according to Regulation (EC) No. 1907/2006 (REACh) Article 31, Annex II as amended



Isopropylamine 10350

Version / Revision Supersedes Version

6 5.02 Revision Date Issuing date 27-Oct-2022 27-Oct-2022

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

1.1. Product identifier

Identification of the substance/preparation

Isopropylamine

 CAS-No
 75-31-0

 EC No.
 200-860-9

 Registration number (REACh)
 01-2119463274-39

1.2. Relevant identified uses of the substance or mixture and uses advised against

Identified uses	Formulation
Uses advised against	None

1.3. Details of the supplier of the safety data sheet

Company/Undertaking Identification	OQ Chemicals GmbH Rheinpromenade 4A D-40789 Monheim Germany
Product Information	Product Stewardship FAX: +49 (0)208 693 2053 email: sc.psq@oq.com

1.4. Emergency telephone number

Emergency telephone number	+44 (0) 1235 239 670 (UK) available 24/7
National emergency telephone number	National Poisons Information Centre +353 (0)1 809 2166
	available to the public 8 am - 10 pm +353 (0)1 809 2566 available 24/7 for medical professionals

SECTION 2: Hazards identification

2.1. Classification of the substance or mixture

This substance is classified based on Directive 1272/2008/EC and its amendments (CLP Regulation)

Flammable liquid Category 1, H224 Acute oral toxicity Category 3, H301 Acute dermal toxicity Category 3, H311 Acute inhalation toxicity Category 3, H331 Skin corrosion/irritation Category 2, H315 Serious eye damage/eye irritation Category 2, H319 Target Organ Systemic Toxicant - Single exposure Category 3, H335

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In addition to the CLP classification based on OQ data this product should also be regarded as: Skin corrosion/irritation: category 1A-1C

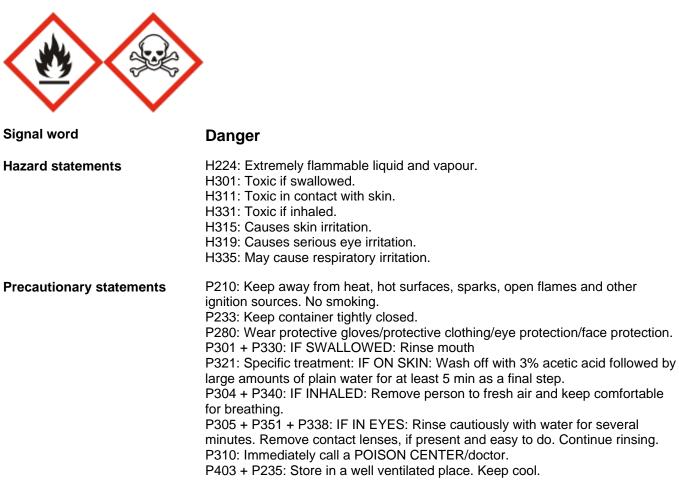
Additional information

For full text of Hazard- and EU Hazard-statements see SECTION 16.

2.2. Label elements

Labelling according to Regulation 1272/2008/EC and its amendments (CLP Regulation).

Hazard pictograms



2.3. Other hazards

Vapours may form explosive mixture with air Vapour is heavier than air and can travel considerable distance to a source of ignition and flashback Components of the product may be absorbed into the body by inhalation, ingestion and through the skin

PBT and vPvB assessment	This substance is not considered to be persistent, bioaccumulating nor toxic (PBT), nor very persistent nor very bioaccumulating (vPvB)
Endocrine disrupting assessments	The substance is not listed on the candidate list according to Art. 59(1), REACh. The substance was not assessed as having endocrine disrupting properties according to regulation 2017/2100/EU or 2018/605/EU.

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SECTION 3: Composition / information on ingredients

3.1. Substances

Component	CAS-No	REACh-No	1272/2008/EC	Concentration (%)
Isopropylamine	75-31-0	01-2119463274-39	Flam. Liq. 1; H224	> 99,7
			Acute Tox. 3; H301	
			Acute Tox. 3; H311	
			Acute Tox. 3, H331	
			Skin Irrit. 2; H315	
			Eye Irrit. 2; H319	
			STOT SE 3; H335	
			ATE = 173 mg/kg	
			(oral)	
			ATE = 400 mg/kg	
			(dermal)	
			$ATE = 8,7 \text{ mg/L}^{***}$	
			(inhalation)	
			(vapours)***	

For full text of Hazard- and EU Hazard-statements see SECTION 16.

SECTION 4: First aid measures

4.1. Description of first aid measures

Inhalation

Keep at rest. Aerate with fresh air. Call a physician immediately. Symptoms of poisoning may develop many hours after exposure.

Skin

Wash off with 3% acetic acid followed by large amounts of plain water for at least 5 min as a final step. Immediate medical treatment is necessary as untreated wounds from corrosion of the skin heal slowly and with difficulty.

Eyes

Rinse immediately with plenty of water, also under the eyelids, for at least 15 minutes. Remove contact lenses. Immediate medical attention is required.

Ingestion

Call a physician immediately. Do not induce vomiting without medical advice.

4.2. Most important symptoms and effects, both acute and delayed

Main symptoms

shortness of breath, convulsions, cough, hypertensive effect, narcosis, unconsciousness, discomfort, nausea.

Special hazard

Stomach perforation, Lung oedema, Pneumonia, Dermatitis.

4.3. Indication of any immediate medical attention and special treatment needed

General advice

Remove contaminated, soaked clothing immediately and dispose of safely. First aider needs to protect himself.

