling 15 ppm (79 mg/m ³)
			Skin notation
			Notice of intended change (to TWA 5 ppm (25 mg/m ³))
	OSHA (USA) PEL	2012	TWA 10 ppm (50 mg/m^3)
	EU, 2000/39/EC	2009	Not established
	as amended	2012	
	Germany, MAK		Skin notation
	HSE (UK) WEL	2007	Not established
Solvent naphtha		conta	ppm total hydrocarbon is recommended. Solvent naphtha ins trimethyl benzene. The ACGIH recommends a TLV-TWA ppm (123 g/m ³) for trimethyl benzene.
			ever, other exposure limits may be defined by local regulations nust be observed.
Monitoring methods			ons working with this product for a longer period should have ent blood tests of their cholinesterase levels. If the nesterase level falls below a critical point, no further exposure d be allowed until it has been determined by means of blood that the cholinesterase level has returned to normal.



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Chlorpyrifos DNEL, systemic PNEC, aquatic environment	
Solvent naphtha DNEL, dermal DNEL, inhalation PNEC, aquatic environment	151 mg/m^3
Naphthalene DNEL, dermal DNEL, inhalation PNEC, aquatic environment	25 mg/m^3
8.2. Exposure controls	When used in a closed system, personal protection equipment will not be required. The following is meant for other situations, when the use of a closed system is not possible, or when it is necessary to open the system. Consider the need to render equipment or piping systems non-hazardous before opening.
	The precautions mentioned below are primarily meant for handling of the undiluted product and for preparing the spray solution, but can be recommended for spraying as well.
Respiratory protectio	In the event of an accidental discharge of the material which produces a vapour or mist, workers must put on officially approved respiratory protection equipment with a universal filter type including particle filter.
Protective gloves	Wear chemical resistant gloves, such as barrier laminate, butyl rubber or nitrile rubber. The breakthrough times of these materials for the product are unknown. Generally, however, the use of protective gloves will give only partial protection against dermal exposure. Small tears in the gloves and cross-contamination can easily occur. It is recommended to shift the gloves frequently and to limit the work done manually.

Wear safety glasses. It is recommended to have an eye wash fountain immediately available in the workplace when there is a potential for eye contact.

Wear appropriate chemical resistant clothing to prevent skin contact depending on the extent of exposure. During most normal work situations where exposure to the material cannot be avoided for a limited time span, waterproof pants and apron of chemical resistant material or coveralls of PE will be sufficient. Coveralls of PE must be discarded after use if contaminated. In cases of appreciable or prolonged exposure, coveralls of barrier laminate may be required.

