

Safety Data Sheet

Material Name:R-134A

Issue Date: 11/16/2017

SECTION 1 IDENTIFICATION

Product Identifier

Product name	R-134A
Chemical Name	1,1,1,2-tetrafluoroethane
Synonyms	C2H2F4, CF3CFH2, 1,1,1,2-tetrafluoroethane, propellant R 134A Fluorocarbon HFC HFA 134a, Amerfrost A-134a, Blow Hard O.S. Extra, Dust-Pro Pressurized Duster, BOC R134A, DuPont SUVA 134a Refrigerant, Freon 134a, Koudemiddel R-134a
Proper shipping name	1,1,1,2-Tetrafluoroethane or Refrigerant gas R 134a
Chemical formula	C2H2F4
Other means of identification	Not Available
CAS number	811-97-2

Recommended use of the chemical and restrictions on use

Relevant identified uses	The use of a quantity of material in an unventilated or confined space may result in increased exposure and an irritating atmosphere developing. Before starting consider control of exposure by mechanical ventilation. Wide use in household and commercial refrigeration and automotive air conditioning. Suitable for use in medium temperature food cabinets, water chillers and fountains, heat pumps and dehumidifiers and as a blowing agent for various foams. Other uses include as a propellant for
	aerosol pharmaceuticals, lacquers, deodorants, perfumes, mousses, air fresheners, insecticides, cleaning products and other household products.

Name, address, and telephone number of the chemical manufacturer, importer, or other responsible party

Registered company name	Electronic Fluorocarbons, LLC
Address	3266 Bergey Road, Hatfield, PA 19440
Telephone	215-443-9600
Fax	1-888-817-7161
Website	www.electronicfluorocarbons.com
Email	sales@efgases.com

Emergency phone number

Association / Organisation	Not Available	
Emergency telephone numbers	Not Available	1-800-535-5053 (Infotrac)
Other emergency telephone numbers	Not Available	Outside the US: 1-352-323-3500

SECTION 2 HAZARD(S) IDENTIFICATION

Classification of the substance or mixture

NFPA 704 diamond



Note: The hazard category numbers found in GHS classification in section 2 of this SDSs are NOT to be used to fill in the NFPA 704 diamond. Blue = Health Red = Fire Yellow = Reactivity White = Special (Oxidizer or water reactive substances)

Classification

Simple Asphyxiant, Gas under Pressure (Liquefied gas)

Hazard pictogram(s)	\diamond	
SIGNAL WORD	WARNING	
Hazard statement(s)		
H280	Contains gas under pressure; may explode if heated.	
	May displace oxygen and cause rapid suffocation	
Hazard(s) not otherwise sp	pecified	
Not Applicable		
Not Applicable		
Precautionary statement(s	Precautionary statement(s) Prevention	
Not Applicable	ot Applicable	
Precautionary statement(s) Response		
Not Applicable		
Precautionary statement(s	Precautionary statement(s) Storage	

P410+P403 Protect from sunlight. Store in a well-ventilated place.