Treat as an alkaline substance (similar to ammonia). If ingested, irrigate the stomach. Treat skin and mucous

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membranes with antihistamine and corticoids. In case of lung irritation, first treatment with cortisone spray. Symptoms may be delayed. Later control for pneumonia and lung oedema.

SECTION 5: Firefighting measures

5.1. Extinguishing media

Suitable extinguishing media

alcohol-resistant foam, dry chemical, carbon dioxide (CO2), water spray

Unsuitable Extinguishing Media

Do not use a solid water stream as it may scatter and spread fire.

5.2. Special hazards arising from the substance or mixture

Under conditions giving incomplete combustion, hazardous gases produced may consist of: carbon monoxide (CO) carbon dioxide (CO2) nitrogen oxides (NOx) hydrogen cyanide (hydrocyanic acid) Combustion gases of organic materials must in principle be graded as inhalation poisons Vapour is heavier than air and can travel considerable distance to a source of ignition and flashback Vapours may form explosive mixture with air

5.3. Advice for firefighters

Special protective equipment for firefighters

Fire fighter protection should include a self-contained breathing apparatus (NIOSH-approved or EN 133) and full fire-fighting turn out gear.

Precautions for firefighting

Cool containers / tanks with water spray. Water run-off and vapor cloud may be corrosive. Dike and collect water used to fight fire. Keep people away from and upwind of fire.

SECTION 6: Accidental release measures

6.1. Personal precautions, protective equipment and emergency procedures

For non-emergency personnel: For personal protective equipment see section 8. Avoid contact with skin and eyes. Avoid breathing vapors or mists. Keep people away from and upwind of spill/leak. Ensure adequate ventilation, especially in confined areas. Keep away from heat and sources of ignition. For emergency responders: Personal protection see section 8.

6.2. Environmental precautions

Prevent further leakage or spillage. Do not discharge product into the aquatic environment without pretreatment (biological treatment plant).

6.3. Methods and material for containment and cleaning up

Methods for containment

Stop the flow of material, if possible without risk. Dike spilled material, where this is possible.

Methods for cleaning up

Soak up with inert absorbent material. DO NOT use combustible materials such as sawdust. Keep in suitable,

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closed containers for disposal. If liquid has been spilt in large quantities clean up promptly by scoop or vacuum. Dispose of in accordance with local regulations. Take necessary action to avoid static electricity discharge (which might cause ignition of organic vapours).

6.4. Reference to other sections

For personal protective equipment see section 8.

SECTION 7: Handling and storage

7.1. Precautions for safe handling

Further info may be available in the appropriate Exposure scenarios in the annex to this SDS.

Advice on safe handling

Do not breathe vapours or spray mist. Avoid contact with skin, eyes and clothing. Wash hands before breaks and immediately after handling the product. Do not use compressed air for filling, discharging or handling. Refill and handle product only in closed system. Provide sufficient air exchange and/or exhaust in work rooms.

Hygiene measures

When using, do not eat, drink or smoke. Take off all contaminated clothing immediately. Wash hands before breaks and immediately after handling the product.

Advice on the protection of the environment

See Section 8: Environmental exposure controls.

Incompatible products

acids Halogenated hydrocarbon strong oxidizing agents acid anhydrides acid chlorides

7.2. Conditions for safe storage, including any incompatibilities

Advice on protection against fire and explosion

Keep away from sources of ignition - No smoking. Take necessary action to avoid static electricity discharge (which might cause ignition of organic vapours). In case of fire, emergency cooling with water spray should be available. Ground and bond containers when transferring material. Vapour is heavier than air and can travel considerable distance to a source of ignition and flashback. Vapours may form explosive mixture with air. The pressure in sealed containers can increase under the influence of heat.

Technical measures/Storage conditions

Keep containers tightly closed in a cool, well-ventilated place. Handle and open container with care. Handle under nitrogen, protect from moisture. Containers, storage tanks or drums are having temperature dependent pressure. Vessels with higher temperature must be depressurised into vent gas systems or handled under ventilation.

Suitable material

mild steel, stainless steel

Unsuitable material

Aluminium, copper, zinc, Tin, lead, including their alloys

Temperature class

T2

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7.3. Specific end use(s)

Formulation

For specific end use information see the annex of this safety data sheet

SECTION 8: Exposure controls / personal protection

8.1. Control parameters

Exposure limits European Union

No exposure limits established

Exposure limits Ireland

Ireland OELs

Component	TWA (mg/m³)	TWA (ppm)	STEL (mg/m ³)	STEL (ppm)	Skin Absorption	Sensitizer
Isopropylamine CAS: 75-31-0	12	5	24	10		

Notes

For details and further information please refer to the original regulation.

DNEL & PNEC

Isopropylamine, CAS: 75-31-0 Workers

DN(M)EL - long-term exposure - systemic effects - Inhalation DN(M)EL - acute / short-term exposure - systemic effects - Inhalation

DN(M)EL - long-term exposure - local effects - Inhalation

DN(M)EL - acute / short-term exposure - local effects - Inhalation

DN(M)EL - long-term exposure - systemic effects - Dermal

DN(M)EL - acute / short-term exposure - systemic effects - Dermal

- DN(M)EL long-term exposure local effects Dermal
- DN(M)EL acute / short-term exposure local effects Dermal
- DN(M)EL local effects eyes