SECTION 9: PHYSICAL AND CHEMICAL PROPERTIES

Eye protection

Other skin protection

9.1. Information on physical and chemical properties Appearance

Yellow to light brown liquid



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	Odour Odour threshold	Aromatic
		Not determined
	pH	1% emulsion in water at 25°C: 5.9
	Melting point/freezing point	Below 0°C
	Initial boiling point and boiling range	Decomposes
	Electronic t	Solvent naphtha : 178 - 209°C
	Flash point	70°C (Pensky-Martens closed tester)
	Evaporation rate	(Butyl acetate = 1)
	Γ_{1}	Solvent naphtha : 0.05
	Flammability (solid/gas)	Not applicable (the product is a liquid)
	Upper/lower flammability or explosive limits	Column transition $(0, 0, 0, 0, 0, 0, 0, 0, 0, 0, 0, 0, 0, 0$
	*	Solvent naphtha : 0.6 - 7.0 vol% (≈ 0.6 - 7.0 kPa) Chlorpyrifos : 2.7 x 10 ⁻³ Pa at 25°C
	Vapour pressure	Chlorpyrifos : 2.7 x 10 ⁻³ Pa at 25°C 1.8 x 10 ⁻² Pa at 35°C
		Solvent naphtha : 100 Pa at 20°C
	Voncur donaite	300 Pa at 38°C
	Vapour density	(Air = 1) Solvent naphtha : > 1
	Relative density	Not determined
	Relative delisity	Density: 1.083 g/ml at 20°C
	Solubility(ies)	Chlorpyrifos : miscible with toluene
	Solubility(les)	miscible with dichloromethane
		miscible with acetone
		miscible with acetate
		774 g/l in hexane at 20°C
		290 g/l in methanol at 20°C
		0.94 mg/l in water at 25°C
	Partition coefficient n-octanol/water	Chlorpyrifos $: \log K_{ow} = 4.7$
	Faithfoll coefficient n-octanol/water	Solvent naphtha : some of the main components have $\log K_{ow} =$
		3.4 - 4.1 at 25°C by model calculation
	Autoignition temperature	460°C
	Decomposition temperature	Not determined (however, see subsection 10.2.)
	Viscosity	3.6 mPa.s at 25°C
		2.0 - 2.3 mPa.s at 45°C
	Explosive properties	Not explosive
	Oxidising properties	Not oxidising
9.2.	Other information	
	Miscibility	The product is emulsifiable in water.
	Surface tension	33 mN/m at 25°C
• SEC	TION 10. STABILITY AND REAC	TIVITV

SECTION 10: STABILITY AND REACTIVITY

10.1.	Reactivity	To our knowledge, the product has no special reactivities.
10.2.	Chemical stability	Chlorpyrifos may decompose rapidly when heated, significantly increasing the risk of explosion. Direct local heating of the product such as electric heating or by steam must be avoided.
		In tests, some variation has been found for the decomposition temperature. The variation may be due to impurities and degree of direct exposure to oxygen. Decomposition appears to be relatively slow up to 160°C, but care seems to be warranted at lower temperatures as well.



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		The decomposition is to a considerable extent dependent on time as well as temperature due to self-accelerating exothermic and autocatalytic reactions. The reactions involve rearrangements and polymerisation releasing volatile malodorous and inflammable compounds such as diethyl sulphide and ethyl mercaptan.
10.3.	Possibility of hazardous reactions	None known.
10.4.	Conditions to avoid	Heating of the product will produce harmful and irritant vapours.
10.5.	Incompatible materials	Strong alkalis and strong oxidising compounds. The product can corrode metals (but does not meet the criteria for classification).
10.6.	Hazardous decomposition products	See subsection 5.2.

♣ SECTION 11: TOXICOLOGICAL INFORMATION

11.1. Information on toxicological effects

Product

Product no.

<u>Product</u>		
Acute toxicity		The product is harmful by ingestion and inhalation. It is considered as less harmful by skin contact. The acute toxicity of the product is measured as:
Route(s) of entry	- ingestion	LD ₅₀ , oral, rat (male): 205 mg/kg (method FIFRA 81-1)
	- skin	LD ₅₀ , dermal, rat: > 4000 mg/kg (method FIFRA 81-2)
	- inhalation	LC50, inhalation, rat: 2.16 mg/l/4 h (method FIFRA 81-3)
Skin corrosion/irrita	ation	Moderately irritating to skin (method FIFRA 81-5).
Serious eye damage	/irritation	Moderately irritating to eyes (method FIFRA 81-4)
Respiratory or skin sensitisation		Not allergenic in animal tests (method FIFRA 81-6). Based on available data, the classification criteria are not met. (B.o.a.d.t.c.c.a.n.m.)
Aspiration hazard		The product presents an aspiration pneumonia hazard.
Symptoms and effects, acute and delayed		On contact, the first symptoms to appear may be irritation. Symptoms of cholinesterase inhibition: nausea, headache, vomiting, cramps, weakness, blurred vision, pin-point pupils, tightness in chest, laboured breathing, nervousness, sweating, watering of eyes, drooling or frothing of mouth and nose, muscle spasms and coma.
<u>Chlorpyrifos</u>		
Acute toxicity		The substance is toxic by ingestion. Toxicity by inhalation is not known. It is considered as less harmful by skin contact. The acute toxicity is measured as:
Route(s) of entry	- ingestion	LD ₅₀ , oral, rat: 172 - 320 mg/kg (method FIFRA 81.01)
	- skin	LD ₅₀ , dermal, rat: > 2000 mg/kg (method FIFRA 81.02)
	- inhalation	LC ₅₀ , inhalation, rat: not available



