

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

1.1. Product identifier

Trade name or designation of the mixture ADVAIR HFA

Registration number -

Synonyms ADVAIR HFA INHALATION AEROSOL * SERETIDE INHALER HFA * SERETIDE EVOHALER * BREXOTIDE INHALER HFA 134A * FLIXOVENT INHALER HFA 134A * SERETAIDE INHALER HFA 134A * VIANI EVOHALER * VIANI INHALER HFA * VIANI MITE 25 MCG/50 MCG DOSIER-AEROSOL FCKW-FREI * VIANI 25 MCG/125 MCG DOSIER-AEROSOL FCKW-FREI * VIANI FORTE 25 MCG/250 MCG DOSIER-AEROSOL FCKW-FREI * SALMETEROL/FLUTICASONE PROPIONATE INHALATION AEROSOL * SALMETEROL/FLUTICASONE PROPIONATE INHALER 25/50 MCG 120 ACTN * SALMETEROL/FLUTICASONE PROPIONATE INHALER 25/125 MCG 120 ACTN * SALMETEROL/FLUTICASONE PROPIONATE INHALER 25/250 MCH 120 ACTN * SALMETEROL/FLUTICASONE PROPIONATE 134A 120 ACTN * SALMETEROL XINAFOATE AND FLUTICASONE PROPIONATE, FORMULATED PRODUCT

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

Identified uses Medicinal Product

This safety data sheet is written to provide health, safety and environmental information for people handling this formulated product in the workplace. It is not intended to provide information relevant to medicinal use of the product. In this instance patients should consult prescribing information/package insert/product label or consult their pharmacist or physician. For health and safety information for individual ingredients used during manufacturing, refer to the appropriate safety data sheet for each ingredient.

Uses advised against No other uses are advised.

1.3. Details of the supplier of the safety data sheet

GlaxoSmithKline UK
980 Great West Road
Brentford, Middlesex TW8 9GS UK
UK General Information (normal business hours): +44-20-8047-5000
Email Address: msds@gsk.com
Website: www.gsk.com

1.4. Emergency telephone number

TRANSPORT EMERGENCIES::
UK In-country toll call: +(44)-870-8200418
International toll call: +1 703 527 3887
available 24 hrs/7 days; multi-language response

SECTION 2: Hazards identification

2.1. Classification of the substance or mixture

Classification according to Directive 67/548/EEC or 1999/45/EC as amended

Exempt from requirements - product regulated as a medicinal product, cosmetic product or medical device.

Classification according to Regulation (EC) No 1272/2008 as amended

Exempt from requirements - product regulated as a medicinal product, cosmetic product or medical device.

2.2. Label elements

Label according to Regulation (EC) No. 1272/2008 as amended

Exempt from requirements - product regulated as a medicinal product, cosmetic product or medical device.

Supplemental label information Not applicable.

2.3. Other hazards

This product is non-flammable. Aerosol containers may violently rupture when exposed to the heat of fire.
Caution - Pharmaceutical agent. See section 11 for additional information on health hazards.

SECTION 3: Composition/information on ingredients

3.2. Mixtures

General information

Chemical name	%	CAS-No. / EC No.	REACH Registration No.	INDEX No.	Notes
1,1,1,2-TETRAFLUOROETHANE	99.6 - 99.87	811-97-2 212-377-0	-	-	
Classification:	DSD: - CLP: -				
FLUTICASONE PROPIONATE	0.08 - 0.34	80474-14-2 -	-	-	
Classification:	DSD: Repr. Cat. 2;R61, Repr. Cat. 3;R62, Xn;R48/20/21, R53 CLP: Repr. 1B;H360D, Repr. 2;H361f, STOT RE 2;H373				
SALMETEROL XINAFOATE	0.05	94749-08-3 -	-	-	
Classification:	DSD: Xi;R36/38, N;R51/53 CLP: Skin Irrit. 2;H315, Eye Irrit. 2;H319				

CLP: Regulation No. 1272/2008.

DSD: Directive 67/548/EEC.

M: M-factor

vPvB: very persistent and very bioaccumulative substance.

PBT: persistent, bioaccumulative and toxic substance.

#: This substance has been assigned Community workplace exposure limit(s).

Composition comments The full text for all R- and H-phrases is displayed in section 16.

SECTION 4: First aid measures

General information If you feel unwell, seek medical advice (show the label where possible). Ensure that medical personnel are aware of the material(s) involved, and take precautions to protect themselves.

4.1. Description of first aid measures

Inhalation	Move to fresh air. Call a physician if symptoms develop or persist.
Skin contact	Wash off with soap and water. Get medical attention if irritation develops and persists.
Eye contact	Rinse thoroughly with plenty of water for at least 15 minutes and consult a physician.
Ingestion	Rinse mouth. Get medical attention if symptoms occur.