General population

DN(M)EL - long-term exposure - systemic effects - Inhalation DN(M)EL - acute / short-term exposure - systemic effects - Inhalation DN(M)EL - long-term exposure - local effects - Inhalation DN(M)EL - acute / short-term exposure - local effects - Inhalation DN(M)EL - long-term exposure - systemic effects - Dermal DN(M)EL - acute / short-term exposure - systemic effects - Dermal DN(M)EL - long-term exposure - local effects - Dermal 10 mg/m³ Medium hazard (no threshold derived) 12 mg/m³ 24 mg/m³ 1.9 mg/kg bw/day Medium hazard (no threshold derived) High hazard (no threshold derived) High hazard (no threshold derived) Medium hazard (no threshold derived)

No hazard identified No hazard identified

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DN(M)EL - acute / short-term exposure - local effects - Dermal DN(M)EL - long-term exposure - systemic effects - Oral DN(M)EL - acute / short-term exposure - systemic effects - Oral DN(M)EL - local effects - eyes

Environment

PNEC aqua - freshwater PNEC aqua - marine water PNEC aqua - intermittent releases PNEC STP PNEC sediment - freshwater PNEC sediment - marine water PNEC Air PNEC soil Secondary poisoning

8.2. Exposure controls

Special adaptations (REACh) Not applicable.

Appropriate Engineering controls

General or dilution ventilation is frequently insufficient as the sole means of controlling employee exposure. Local ventilation is usually preferred. Explosion-proof equipment (for example fans, switches, and grounded ducts) should be used in mechanical ventilation systems.

Personal protective equipment

General industrial hygiene practice

Avoid contact with skin, eyes and clothing. Do not breathe vapours or spray mist. Ensure that eyewash stations and safety showers are close to the workstation location.

Hygiene measures

When using, do not eat, drink or smoke. Take off all contaminated clothing immediately. Wash hands before breaks and immediately after handling the product.

Eye protection

Tightly fitting safety goggles. In addition to goggles, wear a face shield if there is a reasonable chance for splash to the face.

Equipment should conform to EN 166

Hand protection

Wear protective gloves. Recommendations are listed below. Other protective material may be used, depending on the situation, if adequate degradation and permeation data is available. If other chemicals are used in conjunction with this chemical, material selection should be based on protection for all chemicals present.

Suitable material	butyl-rubber
Evaluation	according to EN 374: level 2
Glove thickness	approx 0,3 mm
Break through time	approx 20 min
Suitable material Evaluation Glove thickness	polyvinylchloride Information derived from practical experience approx 0,8 mm



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No hazard identified No hazard identified No hazard identified No hazard identified

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19 µg/l 1,9 µg/l 0,19 mg/l 10 mg/l 161,5 µg/kg dw 16,15 µg/kg dw No hazard identified 21,15 mg/kg No potential for bioaccumulation

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Skin and body protection

Impervious clothing. Wear face-shield and protective suit for abnormal processing problems.

Respiratory protection

Respirator with K- filter. Full mask with above mentioned filter according to producers using requirements or self-contained breathing apparatus. Equipment should conform to EN 136 or EN 140 and EN 143.

Environmental exposure controls

Use product only in closed system. If leakage can not be prevented, the substance needs to be suck off at the emersion point, if possible without danger. Observe the exposure limits, clean exhaust air if needed. If recycling is not practicable, dispose of in compliance with local regulations. Inform the responsible authorities in case of leakage into the atmosphere, or of entry into waterways, soil or drains.

Additional advice

Further details on substance data can be found in the registration dossier under the following link: http://echa.europa.eu/information-on-chemicals/registered-substances. For specific exposure controls see the annex to this safety data sheet.

SECTION 9: Physical and chemical properties

9.1. Information on basic physical and chemical properties

Physical state Colour Odour Odour threshold Melting point/free Method Boiling point or ir point and boiling Method Flammability Lower explosion I Flash point Method Autoignition temp Method Decomposition te pH Kinematic Viscos Method Solubility Partition coefficie n-octanol/water (I	hitial boiling range limit limit berature emperature ity	liquid*** colourless ammonia-like 1,2 ppm < -90 °C (PouDIN ISO 301632 °C @ 1013OECD 103Ignitable2 Vol %11,5 Vol % $<= -25 °C @closed cup, IS355 °C @ 10DIN 51794No data avail13,1 (50 g/l i0,470 mm2/sOECD 114***miscible, in w-0,5 @ 25 °C$	ur point) @ 7 5 3 hPa 1013 hPa 50 2719 16 hPa able n water @ 2 @ 20 °C*** ater, OECD	25 °C (77 °F)) DI 105	N 19268
Vapour pressure Values [hPa] 631	Values [kPa] 63,1	0,623	@ °C 20	@ °F 68	Method DIN EN 13016-2
770 Density and/or re l Values 0,6871	77,3 lative density @ 2		25 @ °F 68	77 Method DIN 51757	DIN EN 13016-2

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Relative vapour density Particle characteristics

2,04 (Air = 1) @ 20 °C (68 °F) not applicable

9.2. Other information

Explosive properties

Oxidizing properties

Molecular weight **Molecular formula** log Koc **Dissociation constant Refractive index** Surface tension **Evaporation rate** hygroscopic.

Does not apply, substance is not explosive. There are no chemical groups associated with explosive properties Does not apply, substance is not oxidising. There are no chemical groups associated with oxidizing properties 59,11 C3 H9 N 1,64 OECD 106 read across pKa 10,8 @ 23,5 °C (74,3 °F) OECD 112 1,373 @ 20 °C 68,5 mN/m (1 g/l @ 20°C (68°F)), OECD 115 No data available

SECTION 10: Stability and Reactivity

10.1. Reactivity

The reactivity of the product corresponds to the typical reactivity shown by the substance group as described in any text book on organic chemistry.