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Skin corrosion/irritation		Slightly irritating to skin (method FIFRA 81.05). B.o.a.d.t.c.c.a.n.m.
Serious eye damag	e/irritation	Slightly irritating to eyes (method FIFRA 81.04). B.o.a.d.t.c.c.a.n.m.
Respiratory or skin	sensitisation	Not sensitising (method FIFRA 81.06). B.o.a.d.t.c.c.a.n.m.
Germ cell mutagen	iicity	Chlorpyrifos is not mutagenic (23 studies). B.o.a.d.t.c.c.a.n.m.
Carcinogenicity		No carcinogenic effects have been observed for chlorpyrifos (5 studies). B.o.a.d.t.c.c.a.n.m.
Reproductive toxicity		No effects on fertility are found for chlorpyrifos (3 studies). Chlorpyrifos is not teratogenic (not causing birth defects) in rats at levels up to 15 mg/kg/day (a maternally toxic level)(2 studies). B.o.a.d.t.c.c.a.n.m.
STOT – single exp	osure	Mild and transient neurotoxic effects were observed for chlorpyrifos at dose level 50 mg/kg bw. B.o.a.d.t.c.c.a.n.m.
STOT – repeated exposure		Target organ: nervous system (cholinesterase inhibition) LOAEL: 1 mg/kg bw/day in a 90-day rat study. At this exposure level, minor cholinesterase inhibition was found which generally does not result in observable effects or discomfort. A level for observable effects (LOEL) has not been determined. B.o.a.d.t.c.c.a.n.m.
Solvent naphtha	(petroleum), heav	v aromatic
Acute toxicity		The substance is not considered as harmful. B.o.a.d.t.c.c.a.n.m. The acute toxicity is measured as:
Route(s) of entry	- ingestion	LD ₅₀ , oral, rat: > 5000 mg/kg (method similar to OECD 401)
	- skin	LD ₅₀ , dermal, rat: > 2000 mg/kg (method similar to OECD 402)
	- inhalation	LC ₅₀ , inhalation, rat: > 4.7 mg/l/4 h (vapour, method similar to OECD 403)

	method similar to OECD 403)
Skin corrosion/irritation	Can cause skin dryness (measured on a similar product; method OECD 404).
Serious eye damage/irritation	May cause mild, short-lasting discomfort to eyes (method similar to OECD 405). B.o.a.d.t.c.c.a.n.m.
Respiratory or skin sensitisation	To our knowledge, no indications of allergenic properties have been recorded. Measured on a similar substance: not a skin sensitizer (method similar to OECD 406). B.o.a.d.t.c.c.a.n.m.
Germ cell mutagenicity	Not mutagenic in test according to method similar to OECD 479. B.o.a.d.t.c.c.a.n.m.
Carcinogenicity	For petroleum solvents in general, IARC has considered the evidence for carcinogenicity as inadequate.

The product contains naphthalene, which is a suspected carcinogen.