Precautionary statement(s) Disposal

Not Applicable

SECTION 3 COMPOSITION / INFORMATION ON INGREDIENTS

Substances

CAS No	%[weight]	Name
811-97-2	>=99	<u>R-134A</u>

Mixtures

See section above for composition of Substances

SECTION 4 FIRST-AID MEASURES

Description of first aid measures

Description of first aid mea	asures
	If product comes in contact with eyes remove the patient from gas source or contaminated area.
	 Take the patient to the nearest eye wash, shower or other source of clean water.
	Open the eyelid(s) wide to allow the material to evaporate.
	 Gently rinse the affected eye(s) with clean, cool water for at least 15 minutes. Have the patient lie or sit down and tilt the head back. Hold the eyelid(s) open and pour water slowly over the eyeball(s) at the inner corners, letting the water run out of the outer corners. The actient may be increased and wich to lease along and which to head back.
	The patient may be in great pain and wish to keep the eyes closed. It is important that the material is rinsed from the eyes to prevent further damage.
Eye Contact	 Ensure that the patient looks up, and side to side as the eye is rinsed in order to better reach all parts of the eye(s) Transport to hospital or doctor.
	Even when no pain persists and vision is good, a doctor should examine the eye as delayed damage may occur.
	If the patient cannot tolerate light, protect the eyes with a clean, loosely tied bandage.
	Ensure verbal communication and physical contact with the patient.
	DO NOT allow the patient to rub the eyes
	DO NOT allow the patient to tightly shut the eyes
	DO NOT introduce oil or ointment into the eye(s) without medical advice
	DO NOT use hot or tepid water.
	If skin contact occurs:
	Immediately remove all contaminated clothing, including footwear.
	Flush skin and hair with running water (and soap if available).
	 Seek medical attention in event of irritation.
	In case of cold burns (frost-bite):
	Move casualty into warmth before thawing the affected part; if feet are affected carry if possible
Chin Contract	Bathe the affected area immediately in luke-warm water (not more than 35 deg C) for 10 to 15 minutes, immersing if possible and without
Skin Contact	rubbing
	DO NOT apply hot water or radiant heat.
	Apply a clean, dry, light dressing of "fluffed-up" dry gauze bandage
	If a limb is involved, raise and support this to reduce swelling
	If an adult is involved and where intense pain occurs provide pain killers such as paracetomol
	Transport to hospital, or doctor
	Subsequent blackening of the exposed tissue indicates potential of necrosis, which may require amputation.

Ingestion	 Administer rescue breathing (preferably with a demand-valve resuscitator, bag-valve mask-device, or pocket mask as trained) or CPR if necessary. Not considered a normal route of entry. Avoid giving milk or oils. Avoid giving alcohol. If spontaneous vomiting appears imminent or occurs, hold patient's head down, lower than their hips to help avoid possible aspiration of vomitus.
Inhalation	 Following exposure to gas, remove the patient from the gas source or contaminated area. NOTE: Personal Protective Equipment (PPE), including positive pressure self-contained breathing apparatus may be required to assure the safety of the rescuer. Prostheses such as false teeth, which may block the airway, should be removed, where possible, prior to initiating first aid procedures. If the patient is not breathing spontaneously, administer rescue breathing. If the patient does not have a pulse, administer CPR. If medical oxygen and appropriately trained personnel are available, administer 100% oxygen. Summon an emergency ambulance. If an ambulance is not available, contact a physician, hospital, or Poison Control Centre for further instruction. Keep the patient warm, comfortable and at rest while awaiting medical care. MONITOR THE BREATHING AND PULSE, CONTINUOUSLY.

Most important symptoms and effects, both acute and delayed

See Section 11

Indication of any immediate medical attention and special treatment needed

for intoxication due to Freons/ Halons;

A: Emergency and Supportive Measures

- Maintain an open airway and assist ventilation if necessary
- Treat coma and arrhythmias if they occur. Avoid (adrenaline) epinephrine or other sympathomimetic amines that may precipitate ventricular arrhythmias. Tachyarrhythmias caused by increased myocardial sensitisation may be treated with propranolol, 1-2 mg IV or esmolol 25-100 microgm/kg/min IV.
- Monitor the ECG for 4-6 hours

B: Specific drugs and antidotes:

There is no specific antidote

C: Decontamination

- ▶ Inhalation; remove victim from exposure, and give supplemental oxygen if available.
- Ingestion; (a) Prehospital: Administer activated charcoal, if available. DO NOT induce vomiting because of rapid absorption and the risk of abrupt onset CNS depression. (b) Hospital: Administer activated charcoal, although the efficacy of charcoal is unknown. Perform gastric lavage only if the ingestion was very large and recent (less than 30 minutes)

D: Enhanced elimination:

• There is no documented efficacy for diuresis, haemodialysis, haemoperfusion, or repeat-dose charcoal.