10.2. Chemical stability

Stable under recommended storage conditions.

10.3. Possibility of hazardous reactions

Vapours may form explosive mixture with air.

10.4. Conditions to avoid

Avoid contact with heat, sparks, open flame and static discharge. Avoid any source of ignition.

10.5. Incompatible materials

acids, strong oxidizing agents, halogenated hydrocarbon, acid anhydrides, acid chlorides.

10.6. Hazardous decomposition products

No decomposition if stored and applied as directed. If heated to thermal decomposition the following decomposition products may occur depending on the conditions. carbon monoxide (CO). nitrogen oxides (NOx). cyanides. nitric acid. nitriles.

SECTION 11: Toxicological information

11.1. Information on hazard classes as defined in Regulation (EC) No 1272/2008

Likely routes of exposure Ingestion, Inhalation, Eye contact, Skin contact

Acute toxicity Isopropylamine (75-31-0) according to Regulation (EC) No. 1907/2006 (REACh) Article 31, Annex II as amended



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Routes of Exposure	Endpoint	Values	Species	Method
Oral	LD50	< 173 mg/kg	rat, male	OECD 425
Dermal	LD50	> 400 mg/kg	rat, male/female	OECD 402
Inhalative	LC50	8,7 mg/l (4h)	rat, male/female	OECD 403

Isopropylamine, CAS: 75-31-0

Assessment

The available data lead to the classification given in section 2

Irritation and corrosion

Isopropylamine (75-31-0)					
Target Organ Effects	Species	Result	Method		
Skin	rabbit	corrosive	OECD 404	3 min	
Eyes	rabbit	corrosive	OECD 405	24h	
Respiratory tract	mouse	RD50: 157 ppm	ASTM 981-84	15 min	

Isopropylamine, CAS: 75-31-0

Assessment

The available data lead to the classification given in section 2

Sensitization				
Isopropylamine (75-31-0)				
Target Organ Effects	Species	Evaluation	Method	
Skin	guinea pig	not sensitizing		10 %, aqueous solution

Isopropylamine, CAS: 75-31-0

Assessment

Based on available data, the classification criteria are not met for: Skin sensitization

For respiratory sensitization, no data are available

Subacute, subchronic and prolonged toxicity					
Isopropylamine (75-31-0)					
Туре	Dose	Species	Method		
Subchronic toxicity	NOAEC: 500 mg/m ³ (90 d)	rat, male/female	OECD 413	Inhalation	

Isopropylamine, CAS: 75-31-0

Assessment

Based on available data, the classification criteria are not met for: STOT RE

Carcinogenicity, Mutagenicity, Reproductive toxicity						
Isopropylamine (75-31	-0)					
Туре	Dose	Species	Evaluation	Method		
Developmental Toxicity	NOAEC: 1000 mg/m ³	rat		OECD 414	Teratogenicity Inhalation	
Developmental Toxicity	NOAEC: 500 mg/m ³	rat		OECD 414	Maternal toxicity Inhalation	
Mutagenicity		mouse lymphoma cells	negative (with metabolic activation)	OECD 476 (Mammalian Gene Mutation)	In vitro study	
Mutagenicity		mouse	negative (without	OECD 476	In vitro study	

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		lymphoma cells	metabolic activation)	(Mammalian Gene Mutation)	
Mutagenicity		Salmonella typhimurium	negative (with metabolic activation)	OECD 471 (Ames)	In vitro study
Mutagenicity		Salmonella typhimurium	negative (without metabolic activation)	OECD 471 (Ames)	In vitro study
Mutagenicity		human lymphocytes	negative (with metabolic activation)	OECD 473 (Chromosomal Aberration)	In vitro study
Mutagenicity		human lymphocytes	negative (without metabolic activation)	OECD 473 (Chromosomal Aberration)	In vitro study
Reproductive toxicity	NOAEC: 500 mg/m ³	rat, parental		OECD 415	Inhalation
Reproductive toxicity	NOAEC: 500 mg/m³	rat, 1. Generation, male/female		OECD 415	Inhalation

Isopropylamine, CAS: 75-31-0

CMR Classification

The available data on CMR properties are summarized in the table above. They do not indicate a classification into categories 1A or 1B

Evaluation

In vitro tests did not show mutagenic effects Animal testing did not show any effects on fertility In the absence of specific alerts no cancer testing is required

Isopropylamine, CAS: 75-31-0

Main symptoms

shortness of breath, convulsions, cough, hypertensive effect, narcosis, unconsciousness, discomfort, nausea. Target Organ Systemic Toxicant - Single exposure

STOT SE

respiratory system

The available data lead to the classification given in section 2

Target Organ Systemic Toxicant - Repeated exposure

Based on available data, the classification criteria are not met for: STOT RE

11.2. Information on other hazards

Endocrine disrupting properties

The substance has not been identified as having endocrine disrupting properties in accordance with section 2.3. **Isopropylamine, CAS: 75-31-0**

Other adverse effects

Components of the product may be absorbed into the body by inhalation, ingestion and through the skin. **Note**

Handle in accordance with good industrial hygiene and safety practice. Further details on substance data can be found in the registration dossier under the following link:

http://echa.europa.eu/information-on-chemicals/registered-substances.