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Reproductive toxicity	Not expected to cause harmful effects on reproduction (measured on similar products; methods OECD 414 and 416). B.o.a.d.t.c.c.a.n.m.
STOT – single exposure	Vapour may cause headache and dizziness.
STOT – repeated exposure	Organic solvents generally are suspected to cause irreversible damage to nervous system on repeated exposure. For some of the main components of solvent naphtha (trimethyl benzenes) this effect was found to occur in humans at a concentration level of around 0.3 mg/l during occupational exposure for periods of 10 to 21 days. LOEL: 0.3 mg/l/day.
	Prolonged and/or repeated skin contact may defat the skin resulting in possible irritation and dermatitis.
Aspiration hazard	Solvent naphtha presents an aspiration hazard.
<u>Naphthalene</u> Acute toxicity	The substance is harmful by ingestion. The acute toxicity is measured as:
Route(s) of entry - ingestion	LD_{50} , oral, rat: > 2000 mg/kg (method OECD 401)
	LD ₅₀ , oral, mouse: 710 mg/kg (method similar to OECD 401)
- skin	LD ₅₀ , dermal, rat: > 2500 mg/kg
- inhalation	LC_{50} , inhalation, rat: > 0.4 mg/l/4 h (vapour, method similar to OECD 403)
Skin corrosion/irritation	Not irritating to skin (method similar to OECD 404). B.o.a.d.t.c.c.a.n.m.
Serious eye damage/irritation	Not irritating to eyes (method similar to OECD 405). B.o.a.d.t.c.c.a.n.m.
Respiratory or skin sensitisation	Not a skin sensitizer (method OECD 406). B.o.a.d.t.c.c.a.n.m.
Germ cell mutagenicity	Results of Chinese hamster ovary cell test were mixed (method OECD 473). Numerous other studies indicate that naphthalene is not mutagenic. B.o.a.d.t.c.c.a.n.m.
Carcinogenicity	Naphthalene is a suspected carcinogen (6 studies).
Reproductive toxicity	Naphthalene produces foetotoxicity at maternally toxic doses in animals (3 studies). Data on fertility are not available. B.o.a.d.t.c.c.a.n.m.
STOT – single exposure	Naphthalene may have narcotic effects at high doses. B.o.a.d.t.c.c.a.n.m.
STOT – repeated exposure	Organic solvents generally are suspected to cause irreversible damage to the brain on repeated exposure.
	NOAEL, oral, was 133 mg/kg bw/day in a 90-day mouse study (method OECD 408), based on decreased weight of brain, liver and spleen.



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	LOAEL, inhalation: 2 ppm (10 mg/m ³) in a 90-day (5 days/week, 6 hrs/day) rat study (method OECD 413), based on minor histopathological changes in the nasal epithelium. B.o.a.d.t.c.c.a.n.m.			
Aspiration hazard .		Naphthalene presents an aspiration pneumonia hazard.		
Calcium dodecyll Acute toxicity	penzene sulphono	tte The substance is not considered as harmful by skin contact, ingestion and inhalation. B.o.a.d.t.c.c.a.n.m. The acute toxicity is measured as:		
Route(s) of entry	- ingestion	LD ₅₀ , oral, rat: 4000 mg/kg		
	- skin	LD ₅₀ , dermal, rat: not available		
	- inhalation	LC ₅₀ , inhalation, rat: not available		
Skin corrosion/irrit	ation	Irritating to skin.		
Serious eye damage	e/irritation	Irritating to eyes with the potential to cause permanent eye damage.		
<u>2-Ethylhexan-1-o</u> Acute toxicity		The substance is not considered as harmful. B.o.a.d.t.c.c.a.n.m. The acute toxicity is measured as:		
Route(s) of entry	- ingestion	LD ₅₀ , oral, rat: 3290 mg/kg (method OECD 401)		
	- skin	LD ₅₀ , dermal, rat: > 3000 mg/kg (method OECD 402)		
	- inhalation	LC ₅₀ , inhalation, rat: 0.89 - 5.3 mg/l/4 h (method OECD 403)		
		Not harmful at saturated vapour pressure (approx. 0.89 mg/l). Harmful at 5.3 mg/l, a mixture of vapour and droplets.		
Skin corrosion/irrit	ation	Mildly irritating to skin. B.o.a.d.t.c.c.a.n.m.		
Serious eye damage	e/irritation	Moderately to severely irritating to eyes.		
Respiratory or skin	sensitisation	Not a skin sensitizer. B.o.a.d.t.c.c.a.n.m.		
Germ cell mutagen	icity	Negative in tests on Chinese hamster ovary cells (methods OECD 473 and 479). B.o.a.d.t.c.c.a.n.m.		
Carcinogenicity		Not carcinogenic to rats and mice (method OECD 451). B.o.a.d.t.c.c.a.n.m.		
Reproductive toxicity		Not expected to cause harmful effects on reproduction. NOAEL for maternal toxicity: 130 mg/kg bw/day NOAEL for teratogenicity: 650 mg/kg bw/day (method OECD 414). B.o.a.d.t.c.c.a.n.m.		
STOT – single exp	osure	Vapour may be irritating to the respiratory tract and may cause headache and dizziness. B.o.a.d.t.c.c.a.n.m.		
STOT – repeated exposure		Organic solvents generally are suspected to cause irreversible damage to nervous system on repeated exposure.		
		Prolonged and/or repeated skin contact may defat the skin resulting		



