

Kerosene SAFETY DATA SHEET According to EC 1907/2006

# Section 1: Identification of the substance/mixture and of the company/undertaking

1.1 Product identifier	
Name:	Kerosene
Synonyms/Other Means of Identification:	Paraffin
	Lamp Oil
	Regular Kerosene
MARPOL Annex I Category:	Kerosenes
<b>REACH Registration Number:</b>	01-2119485517-27-0006
1.2 Relevant identified uses of the substance or mixture a	nd uses advised against
Intended Use:	Heating Oil
Uses Advised Against:	Uses other than those covered by the exposure scenarios
	appended to this Safety Data Sheet are not supported.
1.3 Details of the supplier of the substance or mixture	
Manufacturer:	Emo Oil Ltd.
	Clonminam Ind Est, Portlaoise, Laois
SDS Information:	http://www.emo.ie/Products/ProductsSheets.aspx
Email:	operations@emo.ie
1.4 Emergency telephone number	+353 (0)57 8674700
Section 2: Hazards Identification	

### 2.1 Classification of the substance or mixture

CLP Classification (EC No 1272/2008):

H226 -- Flammable liquids -- Category 3

H304 -- Aspiration Hazard -- Category 1

H315 -- Skin corrosion/irritation -- Category 2

H336 -- Specific target organ toxicity (single exposure) -- Category 3

H411 -- Hazardous to the aquatic environment, chronic toxicity -- Category 2

Superseded DSD Classification (67/548/EEC and 1999/45/EC):

R10, Xi;R38, Xn;R65, R67, N;R51/53

### 2.2 Label Elements



### DANGER

H226: Flammable liquid and vapour.

H304: May be fatal if swallowed and enters airways.

- H315: Causes skin irritation.
- H336: May cause drowsiness or dizziness.

H411: Toxic to aquatic life with long lasting effects.

P102: Keep out of reach of children.

P210: Keep away from heat/sparks/open flames/hot surfaces. - No smoking.

P280: Wear protective gloves / protective clothing / eye protection / face protection.

P301+P310: IF SWALLOWED: Immediately call a POISON CENTER or doctor/physician.

- P331: Do NOT induce vomiting.
- P501: Dispose of contents/container to approved disposal facility.





### 2.3 Other hazards

Does not meet the criteria for persistent, bioaccumulative and toxic (PBT) or very persistent, very bioaccumulative (vPvB) substances

# Section 3: Composition / Information on Ingredients

### 3.1 Substance

Component	CASRN	EINECS	REACH Registration No.	Concentration <sup>1</sup>	CLP Classification <sup>2</sup>	DSD Classification <sup>3</sup>
KeroseneC9-16	8008-20-6	232-366-4	01-2119485517-27	100	H304	Xn;R65

<sup>1</sup> All concentrations are percent by weight unless ingredient is a gas. Gas concentrations are in percent by volume

<sup>2</sup> Regulations EC 1272/2008

<sup>3</sup> Superseded Directives 67/548/EEC and 1999/45/EC

### Total Sulphur: < 0.1 wt%

### **Section 4: First Aid Measures**

### 4.1 Description of first aid measures

- **Eye Contact:** If irritation or redness develops from exposure, flush eyes with clean water. If symptoms persist, seek medical attention.
- Skin Contact: Remove contaminated shoes and clothing, and flush affected area(s) with large amounts of water. If skin surface is damaged, apply a clean dressing and seek medical attention. If skin surface is not damaged, cleanse affected area(s) thoroughly by washing with mild soap and water or a waterless hand cleaner. If irritation or redness develops, seek medical attention. Wash contaminated clothing before reuse.
- Inhalation (Breathing): If respiratory symptoms develop, move victim away from source of exposure and into fresh air in a position comfortable for breathing. If breathing is difficult, oxygen or artificial respiration should be administered by qualified personnel. If symptoms persist, seek medical attention.
- **Ingestion (Swallowing):** Aspiration hazard: Do not induce vomiting or give anything by mouth because this material can enter the lungs and cause severe lung damage. If victim is drowsy or unconscious and vomiting, place on the left side with the head down. If possible, do not leave victim unattended and observe closely for adequacy of breathing. Seek medical attention.

### 4.2 Most important symptoms and effects

Acute: Minor respiratory irritation at high vapour concentrations.