SECTION 12: Ecological information

according to Regulation (EC) No. 1907/2006 (REACh) Article 31, Annex II as amended



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12.1. Toxicity

Acute aquatic toxicity			
Isopropylamine (75-31-0)			
Species	Exposure time	Dose	Method
Daphnia magna (Water flea)	48h	EC50: 47,4 mg/l	79/831/EEC.C2
Desmodesmus subspicatus	72h	EC50: 18,9 mg/l (Growth rate)	DIN 38412, part 9
Oncorhynchus mykiss (rainbow trout)	96h	LC50: 40 mg/l	OECD 203
Activated sludge (domestic)	30 min	EC50: >1000 mg/l (Growth inhibition)	OECD 209

Long term toxicity				
Isopropylamine (75-31-0)				
Туре	Species	Dose	Method	
Aquatic toxicity	Desmodesmus subspicatus	NOEC: 1,25 mg/l (3d) Growth inhibition	DIN 38412 / part 9	

12.2. Persistence and degradability

Isopropylamine, CAS: 75-31-0

Biodegradation

70 - 80 % (28 d), activated sludge, aerobic, domestic, OECD 301 F.

Abiotic Degradation

ADIOLIC Degradation			
Isopropylamine (75-31-0)			
Туре	Result	Method	
Hydrolysis	not expected		
Photolysis	No data available		

12.3. Bioaccumulative potential

Isopropylamine (75-31-0)		
Туре	Result	Method
log Pow	-0,5 @ 25 °C (77 °F)	measured, OECD 117
BCF	not expected	

12.4. Mobility in soil

Isopropylamine (75-31-0)		
Туре	Result	Method
Surface tension	68,5 mN/m (1 g/l @ 20°C (68°F))	OECD 115
Adsorption/Desorption	Koc: 43,2	OECD 106 read across
Distribution to environmental	no data available	
compartments		

12.5. Results of PBT and vPvB assessment

according to Regulation (EC) No. 1907/2006 (REACh) Article 31, Annex II as amended



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Isopropylamine, CAS: 75-31-0

PBT and vPvB assessment

This substance is not considered to be persistent, bioaccumulating nor toxic (PBT), nor very persistent nor very bioaccumulating (vPvB)

12.6. Endocrine disrupting properties

The substance has not been identified as having endocrine disrupting properties in accordance with section 2.3.

12.7. Other adverse effects

Isopropylamine, CAS: 75-31-0 No data available

SECTION 13: Disposal considerations

13.1. Waste treatment methods

Product Information

Disposal required in compliance with all waste management related state and local regulations. The choice of the appropriate method of disposal depends on the product composition by the time of disposal as well as the local statutes and possibilities for disposal.

Hazardous waste according to European Waste Catalogue (EWC)

Uncleaned empty packaging

Contaminated packaging should be emptied as far as possible and after appropriate cleansing may be taken for reuse.

SECTION 14: Transport information

ADR/RID

 14.1. UN number or ID number 14.2. UN proper shipping name 14.3. Transport hazard class(es) Subsidiary Risk 14.4. Packing group 14.5. Environmental hazards 14.6. Special precautions for user ADR Tunnel restriction code Classification Code Hazard Number 	UN 1221 Isopropylamine 3 8 I no (C/E) FC 338
ADN	ADN Container
 14.1. UN number or ID number 14.2. UN proper shipping name 14.3. Transport hazard class(es) Subsidiary Risk 14.4. Packing group 	UN 1221 Isopropylamine 3 8 I

Isopropylamine

according to Regulation (EC) No. 1907/2006 (REACh) Article 31, Annex II as amended



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14.5. Environmental hazards	no
14.6. Special precautions for user	
Classification Code	FC
Hazard Number	338
ICAO-TI / IATA-DGR	
14.1. UN number or ID number	UN 1221
14.2. UN proper shipping name	Isopropylamine
14.3. Transport hazard class(es)	3
Subsidiary Risk	8
14.4. Packing group	I
14.5. Environmental hazards	no
14.6. Special precautions for user	no data available
IMDG_	
14.1. UN number or ID number	UN 1221
14.2. UN proper shipping name	Isopropylamine
14.3. Transport hazard class(es)	3
Subsidiary Risk	8
14.4. Packing group	I
14.5. Environmental hazards	no
14.6. Special precautions for user	
EmS	F-E, S-C
14.7. Maritime transport in bulk according	***
to IMO instruments	
Product name	Isopropylamine
Ship type	2
Pollution category Hazard class	Y S/P***

SECTION 15: Regulatory information

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

Regulation 1272/2008, Annex VI

Isopropylamine, CAS: 75-31-0	
Classification	Flam. Liq. 1; H224 Eye Irrit. 2; H319 STOT SE 3; H335
Hazard pictograms	Skin Irrit. 2; H315 GHS02 Flame GHS07 Exclamation mark
Signal word	Danger
Hazard statements	H224, H319, H335, H315
<u>DI 2012/18/EU (Seveso III)</u> Category	Annex I, part 1:

according to Regulation (EC) No. 1907/2006 (REACh) Article 31, Annex II as amended



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H2

P5a - c; depending on conditions

DI 1999/13/EC (VOC Guideline)

Component	Status
Isopropylamine	regulated
CAS: 75-31-0	

International Inventories

Isopropylamine, CAS: 75-31-0

AICS (AU) DSL (CA) **IECSC (CN)** EC-No. 2008609 (EU) ENCS (2)-131 (JP) ISHL (2)-131 (JP) KECI KE-29257 (KR) INSQ (MX) PICCS (PH) TSCA (US) NZIoC (NZ) TCSI (TW)

15.2. Chemical safety assessment

The Chemical Safety Report (CSR) has been generated. For Exposure Scenarios see the annex.

SECTION 16: Other information

Full text of H-Statements referred to under sections 2 and 3

H224: Extremely flammable liquid and vapour.