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chlorpyrifos is measured to be 1375 for whole fish (rainbow trout).

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			in possible irritation and dermatitis.
			Target organs: liver and stomach NOEL: 125 mg/kg bw/day in a 90-day rat study (method OECD 408).
	Aspiration haza	rd	The substance is not of a type normally considered to present an aspiration pneumonia hazard, but it may cause aspiration pneumonia depending on circumstances. B.o.a.d.t.c.c.a.n.m.
♣ SEC	CTION 12: ECO	DLOGICAL INFORM	IATION
12.1.	Toxicity		The product is highly toxic to fish, aquatic invertebrates and insects. It is toxic to aquatic plants, but it is considered as less toxic to birds and not harmful to soil micro- and macroorganisms.
	The acute ecoto	exicity of the product is	measured as:
	- Fish	Rainbow trout (Salmo	9 gairdneri) 96-h LC ₅₀ : 48 µg/l
	- Invertebrates	Daphnids (Daphnia m	nagna) 48-h EC ₅₀ : 2.6 µg/l
	- Algae	Green algae (Selenast	trum capricornutum) 72-h IC ₅₀ : 0.14 mg/l
	- Earthworms	Eisenia foetida foetida	<i>a</i> 14-day LC ₅₀ : 360 mg/kg dry soil
	- Birds	Bobwhite quail (Colin	nus virginianus) LD ₅₀ : 83 mg/kg
	The following h	has been measured on th	ne active ingredient chlorpyrifos:
	- Bees	Honey bees (Apis mel	<i>llifera</i>) LD_{50} , acute oral: 0.36 µg/bee
			LD_{50} , topical: 0.070 µg/bee
12.2.	Persistence and	d degradability	Chlorpyrifos is biodegradable, but does not meet the criteria for being readily biodegradable. It undergoes degradation in the environment and in waste water treatment plants. No adverse effects are found at concentrations up to 100 mg/l in waste water treatment plants. Degradation occurs both aerobically and anaerobically, biologically as well as abiologically.
			Primary degradation half-lives of chlorpyrifos vary with circumstances, but are usually around 4 - 10 weeks in soil and water. pH has a major influence. Degradation will increase at higher pH.
			Solvent naphtha is not readily biodegradable. However, it is expected to be degraded in the environment at a moderate rate.
			The product contains minor amounts of not readily biodegradable components, which may not be degradable in waste water treatment plants.
12.3.	Bioaccumulati	ve potential	See section 9 for octanol-water partition coefficients.
			Chlorpyrifos has the potential to bioaccumulate, but is rapidly excreted (with half-life 2 - 3 days). The bioaccumulation factor of