POISONING and DRUG OVERDOSE, Californian Poison Control System Ed. Kent R Olson; 3rd Edition

- Do not administer sympathomimetic drugs unless absolutely necessary as material may increase myocardial irritability.
- No specific antidote.
- Because rapid absorption may occur through lungs if aspirated and cause systematic effects, the decision of whether to induce vomiting or not should be made by an attending physician.
- ▶ If lavage is performed, suggest endotracheal and/or esophageal control.
- Danger from lung aspiration must be weighed against toxicity when considering emptying the stomach.
- Treatment based on judgment of the physician in response to reactions of the patient

For frost-bite caused by liquefied petroleum gas:

- If part has not thawed, place in warm water bath (41-46 C) for 15-20 minutes, until the skin turns pink or red.
- Analgesia may be necessary while thawing.
- If there has been a massive exposure, the general body temperature must be depressed, and the patient must be immediately rewarmed by whole-body immersion, in a bath at the above temperature.
- Shock may occur during rewarming.
- Administer tetanus toxoid booster after hospitalization.
- Prophylactic antibiotics may be useful.
- The patient may require anticoagulants and oxygen.

[Shell Australia 22/12/87]

For gas exposures:

BASIC TREATMENT

- Establish a patent airway with suction where necessary.
- Watch for signs of respiratory insufficiency and assist ventilation as necessary.
- Administer oxygen by non-rebreather mask at 10 to 15 l/min.
- Monitor and treat, where necessary, for pulmonary oedema .
- Monitor and treat, where necessary, for shock.
- Anticipate seizures.
- _____

ADVANCED TREATMENT

- Consider orotracheal or nasotracheal intubation for airway control in unconscious patient or where respiratory arrest has occurred.
- Positive-pressure ventilation using a bag-valve mask might be of use.
- Monitor and treat, where necessary, for arrhythmias.
- Start an IV D5W TKO. If signs of hypovolaemia are present use lactated Ringers solution. Fluid overload might create complications.
- Drug therapy should be considered for pulmonary oedema.
- Hypotension with signs of hypovolaemia requires the cautious administration of fluids. Fluid overload might create complications.
- Treat seizures with diazepam.
- Proparacaine hydrochloride should be used to assist eye irrigation.

SECTION 5 FIRE-FIGHTING MEASURES

Extinguishing media

SMALL FIRE: Use extinguishing agent suitable for type of surrounding fire. LARGE FIRE: Cool cylinder. DO NOT direct water at source of leak or venting safety devices as icing may occur.

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

Special protective equipment and precautions for fire-fighters

Fire Fighting	GENERAL Alert Fire Department and tell them location and nature of hazard. Wear breathing apparatus and protective gloves. Fight fire from a safe distance, with adequate cover.
Fire/Explosion Hazard	 Use water delivered as a fine spray to control fire and cool adjacent area. Containers may explode when heated - Ruptured cylinders may rocket Fire exposed containers may vent contents through pressure relief devices. High concentrations of gas may cause asphyxiation without warning. May decompose explosively when heated or involved in fire. Contact with gas may cause burns, severe injury and/ or frostbite. Decomposition may produce toxic fumes of: carbon monoxide (CO) Combustion products include: carbon dioxide (CO2) hydrogen fluoride carbon dioxide (CO2) hydrogen fluoride during a substance: Closed containers may rupture due to pressure buildup under fire conditions. Vented gas is more dense than air and may collect in pits, basements. [Although not flammable in air at temperatures up to 100 deg. C at atmospheric temperature, mixtures with high concentrations of air at elevated pressure and / or temperature can become combustible in the presence of an ignition source. The material can also become combustible in an oxygen enriched environment (oxygen concentrations greater than in air). Whether air-mixtures or oxygen-mixtures become combustible depends on temperature, pressure and oxygen concentration.