- H301: Toxic if swallowed.
- H311: Toxic in contact with skin.
- H331: Toxic if inhaled.
- H315: Causes skin irritation.
- H319: Causes serious eye irritation.

H335: May cause respiratory irritation.

Abbreviations

A table of terms and abbreviations can be found under the following link: http://echa.europa.eu/documents/10162/13632/information_requirements_r20_en.pdf

Training advice

For effective first-aid, special training / education is needed.

Sources of key data used to compile the datasheet

Information contained in this safety data sheet is based on OQ owned data and public sources deemed valid or acceptable. The absence of data elements required by OSHA, ANSI or Annex II, Regulation 1907/2006/EC indicates, that no data meeting these requirements is available.

Further information for the safety data sheet

according to Regulation (EC) No. 1907/2006 (REACh) Article 31, Annex II as amended



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Changes against the previous version are marked by ***. Observe national and local legal requirements. For more information, other material safety data sheets or technical data sheets please consult the OQ homepage (www.chemicals.oq.com).

Disclaimer

For industrial use only. The information contained herein is accurate to the best of our knowledge. We do not suggest or guarantee that any hazards listed herein are the only ones which exist. OQ Chemicals makes no warranty of any kind, express or implied, concerning the safe use of this material in your process or in combination with other substances. User has the sole responsibility to determine the suitability of the materials for any use and the manner of use contemplated. User must meet all applicable safety and health standards.

End of Safety Data Sheet

Annex to the extended Safety Data Sheet (eSDS)

General information

A quantitative approach used to conclude safe use for: Environmental compartment Long-term Systemic effects via inhalation Long term local hazards via inhalation Acute local hazards via inhalation A qualitative approach used to conclude safe use for: Acute systemic hazards via skin Acute local hazards via skin Long term local hazards via skin Acute systemic hazards via skin Local hazards via skin Local hazards via eyes

Operational conditions and risk management measures

Following operational conditions and risk management measures, are based on qualitative risk characterisation: Wear suitable face shield. Substance/task appropriate gloves Full skin coverage with appropriate light-weight barrier material

Chemical goggles or safety glasses Any measure to eliminate exposure should be considered

Containment of source except for short term exposure (e.g. taking sample)

Design closed system to allow for easy maintenance

If possible keep equipment under negative pressure

Control staff entry to work area

Ensure all equipment well maintained

Permit to work for maintenance work

Regular cleaning of equipment and work area

Training for staff on good practice

Procedures and training for emergency decontamination and disposal

Good standard of personal hygiene

Recording of any 'near miss' situations

Supervision in place to check that the RMMs in place are being used correctly and OCs followed.

Exposure scenario identification

1 Formulation & (re)packing of substances and mixtures Exposure estimation and reference to its source

according to Regulation (EC) No. 1907/2006 (REACh) Article 31, Annex II as amended



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Short title of the exposure scenario

Formulation & (re)packing of substances and mixtures

Sector of uses [SU]

SU3: Industrial uses: Uses of substances as such or in preparations at industrial sites SU10: Formulation [mixing] of preparations and/or re-packaging (excluding alloys)

Process categories [PROC]

PROC1: Use in closed process, no likelihood of exposure

PROC2: Use in closed, continuous process with occasional controlled exposure

PROC3: Use in closed batch process (synthesis or formulation)

PROC4: Use in batch and other process (synthesis) where opportunity for exposure arises

PROC5: Mixing or blending in batch processes for formulation of preparations and articles (multistage and/or significant contact)

PROC8a: Transfer of substance or preparation (charging/discharging) from/to vessels/large containers at non-dedicated facilities

PROC8b: Transfer of substance or preparation (charging/discharging) from/to vessels/large containers at dedicated facilities PROC9: Transfer of substance or preparation into small containers (dedicated filling line, including weighing) PROC15: Use as laboratory reagent

Processes and activities covered by the exposure scenario

Formulation, packing and re-packing of the substance and its mixtures in batch or continuous operations, including storage, materials transfers, mixing, tabletting, compression, pelletisation, extrusion, large and small scale packing, sampling, maintenanance and associated laboratory activities.

Further explanations

Industrial use Assessment tool used: Chesar 3.5 liquid Assumes use at not more than 20°C above ambient temperature (unless stated differently) Covers percentage substance in the product up to 100 % (unless stated differently). Assumes an advanced standard of occupational Health and Safety Management System Number of the contributing scenario Contributing exposure scenario controlling environmental exposure for ERC 2

Further specification

assessment tool used:, Chesar 3.5, release factors for (Sp)ERC were modified. Amounts used Daily amount per site: 10 to Annual amount per site: 1000 to Fraction of Regional tonnage used locally: 1 Technical conditions and measures at process level (source) to prevent release Release fraction to air from process: 2,5% Release fraction to wastewater from process: 0.025% Release fraction to soil from process: 0,01% Conditions and measures related to municipal sewage treatment plant Size of industrial sewage treatment plant (m3/d): 2000 The minimum grade of elimination in the sewage plant is (%): 87,74 Do not apply industrial sludge to natural soils Number of the contributing scenario 2 Contributing exposure scenario controlling worker exposure for PROC 1

Frequency and duration of use 8 h (full shift)

according to Regulation (EC) No. 1907/2006 (REACh) Article 31, Annex II as amended