**Delayed:** Dry skin and possible irritation with repeated or prolonged exposure

# 4.3 Indication of immediate medical attention and special treatment needed

Other Comments: None

### Section 5: Fire-Fighting Measures

### 5.1 Extinguishing media

Dry chemical, carbon dioxide, or foam is recommended. Water spray is recommended to cool or protect exposed materials or structures. Carbon dioxide can displace oxygen. Use caution when applying carbon dioxide in confined spaces. Simultaneous use of foam and water on the same surface is to be avoided as water destroys the foam. Water may be ineffective for extinguishment, unless used under favourable conditions by experienced fire fighters.





### 5.2 Special hazards arising from the substance or mixture

**Unusual Fire & Explosion Hazards:** Flammable. This material can be ignited by heat, sparks, flames, or other sources of ignition (e.g., static electricity, pilot lights, mechanical/electrical equipment, and electronic devices such as cell phones, computers, calculators, and pagers which have not been certified as intrinsically safe). Vapours may travel considerable distances to a source of ignition where they can ignite, flash back, or explode. May create vapour/air explosion hazard indoors, in confined spaces, outdoors, or in sewers. This product will float and can be reignited on surface water. Vapours are heavier than air and can accumulate in low areas. If container is not properly cooled, it can rupture in the heat of a fire.

**Hazardous Combustion Products:** Combustion may yield smoke, carbon monoxide, and other products of incomplete combustion. Oxides of nitrogen and sulphur may also be formed.

### 5.3 Special protective actions for fire-fighters

For fires beyond the initial stage, emergency responders in the immediate hazard area should wear protective clothing. When the potential chemical hazard is unknown, in enclosed or confined spaces, a self-contained breathing apparatus should be worn. In addition, wear other appropriate protective equipment as conditions warrant (see Section 8). Isolate immediate hazard area and keep unauthorized personnel out. Stop spill/release if it can be done safely. Move undamaged containers from immediate hazard area if it can be done safely. Water spray may be useful in minimizing or dispersing vapours and to protect personnel. Avoid spreading burning liquid with water used for cooling purposes. Cool equipment exposed to fire with water, if it can be done safely.

### See Section 9 for Flammable Properties including Flash Point and Flammable (Explosive) Limits

### Section 6: Accidental Release Measures

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

Flammable. Spillages of liquid product will create a fire hazard and may form an explosive atmosphere. Keep all sources of ignition and hot metal surfaces away from spill/release if safe to do so. The use of explosion-proof electrical equipment is recommended. Stay upwind and away from spill/release. Avoid direct contact with material. For larges spillages, notify persons downwind of the spill/release, isolate immediate hazard area and keep unauthorized personnel out. Wear appropriate protective equipment, including respiratory protection, as conditions warrant (see Section 8). See Sections 2 and 7 for additional information on hazards and precautionary measures.

### 6.2 Environmental precautions

Stop spill/release if it can be done safely. Prevent spilled material from entering sewers, storm drains, other unauthorized drainage systems, and natural waterways. Use foam on spills to minimize vapours. Use water sparingly to minimize environmental contamination and reduce disposal requirements. If spill occurs on water notify appropriate authorities and advise shipping of any hazard.

### 6.3 Methods and material for containment and cleaning up

Notify relevant authorities in accordance with all applicable regulations. Immediate clean-up of any spill is recommended. Dike far ahead of spill for later recovery or disposal. Absorb spill with inert material such as sand or vermiculite, and place in suitable container for disposal. If spilled on water remove with appropriate methods (e.g. skimming, booms or absorbents). In case of soil contamination, remove contaminated soil for remediation or disposal, in accordance with local regulations.

Recommended measures are based on the most likely spillage scenarios for this material; however local conditions and regulations may influence or limit the choice of appropriate actions to be taken.





### Section 7: Handling and Storage

#### 7.1 Precautions for safe handling

Keep away from ignition sources such as heat/sparks/open flame – No smoking. Take precautionary measures against static discharge. Nonsparking tools should be used. Wear protective gloves/clothing and eye/face protection. Wash thoroughly after handling. Use good personal hygiene practices and wear appropriate personal protective equipment.

Flammable. May vaporize easily at ambient temperatures. The vapour is heavier than air and may create an explosive mixture of vapour and air. Beware of accumulation in confined spaces and low lying areas. Open container slowly to relieve any pressure. Electrostatic charge may accumulate and create a hazardous condition when handling or processing this material. To avoid fire or explosion, dissipate static electricity during transfer by grounding and bonding containers and equipment before transferring material. The use of explosion-proof electrical equipment is recommended and may be required (see appropriate fire codes for specific bonding/grounding requirements). Do not enter confined spaces such as tanks or pits without following proper entry procedures. Do not wear contaminated clothing or shoes. Keep contaminated clothing away from sources of ignition such as sparks or open flames.