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		Solvent naphtha has a moderate potential to bioaccumulate if continuous exposure is maintained. Most components can be metabolised by many organisms. Bioaccumulation factors (BCFs) of some of the main components are 246 - 810 by model calculation.
12.4.	Mobility in soil	Chlorpyrifos is not mobile in the environment, but is strongly absorbed to soil
		Solvent naphtha is not mobile in the environment, but it is highly volatile and will rapidly evaporate to the air if released onto water or on the surface of soil. It floats and can migrate to sediment.
12.5.	Results of PBT and vPvB	New of the investigation of the original factor is DDT and DDT
	assessment	None of the ingredients meets the criteria for being PBT or vPvB.
12.6.	Other adverse effects	Other relevant hazardous effects in the environment are not known.
SECT	TON 13: DISPOSAL CONSIDERAT	IONS
13.1.	Waste treatment methods	Remaining quantities of the material and empty but unclean packaging should be regarded as hazardous waste.
		Disposal of waste and packagings must always be in accordance with all applicable local regulations.
	Disposal of product	According to the Waste Framework Directive (2008/98/EC), possibilities for reuse or reprocessing should first be considered. If this is not feasible, the material can be disposed of by removal to a licensed chemical destruction plant or by controlled incineration with flue gas scrubbing.
		Chlorpyrifos is rapidly hydrolysed at $pH > 8.0$.
		Do not contaminate water, foodstuffs, feed or seed by storage or disposal. Do not discharge to sewer systems.
	Disposal of packaging	Containers can be triply rinsed (or equivalent) and offered for recycling or reconditioning. Alternatively, the packaging can be punctured to make it unusable for other purposes and then be disposed of in a sanitary landfill. Controlled incineration with flue gas scrubbing is possible for combustible packaging materials.
▲ SE	CTION 14: TRANSPORT INFORM	ATION

♣ SECTION 14: TRANSPORT INFORMATION

14.1.	UN number	3018
14.2.	UN proper shipping name	Organophosphorus pesticide, liquid, toxic (chlorpyrifos and alkyl(C3-C5)benzenes)
14.3.	Transport hazard class(es)	6.1
14.4.	Packing group	III
14.5.	Environmental hazards	Marine pollutant



Special precautions for user

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Do not discharge to the environment.

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14.7. Transport in bulk according to Annex II of MARPOL 73/78 and the IBC code The product is not transported in bulk tankers. **SECTION 15: REGULATORY INFORMATION** 15.1. Safety, health and environmental Seveso category in Annex I, part 2, to Dir. 96/82/EC: dangerous for regulations/legislation specific for the environment the substance or mixture The Young Worker Directive (94/33/EC) prohibits people under the age of 18 to work with this product. All ingredients in this product are covered by EU chemical legislation. 15.2. A chemical safety assessment has not been performed. Chemical safety assessment **♣** SECTION 16: OTHER INFORMATION

SECTION 16: OTHER INFORMATION

Relevant changes in the SDS	Minor corrections only.		
List of abbreviations	ACGIH	American Conference of Governmental Industrial Hygienists	
	BAT	Biologische Arbeitsstoff-Toleranzwert	
	BEI	Biological Exposure Index	
		c.c.a.n.m.: Based on available data, the classification	
	Dioiaiditie	criteria are not met.	
	CAS	Chemical Abstracts Service	
	CLP	Classification, Labelling and Packaging; refers to EU	
		regulation 1272/2008 as amended	
	Dir.	Directive	
	DNEL	Derived No Effect Level	
	DPD	Dangerous Preparation Directive; refers to Dir.	
		1999/45/EC as amended	
	DSD	Dangerous Substance Directive; refers to Dir.	
		67/548/EEC as amended	
	EC	European Community, or	
		Emulsifiable Concentrate	
	EC_{50}	50% Effect Concentration	
	EINECS	European Inventory of Existing Commercial Chemical	
		Substances	
	FIFRA	Federal Insecticide, Fungicide and Rodenticide Act	
	GHS	Globally Harmonized classification and labelling	
		System of chemicals, Fourth revised edition 2011	
	HSE	Health & Safety Executive, UK	
	IARC	International Agency for Research on Cancer	
	IBC	International Bulk Chemical code	
	IC_{50}	50% Inhibition Concentration	
	ISO	International Organisation for Standardisation	
	IUPAC	International Union of Pure and Applied Chemistry	
	LC_{50}	50% Lethal Concentration	
	LD ₅₀	50% Lethal Dose	
	LOAEL	Lowest Observed Adverse Effect Level	
	LOEL	Lowest Observed Effect Level	