SECTION 6 ACCIDENTAL RELEASE MEASURES

Personal precautions, protective equipment and emergency procedures

See section 8

Environmental precautions

See section 12

Methods and material for containment and cleaning up

Minor Spills	 Avoid breathing vapour and any contact with liquid or gas. Protective equipment including respirator should be used. DO NOT enter confined spaces where gas may have accumulated. Increase ventilation.
Major Spills	 Clear area of all unprotected personnel and move upwind. Alert Emergency Authority and advise them of the location and nature of hazard. Wear breathing apparatus and protective gloves. Prevent by any means available, spillage from entering drains and water-courses. Remove leaking cylinders to a safe place. Fit vent pipes. Release pressure under safe, controlled conditions Burn issuing gas at vent pipes. DO NOT exert excessive pressure on valve; DO NOT attempt to operate damaged valve.

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

SECTION 7 HANDLING AND STORAGE

Safe handling	 Consider use in closed pressurised systems, fitted with temperature, pressure and safety relief valves which are vented for safe dispersal. Use only properly specified equipment which is suitable for this product, its supply pressure and temperature The tubing network design connecting gas cylinders to the delivery system should include appropriate pressure indicators and vacuum or suction lines. Fully-welded types of pressure gauges, where the bourdon tube sensing element is welded to the gauge body, are recommended. Before connecting gas cylinders, ensure manifold is mechanically secure and does not containing another gas. DO NOT transfer gas from one cylinder to another. [Contact of welding or soldering torch flame with high concentration of refrigerant can result in visible changes in the size and colour of torch flames. This flame effect will only occur in concentrations of product well above the recommended exposure limit.; therefore stop all work and ventilate to disperse refrigerant vapours from the work are before using any open flames.
Other information	 Cylinders should be stored in a purpose-built compound with good ventilation, preferably in the open. Such compounds should be sited and built in accordance with statutory requirements. The storage compound should be kept clear and access restricted to authorised personnel only. Cylinders stored in the open should be protected against rust and extremes of weather.

Conditions for safe storage, including any incompatibilities

Suitable container	 DO NOT use aluminium or galvanised containers Cylinder: Ensure the use of equipment rated for cylinder pressure. Ensure the use of compatible materials of construction. Valve protection cap to be in place until cylinder is secured, connected. Cylinder must be properly secured either in use or in storage.
Storage incompatibility	 As a general rule, hydrofluorocarbons tend to be flammable unless they contain more fluorine atoms than hydrogen atoms. Haloalkanes: are highly reactive:some of the more lightly substituted lower members are highly flammable; the more highly substituted may be used as fire suppressants, not always with the anticipated results. may react with the lighter divalent metals to produce more reactive compounds analogous to Grignard reagents. may produce explosive compounds following prolonged contact with metallic or other azides may react on contact with potassium or its alloys - although apparently stable on contact with a wide rage of halocarbons, reaction products may be shock-sensitive and may explode with great violence on light impact; severity generally increases with the degree of halocarbon substitution and potassium-sodium alloys give extremely sensitive mixtures . BRETHERICK L: Handbook of Reactive Chemical Hazards react with metal halides and active metals, eg. sodium (Na), potassium (K), lithium (Li),calcium (Ca), zinc (Zn), powdered aluminium (Al) and aluminium alloys, magnesium (Mg) and magnesium alloys. Compressed gases may contain a large amount of kinetic energy over and above that potentially available from the energy of reaction produced by the gas in chemical reaction with other substances [Avoid reaction with alkali metals, zinc, aluminium alloys (gt; 2% magnesium).]Avoid contact with plastics such as methacrylate polymers, polyethylene and polystyrene.