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Other given operational conditions affecting work Indoor and outdoor use Technical conditions and measures to control dis provide a good standard of general ventilation (not les Conditions and measures related to personal prot Wear suitable gloves (tested to EN374) and eye prote	persion from source towards the worker as than 3 to 5 air changes per hour). aection, hygiene and health evaluation
Number of the contributing scenario Contributing exposure scenario controlling v PROC 2	3 vorker exposure for
Frequency and duration of use 8 h (full shift) Other given operational conditions affecting work Indoor use Technical conditions and measures to control dis Effectiveness of LEV (local exhaust ventilation): 90 % ventilation (not less than 3 to 5 air changes per hour). Conditions and measures related to personal prof Wear suitable gloves (tested to EN374) and eye prote Number of the contributing scenario Contributing exposure scenario controlling v PROC 3	persion from source towards the worker (inhalative); 90 % (dermal). provide a good standard of general ection, hygiene and health evaluation ection. Wear respiratory protection (Efficiency: 95 %). 4
Frequency and duration of use 8 h (full shift) Other given operational conditions affecting work Indoor use Technical conditions and measures to control dis Effectiveness of LEV (local exhaust ventilation): 90 % ventilation (not less than 3 to 5 air changes per hour). Conditions and measures related to personal prof Wear suitable gloves (tested to EN374) and eye prote Number of the contributing scenario Contributing exposure scenario controlling w PROC 4	persion from source towards the worker (inhalative); 90 % (dermal). provide a good standard of general ection, hygiene and health evaluation ection. Wear respiratory protection (Efficiency: 95 %). 5
ventilation (not less than 3 to 5 air changes per hour). Conditions and measures related to personal prof	 persion from source towards the worker 6 (inhalative); 90 % (dermal). provide a good standard of general cection, hygiene and health evaluation a combination with specific activity training. Wear respiratory protection 1374) and eye protection.
Frequency and duration of use 8 h (full shift) Other given operational conditions affecting work Indoor use Technical conditions and measures to control dis Effectiveness of LEV (local exhaust ventilation): 90 %	•

ventilation (5 to 10 air changes per hour) .

Conditions and measures related to personal protection, hygiene and health evaluation Wear chemically resistant gloves (tested to EN374) in combination with specific activity training. Wear respiratory protection (Efficiency: 95 %). Wear suitable gloves (tested to EN374) and eye protection.

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Number of the contributing scenario Contributing exposure scenario controlling wor PROC 8a	7 ker exposure for
Frequency and duration of use 8 h (full shift)	
Other given operational conditions affecting workers Indoor use	
Technical conditions and measures to control disper- Effectiveness of LEV (local exhaust ventilation): 90 % (inf ventilation (5 to 10 air changes per hour).	sion from source towards the worker nalative); 90 % (dermal). provide a good standard of controlled
Conditions and measures related to personal protect	nically resistant gloves (tested to EN374) in combination with specific
Number of the contributing scenario	8
Contributing exposure scenario controlling wor PROC 8b	ker exposure for
Frequency and duration of use	
8 h (full shift) Other given operational conditions affecting workers Indoor use	exposure
Technical conditions and measures to control disper- Effectiveness of LEV (local exhaust ventilation): 95 % (inf ventilation (not less than 3 to 5 air changes per hour).	nalative); 95 % (dermal). provide a good standard of general
Conditions and measures related to personal protect Wear suitable gloves and eye/face protection. Wear chem activity training. Wear respiratory protection (Efficiency: 9	nically resistant gloves (tested to EN374) in combination with specific
Number of the contributing scenario	9
Contributing exposure scenario controlling wor PROC 9	ker exposure for
Product characteristics Covers percentage substance in the product up to 100 %	(unloss stated differently)
Frequency and duration of use 8 h (full shift)	(unless stated unerently)
Other given operational conditions affecting workers Indoor use	·
Technical conditions and measures to control disper Effectiveness of LEV (local exhaust ventilation): 90 % (in ventilation (not less than 3 to 5 air changes per hour).	sion from source towards the worker halative); 90 % (dermal). provide a good standard of general
Conditions and measures related to personal protect Wear suitable gloves and eye/face protection. Wear chem	nically resistant gloves (tested to EN374) in combination with specific
activity training. Wear respiratory protection (Efficiency: 9 Number of the contributing scenario	5 %). 10
Contributing exposure scenario controlling wor PROC 15	
Product characteristics Liquid	
Frequency and duration of use 1 h per shift	
Other given operational conditions affecting workers Indoor use	
ventilation): 90 % (inhalative); 90 % (dermal).	air changes per hour) . Effectiveness of LEV (local exhaust
Conditions and measures related to personal protect Wear chemically resistant gloves (tested to EN374) in con-	
Number of the contributing scenario	11

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Contributing exposure scenario controlling worker exposure for PROC 8a

Product characteristics Liquid Frequency and duration of use 1 h per shift Other given operational conditions affecting workers exposure Indoor use Technical conditions and measures to control dispersion from source towards the worker provide a basic standard of general ventilation (1 to 3 air changes per hour). Effectiveness of LEV (local exhaust ventilation): 90 % (inhalative). Conditions and measures related to personal protection, hygiene and health evaluation Wear respiratory protection (Efficiency: 95 %). Wear chemically resistant gloves (tested to EN374) in combination with specific activity training. Fresh Water (Pelagic) PEC: 0,015 mg/l; RCR: 0,806 Fresh Water (Sediment) PEC: 0.121 mg/kg dw; RCR: 0.751 Marine Water (Pelagic) PEC: 1.53E-3 mg/l; RCR: 0.806 Marine Water (Sediment) PEC: 0.012 mg/kg dw; RCR: 0.751 Agricultural Soil PEC: 3.68E-3 mg/kg dw; RCR: 0.174 PEC: 0,153 mg/l; RCR: 0.015 Sewage Treatment Plant (Effluent) Man via environment – Inhalation Concentration in air: 0.019 mg/m3: RCR: 0.011 Man via environment – Oral Exposure via food consumption: 4,68E-4 mg/kg bw/day; RCR: 0,01 Man via environment - combined RCR: 0,011 routes

Human exposure prediction (oral, dermal, inhalative)

Oral exposure is not expected to occur. EE(inhal): Estimated inhalative exposure [mg/m³]. EE(derm): Estimated dermal exposure [mg/kg b.w./d]. Exposure estimates are given for short-term or long-term, systemic or local exposure depending on which lead to more conservative risk characterization ratios. The RMMs described above suffice to control risks for both local and systemic effects.