The use of hydrocarbon fuel in an area without adequate ventilation may result in hazardous levels of incomplete combustion products (e.g. carbon monoxide, oxides of sulphur and nitrogen, benzene and other hydrocarbons) and/or dangerously low oxygen levels.

#### 7.2 Conditions for safe storage, including any incompatibilities

Keep container(s) tightly closed and properly labelled. Use and store this material in cool, dry, well-ventilated areas away from heat, direct sunlight, hot metal surfaces, and all sources of ignition. Store only in approved containers. Post area "No Smoking or Open Flame." Keep away from any incompatible material (see Section 10). Protect container(s) against physical damage.

"Empty" containers retain residue and may be dangerous. Do not pressurize, cut, weld, braze, solder, drill, grind, or expose such containers to heat, flame, sparks, or other sources of ignition. They may explode and cause injury or death. "Empty" drums should be completely drained, properly bunged, and promptly shipped to the supplier or a drum reconditioner. All containers should be disposed of in an environmentally safe manner and in accordance with governmental regulations. Before working on or in tanks which contain or have contained this material, refer to appropriate guidance pertaining to cleaning, repairing, welding, or other contemplated operations. Outdoor or detached storage is preferred. Indoor storage should meet Country or Committee standards and appropriate fire codes.

#### 7.3 Specific end use(s)

Refer to supplemental exposure scenarios if attached.

# Section 8: Exposure Controls / Personal Protection

#### 8.1 Control parameters

	Occupational E	Exposure Limits	
Component	US-ACGIH	Ireland-HSA	Other
KeroseneC9-16	TWA: 200 mg/m3 Skin	None	None

STEL = Short Term Exposure Limit (15 minutes): TWA – Time Weighted Average (8 hours): None – No Occupational Exposure Limit

	Biological Limit Values	
Component	US-ACGIH	EU 98/24/EC
KeroseneC9-16	None	None

None = No Biological Limit Value

Relevant DNEL and PNEC: Pending

Kerosene Date of Issue: 01- Dec – 2014







### 8.2 Exposure controls

**Engineering controls:** If current ventilation practices are not adequate to maintain airborne concentrations below the established exposure limits, additional engineering controls may be required.

**Eye/Face Protection:** The use of eye protection that meets or exceeds EN 166 is recommended to protect against potential eye contact, irritation, or injury. Depending on conditions of use, close fitting eye protection and a face shield may be necessary.

**Skin/Hand Protection:** The use of gloves impervious to the specific material handled is advised to prevent skin contact. Users should check with manufacturers to confirm the breakthrough performance of their products. Depending on exposure and use conditions, additional protection may be necessary to prevent skin contact including use of items such as chemical resistant boots, aprons, arm covers, hoods, coveralls, or encapsulated suits. Suggested protective materials: Nitrile

**Respiratory Protection:** Where there is potential for airborne exposure above the exposure limit an approved air purifying respirator equipped with Type A, organic gases and vapour filters (as specified by the manufacturer) may be used.

A respiratory protection program that follows recommendations for the selection, use, care and maintenance of respiratory protective devices in EN 529:2005 should be followed whenever workplace conditions warrant a respirator's use. Air purifying respirators provide limited protection and cannot be used in atmospheres that exceed the maximum use concentration (as directed by regulation or the manufacturer's instructions), in oxygen deficient (less than 19.5 percent oxygen) situations, or under conditions that are immediately dangerous to life and health.

**Other Protective Equipment:** Eye wash and quick-drench shower facilities should be available in the work area. Thoroughly clean shoes and wash contaminated clothing before reuse.

**Environmental Exposure Controls:** Refer to Sections 6, 7, 12 and 13.

Suggestions provided in this section for exposure control and specific types of protective equipment are based on readily available information. Users should consult with the specific manufacturer to confirm the performance of their protective equipment. Specific situations may require consultation with industrial hygiene, safety, or engineering professionals.





# **Section 9: Physical and Chemical Properties**

# 9.1 Information on basic physical and chemical properties

Data represent typical values and are not intended to be specifications. N/A = Not Applicable; N/D = Not Determined

Appearance:	Colourless
Physical Form:	Liquid
Odour:	Mild paraffinic
Odour Threshold:	N/D
	,
pH:	N/A
Melting/Freezing Point:	N/D
Initial Boiling Point/Range:	150-290°C
Flash Point:	>38°C
Evaporation Rate (nBuAc=1):	N/D
Flammability (solid, gas):	Flammable
Upper Explosive Limits (vol % in air):	6.0
Lower Explosive Limits (vol % in air):	0.5
Vapour Pressure:	3 kPa @20°C
Relative Vapour Density (air=1):	>1
Relative Density (water=1):	0.77-0.82 @ 15°C
Solubility (ies):	Solubility in water: Negligible
Partition Coefficient (n-octanol/water) (Kow):	N/D
Auto-ignition Temperature:	250°C
Decomposition Temperature:	N/D
Viscosity:	1.0-2.0 mm²/s @ 20°C
Explosive Properties:	N/A
Oxidising Properties:	N/A