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	MAK	Maximale Arbeitspaltz-Konzentration		
		Set of rules from the International Maritime		
		Organisation (IMO) for prevention of sea pollution		
	NOAEL	No Observed Adverse Effect Level		
	NOEL	No Observed Effect Level		
	OECD	Organisation for Economic Cooperation and		
		Development		
	OSHA	Occupational Safety and Health Administration		
	PBT	Persistent, Bioaccumulative, Toxic		
	PE	Polyethylene		
	PEL	Personal Exposure Limit		
	PNEC	Predicted No Effect Concentration		
	Reg.	Regulation		
		Risk phrase		
	SDS	Safety Data Sheet		
	SE	Single Exposure		
	SP	Safety Precaution		
	S-phrase	Safety phrase		
	STEL	Short-Term Exposure Limit		
	STOT	Specific Target Organ Toxicity		
	TLV	Threshold Limit Value		
	TWA	Time Weighed Average		
	vPvB	very Persistent, very Bioaccumulative		
	WEL	Workplace Exposure Limit		
	WHO	World Health Organisation		
		World Houth Organisation		
References Data measured on a the product a		sured on a the product are unpublished company data.		
		gredients are available from published literature and can		
		several places.		
Method for classification	Acute ora	l toxicity: test data		
	Inhalation toxicity: test data			
		ion: test data		
		enicity: calculation method		
		n toxicity: calculation method		
		o the aquatic environment, acute: test data		
	The second second	chronic: calculation method		
		en onie. eureuluion neulou		
Used R-phrases	R20/22	Harmful by inhalation and if swallowed.		
r	R22	Harmful if swallowed.		
	R25	Toxic if swallowed.		
	R36	Irritating to eyes.		
	R38	Irritating to skin.		
	R40	Limited evidence of a carcinogenic effect.		
	R41	Risk of serious damage to eyes.		
	R50/53	Very toxic to aquatic organisms, may cause long-term		
		adverse effects in the aquatic environment.		
	R51/53	Toxic to aquatic organisms, may cause long-term		
	1.0 1,00	adverse effects in the aquatic environment.		
	R65	Harmful: may cause lung damage if swallowed.		
	R66	Repeated exposure may cause skin dryness and		
	100	cracking.		
	R67	Vapours may cause drowsiness and dizziness.		
	1.07	· apouto maj oudoo dio nomedo and dizzinedo.		
Used CLP hazard statements	H301	Toxic if swallowed.		
	H302	Harmful if swallowed.		
	H302	Harmful if swallowed.		



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	H304	May be fatal if swallowed and enters airways.
	H315	Causes skin irritation.
	H318	Causes serious eye damage.
	H319	Causes serious eye irritation.
	H332	Harmful if inhaled.
	H336	May cause drowsiness or dizziness.
	H351	Suspected of causing cancer.
	H400	Very toxic to aquatic life.
	H410	Very toxic to aquatic life with long lasting effects.
	H411	Toxic to aquatic life with long lasting effects.
	EUH066	Repeated exposure may cause skin dryness and cracking.
	EUH401	To avoid risks to human health and the environment, comply with the instructions of use.
Advice on training	This material should only be used by persons who are made aware of its hazardous properties and have been instructed in the required safety precautions.	

The information provided in this safety data sheet is believed to be accurate and reliable, but uses of the product vary and situations unforeseen by Cheminova A/S may exist. The user has to check the validity of the information under local circumstances.

Prepared by: Cheminova A/S Safety, Health, Environment & Quality Department / GHB