SECTION 8 EXPOSURE CONTROLS / PERSONAL PROTECTION

Control parameters

INGREDIENT DATA

Not Available

EMERGENCY LIMITS

Ingredient	Material name	TEEL-1		TEEL-2	TEEL-3
R-134A	HFC 134a; (Tetrafluoroethane, 1,1,1,2-)	Not Available		Not Available	Not Available
Ingredient	Original IDLH	Revised IDLH			
R-134A	Not Available		Not Available		

Exposure controls

Appropriate engineering controls	Engineering controls are used to remove a hazard or place a barrier between the worker and the hazard. Well-designed engineering controls can be highly effective in protecting workers and will typically be independent of worker interactions to provide this high level of protection. The basic types of engineering controls are: Process controls which involve changing the way a job activity or process is done to reduce the risk. Enclosure and/or isolation of emission source which keeps a selected hazard "physically" away from the worker and ventilation that strategically "adds" and "removes" air in the work environment.
Personal protection	
Eye and face protection	 Chemical goggles. Full face shield may be required for supplementary but never for primary protection of eyes. Contact lenses may pose a special hazard; soft contact lenses may absorb and concentrate irritants. A written policy document, describing the wearing of lenses or restrictions on use, should be created for each workplace or task.
Skin protection	See Hand protection below

Hands/feet protection	 When handling sealed and suitably insulated cylinders wear cloth or leather gloves. Insulated gloves: NOTE: Insulated gloves should be loose fitting so that may be removed quickly if liquid is spilled upon them. Insulated gloves are not made to permit hands to be placed in the liquid; they provide only short-term protection from accidental contact with the liquid.
Body protection	See Other protection below
Other protection	 Protective overalls, closely fitted at neck and wrist. Eye-wash unit. Ensure availability of lifeline in confined spaces. Staff should be trained in all aspects of rescue work.
Thermal hazards	Not Available

Respiratory protection

Type AX Filter of sufficient capacity. (AS/NZS 1716 & 1715, EN 143:2000 & 149:2001, ANSI Z88 or national equivalent)

Where the concentration of gas/particulates in the breathing zone, approaches or exceeds the "Exposure Standard" (or ES), respiratory protection is required. Degree of protection varies with both face-piece and Class of filter; the nature of protection varies with Type of filter.

Required Minimum Protection Factor	Half-Face Respirator	Full-Face Respirator	Powered Air Respirator
up to 10 x ES	AX-AUS	-	AX-PAPR-AUS / Class 1
up to 50 x ES	-	AX-AUS / Class 1	-
up to 100 x ES	-	AX-2	AX-PAPR-2 ^

^ - Full-face

A(All classes) = Organic vapours, B AUS or B1 = Acid gasses, B2 = Acid gas or hydrogen cyanide(HCN), B3 = Acid gas or hydrogen cyanide(HCN), E = Sulfur dioxide(SO2), G = Agricultural chemicals, K = Ammonia(NH3), Hg = Mercury, NO = Oxides of nitrogen, MB = Methyl bromide, AX = Low boiling point organic compounds(below 65 degC)

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.

Positive pressure, full face, air-supplied breathing apparatus should be used for work in enclosed spaces if a leak is suspected or the primary containment is to be opened (e.g. for a cylinder change)

+ Air-supplied breathing apparatus is required where release of gas from primary containment is either suspected or demonstrated.

SECTION 9 PHYSICAL AND CHEMICAL PROPERTIES

Information on basic physical and chemical properties

Appearance	Colourless gas with slight ethereal odour; does not mix well with water (0.09 wt %, 25 C, 1 Bar).		
Physical state	Liquified Gas	Relative density (Water = 1)	1.21
Odour	Not Available	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	> 743
pH (as supplied)	Not Applicable	Decomposition temperature	Not Available
Melting point / freezing point (°C)	-101	Viscosity (cSt)	0.210
Initial boiling point and boiling range (°C)	-26.2	Molecular weight (g/mol)	102.03
Flash point (°C)	Not Available	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Not Available	Oxidising properties	Not Available
Upper Explosive Limit (%)	Not Applicable	Surface Tension (dyn/cm or mN/m)	Not Available
Lower Explosive Limit (%)	Not Applicable	Volatile Component (%vol)	100
Vapour pressure (kPa)	560.5	Gas group	Not Available
Solubility in water (g/L)	Immiscible	pH as a solution (1%)	Not Applicable
Vapour density (Air = 1)	3.5	VOC g/L	Not Available

SECTION 10 STABILITY AND REACTIVITY

Reactivity	See section 7
Chemical stability	 Unstable in the presence of incompatible materials. Product is considered stable. Hazardous polymerisation will not occur.