### 9.2 Other Information

### **Pour Point:**

>-25°C

Section	10:	Stability	/ and	Reacti	vitv
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10.1 Reactivity:	Not chemically reactive.
10.2 Chemical stability:	Stable under normal ambient and anticipated conditions of use.
10.3 Possibility of hazardous reactions:	Hazardous reactions not anticipated.
10.4 Conditions to avoid:	Avoid high temperatures and all sources of ignition. Prevent vapour accumulation.
10.5 Incompatible materials:	Avoid contact with strong oxidizing agents and strong reducing agents.
10.6 Hazardous decomposition products:	Not anticipated under normal conditions of use.





# **Section 11: Toxicological Information**

### 11.1 Information on Toxicological Effects of Substance/Mixture

Substance / Mixture					
Acute Toxicity	Hazard		Additional Information	LC50/LD50 Data	
Inhalation	Unlikely to be harmful			> 5.2 mg/L (mist)	
Skin Absorption	Unlikely to	be harmful		> 2 g/kg	
Ingestion (Swallowing)	Unlikely to	be harmful		> 5 g/kg	
Aspiration Hazard:		May be fatal if s	wallowed and enters airways		
Skin Corrosion/Irrit	ation:	Causes skin irrita	Causes skin irritation. Repeated exposure may cause skin dryness or cracking.		
Serious Eye Damage	Serious Eye Damage/Irritation: C		Causes mild eye irritation.		
can cause n of coordina		can cause minor of coordination,	respiratory irritation, heada	not likely, high concentrations che, drowsiness, dizziness, loss ngestion can cause irritation of omiting.	
Skin Sensitization:	Skin Sensitization: Not expected to be a skin sensitizer.				
<b>Respiratory Sensitiz</b>	Respiratory Sensitization: No information available.				
Specific Target Orga	an Toxicity (Si	ngle Exposure):	May cause drowsiness and	dizziness.	
Specific Target Orga	an Toxicity (Re	peated Exposure	: Not expected to cause orga exposure.	in effects from repeated	
Carcinogenicity:		to cause skin tur Follow-up studie nongenotoxic m and that they ar skin irritation. M	mours in mice following repe es have shown that these tun echanism associated with fre e not likely to cause tumours	ddle distillates have been shown ated and prolonged skin contact. nours are produced through a equent cell damage and repair, in the absence of prolonged ynuclear aromatic hydrocarbon ogen by IARC.	
Germ Cell Mutagen	icity:	Not expected to	cause heritable genetic effe	cts.	
Reproductive Toxic	ity:	Not expected to	cause reproductive toxicity.		
Section 12: Ecological In	formation				

# Section 12: Ecological Information

### 12.1 Toxicity

Acute aquatic toxicity studies on samples of jet fuel and kerosene streams show acute toxicity values greater than 1 mg/L and mostly in the range 1-100 mg/L. These tests were carried out on water accommodated fractions, in closed systems to prevent evaporative loss. Results are consistent with the predicted aquatic toxicity of these substances based on their hydrocarbon composition. Kerosenes should be regarded as toxic to aquatic organisms, with the potential to cause long term adverse effects in the aquatic environment. Classification: H411; Chronic Cat 2.

### 12.2 Persistence and degradability

The hydrocarbons in this material are not readily biodegradable but are regarded as inherently biodegradable since their hydrocarbon components can be degraded by microorganisms.

# Persistence per IOPC Fund definition: Non-Persistent





### 12.3 Bioaccumulative potential

Hydrocarbon constituents of kerosene show measured or predicted Log Kow values ranging from 3 to 6 and above and therefore would be regarded as having the potential to bioaccumulate. In practice, metabolic processes may reduce bioconcentration.

### 12.4 Mobility in soil and environmental fate

On release to water, hydrocarbons will float on the surface and since they are sparingly soluble, the only significant loss is volatilization to air. It is possible that some of the higher molecular weight hydrocarbons will be adsorbed on sediment. Biodegradation in water is a minor loss process. In air, these hydrocarbons are photo degraded by reaction with hydroxyl radicals with half-lives varying from 0.1 to 0.7 days.