EE(inhal): 0,069; EE(derm): 6.8E-3
EE(inhal): 0,862; EE(derm): 0,027
EE(inhal): 1,724; EE(derm): 0,014
EE(inhal): 3,448; EE(derm): 0,034
EE(inhal): 3,694; EE(derm): 0,069
EE(inhal): 3.694; EE(derm): 0.069 - Contributing Scenarios 7
EE(inhal): 12.31; EE(derm): 0.137 - Contributing Scenarios 11
EE(inhal): 2,586; EE(derm): 0,034
EE(inhal): 6,896; EE(derm): 0.034
EE(inhal): 14.77; EE(derm): 1.36E-3

Risk characterisation

RCR(inhal): inhalative risk characterisation ratio; RCR(derm): dermal risk characterisation ratio; total RCR= RCR(inhal) +RCR(derm). Where required local and systemic effects were evaluated both for short-term and long-term exposure. The RCR's given correspond in each case to the most conservative calculated values.

Proc 1	RCR(inhal): 0,01; RCR(derm): 0,01
Proc 2	RCR(inhal): 0,036; RCR(derm): 0,014
Proc 3	RCR(inhal): 0,072; RCR(derm): 0,01
Proc 4	RCR(inhal): 0,144; RCR(derm): 0,018
Proc 5	RCR(inhal): 0,154; RCR(derm): 0,036
Proc 8a	RCR(inhal): 0,154; RCR(derm): 0,036 - Contributing Scenarios 7
	RCR(inhal): 0.513; RCR(derm): 0.072 - Contributing Scenarios 11
Proc 8b	RCR(inhal): 0,108; RCR(derm): 0,018
Proc 9	RCR(inhal): 0,287; RCR(derm): 0,018
Proc 15	RCR(inhal): 0.616; RCR(derm): 0,01

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Guidance to Downstream User to evaluate whether he works inside the boundaries set by the ES

Usage of relase factors allows downstream users to verify in a first approximation, if the combination of local usage and production conditions meets the defined release quantities resulting from this exposure scenario (calculated as M(site) [see amounts used, contributing scenario 1] x release factor [Technical conditions and measures at process level (source) to prevent release; contributing scenario 1])

associated uses:

Other combinations of operational conditions may also be safe. Please contact OQ in case your local operational conditions differ from the ones described above and you are unsure if they are also safe

Human exposure prediction (oral, dermal, inhalative)

Oral exposure is not expected to occur. EE(inhal): Estimated inhalative long-term exposure [mg/m³]; EE(derm): Estimated dermal long-term exposure [mg/kg b.w./d]. Exposure estimates are given for either short-term or long-term exposure depending on which lead to more conservative risk characterisation ratios. The RMMs described above suffice to control risks for both local and systemic effects.

Proc 1	EE(inhal): 0.025 ; EE(derm): 0.069
Proc 2	EE(inhal): 2.463 ; EE(derm): 0.027
Proc 3	EE(inhal): 6.157 ; EE(derm): 0.007
Proc 4	EE(inhal): 4.926 ; EE(derm): 0.137
Proc 8a	EE(inhal): 7.389 ; EE(derm): 0.027
Proc 8b	EE(inhal): 3.694 ; EE(derm): 0.137
Proc 9	EE(inhal): 1.231 ; EE(derm): 0.137
Proc 15	EE(inhal): 2.463 ; EE(derm): 0.007

Risk characterisation

RCR(inhal): inhalative risk characterisation ratio; RCR(derm): dermal risk characterisation ratio; total RCR= RCR(inhal) +RCR(derm). Where required local and systemic effects were evaluated both for short-term and long-term exposure. The RCR's given correspond in each case to the most conservative calculated values.

Proc 1 Proc 2	RCR(inhal): 0.002 ; RCR(derm): 0.014 RCR(inhal): 0.205 ; RCR(derm): 0.006
Proc 3	RCR(inhal): 0.513 ; RCR(derm): 0.001
Proc 4	RCR(inhal): 0.411 ; RCR(derm): 0.029
Proc 8a	RCR(inhal): 0.616 ; RCR(derm): 0.006
Proc 8b	RCR(inhal): 0.308 ; RCR(derm): 0.029
Proc 9	RCR(inhal): 0.103 ; RCR(derm): 0.029
Proc 15	RCR(inhal): 0.205 ; RCR(derm): 0.001

Guidance to Downstream User to evaluate whether he works inside the boundaries set by the ES Usage of relase factors allows downstream users to verify in a first approximation, if the combination of local usage and production conditions meets the defined release quantities resulting from this exposure scenario (calculated as M(site) [see amounts used, contributing scenario 1] x release factor [Technical conditions and measures at process level (source) to prevent release; contributing scenario 1])

associated uses:

Other combinations of operational conditions may also be safe. Please contact OQ in case your local operational conditions differ from the ones described above and you are unsure if they are also safe

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