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

	Inhalation of vapours may cause drowsiness and dizz	iness. This may be accompanied b	y sleepiness, reduced alertness, loss of reflexes, lack of	
co-ordination, and vertigo. Inhalation of vapours or aerosols (mists, fumes), generated by the material during the course of normal handling				
	of the individual.			
	There is some evidence to suggest that the material c cause further lung damage.	an cause respiratory irritation in s	ome persons. The body's response to such irritation can	
	Exposure to fluorocarbons can produce non-specific		fever, weakness, muscle pain, headache, chest ise irregular heartbeats and a stepwise reduction in lung	
	capacity.			
Inhaled	Material is highly volatile and may quickly form a cond	centrated atmosphere in confined	or unventilated areas. The vapour may displace and	
	replace air in breathing zone, acting as a simple asphy			
	Symptoms of asphyxia (suffocation) may include hea ears. If the asphyxia is allowed to progress, there may		ath, muscular weakness, drowsiness and ringing in the	
	convulsions, coma and death.	be hausea and vonnting, further		
			eased exposure and an irritating atmosphere developing	
	Before starting consider control of exposure by mech Acute intoxication by halogenated alighatic hydrocart		o stages. Signs of a reversible narcosis are evident in the	
	first stage and in the second stage signs of injury to o			
	Overexposure is unlikely in this form.			
Ingestion	Not normally a hazard due to physical form of product. Considered an unlikely route of entry in commercial/industrial environments			
	Skin contact is not thought to have harmful health eff	•	ves); the material may still produce health damage	
	following entry through wounds, lesions or abrasions. Repeated exposure may cause skin cracking, flaking or drying following normal handling and use.			
	There is some evidence to suggest that this material		-	
	Fluorocarbons remove natural oils from the skin, causing irritation, dryness and sensitivity.			
Skin Contact	Open cuts, abraded or irritated skin should not be exposed to this material Entry into the blood-stream, through, for example, cuts, abrasions or lesions, may produce systemic injury with harmful effects. Examine the			
	skin prior to the use of the material and ensure that a	ny external damage is suitably pro	tected.	
	Vapourising liquid causes rapid cooling and contact may cause cold burns, frostbite, even through normal gloves. Frozen skin tissues are painless and appear waxy and yellow. Signs and symptoms of frost-bite may include "pins and needles", paleness followed by numbness, a			
			, (first white, then mottled and blue and eventually black	
	on recovery, red, hot, painful and blistered).			
Eye	Although the material is not thought to be an irritant discomfort characterised by tearing or conjunctival re		ect contact with the eye may produce transient	
Lye	Not considered to be a risk because of the extreme vo			
			following repeated or long-term occupational exposure.	
	Main route of exposure to the gas in the workplace is by inhalation. Fluorocarbons can cause an increased risk of cancer, spontaneous abortion and birth defects.			
Chronic	No effects have been seen in rats exposed to up to 50,000 ppm for 90 days. It is not teratogenic in rats or rabbits. Short term screening tests			
			stered by gavage to rats at a dose of 300 mg/kg/day for	
	one year and the rats held for the remainder of their l	në span.		
R-134A	ΤΟΧΙCITY	IRRITATION		
N-134A	Inhalation (rat) LC50: >500000 ppm/4hr ^[1]	Not Available	2	
Legend:			obtained from manufacturer's SDS. Unless otherwise	
	specified data extracted from RTECS - Register of Tox	ic Effect of chemical Substances		
		, , , , , , , , , , , , , , , , , , ,	· · · · · · · · ·	
	Disinfection byproducts (DBPs) are formed when disinfectants such as chlorine, chloramines and ozone react with organic and inorganic matter in water. Animal studies have shown that some DBPs cause cancer. To date, several hundred DBPs have been identified.			
R-134A	Numerous haloalkanes and haloalkenes have been tested for cancer-causing and mutation-causing activities.			
	* with added oxygen - ZhongHao New Chemical Mate concentrations of decomposition products can cause		n can have a narcotic effect; inhalation of high	
		-	0	
Acute Toxicity	×	Carcinogenicity	0	
Skin Irritation/Corrosion	©	Reproductivity	©	
Serious Eye	0	STOT - Single Exposure	\otimes	