### 12.5 Results of PBT and vPvB Assessment

Not a PBT or vPvB substance.

# **12.6 Other Adverse Effects**

None anticipated.

### Section 13: Disposal Considerations

### 13.1 Waste treatment methods

### **European Waste Code:** 13 07 03\* other fuels (including mixtures)

This material, if discarded as produced, would be considered as hazardous waste pursuant to Directive 91/689/EEC on hazardous waste, and subject to the provisions of that Directive unless Article 1(5) of that Directive applies. This code has been assigned based upon the most common uses for this material and may not reflect contaminants resulting from actual use. Waste generators/producers are responsible for assessing the actual process used when generating the waste and it's contaminants in order to assign the proper waste disposal code.

Disposal must be in accordance with Directive 2006/12/EC and other applicable national or regional provisions, and based upon material characteristics at time of disposal. For incineration of waste, follow Directive 2000/76/EC. For landfill of waste, follow Directive 1999/31/EC. Product is suitable for burning in an enclosed controlled burner for fuel value if >5000 BTU, or disposal by supervised incineration at very high temperatures to prevent formation of undesirable combustion products. Follow Directive 2000/76/EC.

**Empty Containers:** Container contents should be completely used and containers emptied prior to discard. Empty drums should be properly sealed and promptly returned to a drum reconditioner. All containers should be disposed of in an environmentally safe manner and in accordance with applicable regulations.

### **Section 14: Transport Information**

14.1 UN number	UN1223
14.2 UN proper shipping name	HEATING OIL, LIGHT
14.3 Transport hazard class(es)	3
14.4 Packing group	III
14.5 Environmental hazards	Marine pollutant
14.6 Special precautions for user	If transported in bulk by marine vessel in international waters, product is being carried under the scope of MARPOL Annex I.

14.7 Transport in bulk according to Annex II of MARPOL 73/78 and the IBC Code Not Applicable

### **Section 15: Regulatory Information**





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

EC 1272/2008 - Classification, labelling and packaging of substances and mixtures EN166:2002 Eye Protection EN 529:2005 Respiratory Protective devices BS EN 374-1:2003 Protective gloves against chemicals and micro-organisms Occupational Exposure Limits, Health and Safety Authority Federal Water Act on the Classification of Substances Hazardous to Waters Directive 91/689/EEC on hazardous waste (European Waste Codes) Directive 2000/76/EC on incineration of waste Directive 1999/31/EC on landfill of waste **Export Rating:** NLR (No License Required)

### **15.2 Chemical Safety Assessment**

A chemical safety assessment has been carried out for the substance/mixture.

### **Section 16: Other Information**

Date of Issue:
Status: Previous Issue Date:
Revised Sections or Basis for Revision:

01-Dec-2014 FINAL 01-Dec-2011 Format change Composition (Section 3) English

#### Language:

### List of Relevant Hazard Statements:

H226: Flammable liquid and vapour

H304: May be fatal if swallowed and enters airways

H315: Causes skin irritation

H336: May cause drowsiness or dizziness

H411: Toxic to aquatic life with long lasting effects

R10: Flammable.

R38: Irritating to skin.

R65: Harmful: may cause lung damage if swallowed.

R67: Vapours may cause drowsiness and dizziness.

R51/53: Toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment.

### Guide to Abbreviations:

ACGIH = American Conference of Governmental Industrial Hygienists; ADR = Agreement on Dangerous Goods by Road; BMGV = Biological Monitoring Guidance Value; CASRN = Chemical Abstracts Service Registry Number; CEILING = Ceiling Limit (15 minutes); EINECS – European Inventory of Existing Commercial Chemical Substances; EPA = [US] Environmental Protection Agency; Germany-TRGS = Technical Rules for Dangerous Substances; IARC = International Agency for Research on Cancer; ICAO/IATA = International Civil Aviation Organization / International Air Transport Association; IMDG = International Maritime Dangerous Goods; Ireland-HSA = Ireland's National Health and Safety Authority; LEL = Lower Explosive Limit; N/A = Not Applicable; N/D = Not Determined; NTP = [US] National Toxicology Program; PBT = Persistent, Bioaccumulative and Toxic; RID = Regulations Concerning the International Transport of Dangerous Goods by Rail; STEL = Short Term Exposure Limit (15 minutes); TLV = Threshold Limit Value; TWA = Time Weighted Average (8 hours); UEL = Upper Explosive Limit; UK-EH40 = United Kingdom EH40/2005 Workplace Exposure Limits; vPvB = very Persistent, very Bioaccumulative

### **Disclaimer of Expressed and implied Warranties:**

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