Damage/Irritation

Respiratory or Skin sensitisation

Mutagenicity

 \odot

Aspiration Hazard

Legend: 🔀

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

Data available to make classification

0

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🚫 – Data Not Available to make classification

SECTION 12 ECOLOGICAL INFORMATION

Toxicity

	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
	LC50	96	Fish	29.671mg/L	3
	EC50	48	Crustacea	980mg/L	5
R-134A	EC50	96	Algae or other aquatic plants	97.260mg/L	3
	EC50	384	Crustacea	7.065mg/L	3
	NOEC	72	Algae or other aquatic plants	ca.13.2mg/L	2
Legend:	V3.12 (QSAR)	, , , , , , , , , , , , , , , , , , , ,	ed Substances - Ecotoxicological Information - Aqu itox database - Aquatic Toxicity Data 5. ECETOC Aq	,	

In addition to carbon dioxide (CO2), methane (CH4) and nitrous oxide (N2O), the greenhouse gases mentioned in the Kyoto Protocol include synthetic substances that share the common feature of being highly persistent in the atmosphere and inhibit radiation from escaping out of the atmosphere. These synthetic substances include hydrocarbons that are partially fluorinated (HCFs) or totally fluorinated (PFCs) as well as sulfur hexafluoride (SF6). The greenhouse potential of these substances, expressed as multiples of that of CO2, are within the range of 140 to 11,700 for HFCs, from 6500 to 9,200 for PFCs and 23,900 for SF6. Once emitted into the atmosphere, these substances have an impact on the environment for decades, centuries, or even for thousands of years.

DO NOT discharge into sewer or waterways.

|Persistence and Degradation:|Ozone Destruction Potential PDO) = 0 (R11=1)|Greenhouse Effect Potential (ESP) = 0.27 (R11=1)|Decomposed comparatively rapidly in the lower atmosphere (troposphere).|Atmospheric lifetime is 15.6 years. Products of decomposition will be highly dispersed and hence will have a very low concentration. Does not influence photochemical smog (i.e. is not a VOC under the terms of the UNECE agreement). Does not deplete ozone.|Effect on Effluent Treatment:|Discharges of the product will enter the atmosphere and will not result in long term aqueous contamination. [ICI]|Ecotoxicity:|Fish LC50 (96 h): Salmo gairdneri 450 mg/l ; NOEC 300 mg/l (mortality) (semi-static tests)|Daphnia EC50 (48 h): 980 mg/l|Bacteria EC10 (6 h): Pseudomonas putida gt;730 mg/l (growth)|Mobility|Air: Henrys Law constant (H): 65 kPa.m3/ml (20 C/ calculated) - considerable volatility|Soil/sediment log Koc 1.5 approx (adsorption - calculated)|Persistence and biodegradability|Abiotic degradation|Air, indirect photo-oxidation 11/2=10.9 years|Conditions: sensitiser: OH radicals|Degradation products: carbon dioxide/ fluorhydric acid/ trichloracetic acid [Air, photolysis, ODP=0]No effect on stratospheric ozone|Reference value for CFC 11: ODP=1|Air, greenhouse effect, GWP=0.25|Reference value for CFC 11: GWP=1|Biotic degradation|Aerobic, test ready biodegradability/ closed bottle, degradation from 2-3% 28 days|Result: not readily biodegradable|Aerobic, test biodegradabil on ymethane oxidation|Result: non-biodegradable|Conditions: inoculum: Methylosinus trichosporium OB3b|Bioaccumulative potential; bioconcentration log POW=1.06|Product is persistent in air (atmospheric lifetime: 15.7 years)|Product is not significantly hazardous for the aquatic environment as: very low toxicity for aquatic organisms|considerable volatility|no bioaccumulation

Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
R-134A	HIGH	HIGH

Bioaccumulative potential

Ingredient	Bioaccumulation
R-134A	LOW (LogKOW = 1.68)

Mobility in soil

Ingredient	Mobility
R-134A	LOW (KOC = 96.63)

SECTION 13 DISPOSAL CONSIDERATIONS

Vaste treatment methods	
Product / Packaging disposal	 Evaporate residue at an approved site. Return empty containers to supplier. If containers are marked non-returnable establish means of disposal with manufacturer prior to purchase. Ensure damaged or non-returnable cylinders are gas-free before disposal.

SECTION 14 TRANSPORT INFORMATION



Marine Pollutant

Land transport (DOT)

UN number	3159
UN proper shipping name	1,1,1,2-Tetrafluoroethane or Refrigerant gas R 134a
Transport hazard class(es)	Class 2.2 Subrisk Not Applicable
Packing group	Not Applicable
Environmental hazard	Not Applicable
Special precautions for user	Hazard Label2.2Special provisionsT50

Air transport (ICAO-IATA / DGR)

UN number	3159	
UN proper shipping name	1,1,1,2-TETRAFLUOROETHANE (REFRIGERANT GAS R 134a)	
Transport hazard class(es)	ICAO/IATA Class 2.2 ICAO / IATA Subrisk Not Applicable ERG Code 2L	
Packing group	Not Applicable	
Environmental hazard	Not Applicable	
Special precautions for user	Special provisions Cargo Only Packing Instructions Cargo Only Maximum Qty / Pack Passenger and Cargo Packing Instructions Passenger and Cargo Maximum Qty / Pack Passenger and Cargo Limited Quantity Packing Instructions	Not Applicable 200 150 kg 200 75 kg Forbidden
	Passenger and Cargo Limited Maximum Qty / Pack	Forbidden

Sea transport (IMDG-Code / GGVSee)

UN number	3159
UN proper shipping name	Refrigerant gas R 134a
Transport hazard class(es)	IMDG Class 2.2 IMDG Subrisk Not Applicable
Packing group	Not Applicable
Environmental hazard	Not Applicable
Special precautions for user	EMS Number F-C, S-V Special provisions Not Applicable Limited Quantities 120 mL

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

Not Applicable

SECTION 15 REGULATORY INFORMATION

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

R-134A(811-97-2) IS FOUND ON THE FOLLOWING REGULATORY LISTS

US - Washington Toxic air pollutants and their ASIL, SQER and de minimis emission values

US AIHA Workplace Environmental Exposure Levels (WEELs)

Federal Regulations

Superfund Amendments and Reauthorization Act of 1986 (SARA)

SECTION 311/312 HAZARD CATEGORIES

Immediate (acute) health hazard	No
Delayed (chronic) health hazard	No
Fire hazard	No
Pressure hazard	Yes
Reactivity hazard	No

US. EPA CERCLA HAZARDOUS SUBSTANCES AND REPORTABLE QUANTITIES (40 CFR 302.4)

None Reported

State Regulations

US. CALIFORNIA PROPOSITION 65

None Reported

National Inventory	Status
Australia - AICS	Υ
Canada - DSL	Υ
Canada - NDSL	N (R-134A)
China - IECSC	Υ
Europe - EINEC / ELINCS / NLP	Y
Japan - ENCS	Υ
Korea - KECI	Υ
New Zealand - NZIoC	Υ
Philippines - PICCS	Υ
USA - TSCA	Υ
Legend:	Y = All ingredients are on the inventory N = Not determined or one or more ingredients are not on the inventory and are not exempt from listing(see specific ingredients in brackets)

SECTION 16 OTHER INFORMATION

Other information

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
- 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