4.2. Most important symptoms and effects, both acute and delayed The following adverse effects have been noted with therapeutic use of this material: increased susceptibility to infection; headache; inflamed nasal cavity; back pain; joint pain; coughing; nausea; vomiting.

4.3. Indication of any immediate medical attention and special treatment needed Provide general supportive measures and treat symptomatically.

SECTION 5: Firefighting measures

General fire hazards This product is non-flammable. Aerosol containers may violently rupture when exposed to the heat of fire.

5.1. Extinguishing media

Suitable extinguishing media	Water fog. Foam. Dry chemical powder. Carbon dioxide (CO ₂).
Unsuitable extinguishing media	None known.

5.2. Special hazards arising from the substance or mixture During fire, gases hazardous to health may be formed.

5.3. Advice for firefighters

Special protective equipment for firefighters	Self-contained breathing apparatus and full protective clothing must be worn in case of fire.
Special fire fighting procedures	Use standard firefighting procedures and consider the hazards of other involved materials. In the event of fire, cool tanks with water spray. Move containers from fire area if you can do so without risk.

SECTION 6: Accidental release measures

6.1. Personal precautions, protective equipment and emergency procedures

For non-emergency personnel Keep unnecessary personnel away. Keep people away from and upwind of spill/leak. Wear appropriate personal protective equipment. Ensure adequate ventilation. Local authorities should be advised if significant spillages cannot be contained. For personal protection, see section 8.

For emergency responders Keep unnecessary personnel away. Use personal protection recommended in Section 8 of the MSDS.

6.2. Environmental precautions Avoid release to the environment. Contact local authorities in case of spillage to drain/aquatic environment. Prevent further leakage or spillage if safe to do so. Do not contaminate water. Avoid discharge into drains, water courses or onto the ground.

6.3. Methods and material for containment and cleaning up Stop the flow of material, if this is without risk. Collect spillage. Dike far ahead of spill for later disposal. Prevent product from entering drains. Following product recovery, flush area with water.

Never return spills in original containers for re-use.

6.4. Reference to other sections For personal protection, see section 8. For waste disposal, see section 13.

SECTION 7: Handling and storage

7.1. Precautions for safe handling Avoid prolonged exposure. Provide adequate ventilation. Wear appropriate personal protective equipment. Observe good industrial hygiene practices. Avoid release to the environment. Do not empty into drains.

7.2. Conditions for safe storage, including any incompatibilities Store in original tightly closed container. Store in a cool, dry place out of direct sunlight. Store away from incompatible materials (see Section 10 of the MSDS). The recommended temperature for storage is 15-25 °C.

7.3. Specific end use(s) Medicinal Product

SECTION 8: Exposure controls/personal protection

8.1. Control parameters

Occupational exposure limits

GSK Components	Type	Value	Note
FLUTICASONE PROPIONATE (CAS 80474-14-2)	8 HR TWA	3 mcg/m3	
	OHC	4 4	Skin Reproductive hazard
SALMETEROL XINAFOATE (CAS 94749-08-3)	8 HR TWA	1 mcg/m3	
	OHC	5	
UK. EH40 Workplace Exposure Limits (WELs) Components	Type	Value	
1,1,1,2-TETRAFLUOROETHANE (CAS 811-97-2)	TWA	4240 mg/m3	
		1000 ppm	

Recommended monitoring procedures Follow standard monitoring procedures.

Derived No Effect Level (DNEL) Not available.

Predicted no effect concentrations (PNECs) Not available.

8.2. Exposure controls

Appropriate engineering controls Good general ventilation (typically 10 air changes per hour) should be used. Ventilation rates should be matched to conditions. If applicable, use process enclosures, local exhaust ventilation, or other engineering controls to maintain airborne levels below recommended exposure limits. If exposure limits have not been established, maintain airborne levels to an acceptable level. An Exposure Control Approach (ECA) is established for operations involving this material based upon the OEL/Occupational Hazard Category and the outcome of a site- or operation-specific risk assessment.

Individual protection measures, such as personal protective equipment

General information Personal protection equipment should be chosen according to the CEN standards and in discussion with the supplier of the personal protective equipment. Follow all local regulations if personal protective equipment (PPE) is used in the workplace.

Eye/face protection If contact is likely, safety glasses with side shields are recommended. (eg. EN 166)

Skin protection	
- Hand protection	The choice of an appropriate glove does not only depend on its material but also on other quality features and is different from one producer to the other. Glove selection must take into account any solvents and other hazards present. With respect to the above precautions select suitable chemical resistant protective gloves (EN 374) with a protective index 6 (>480min permeation time).
- Other	Wear suitable protective clothing. (EN 14605 for splashes, EN ISO 13982 for dust)
Respiratory protection	When workers are facing concentrations above the exposure limit they must use appropriate certified respirators. Where breathable aerosols/dust are formed, use suitable combination filter for gases/vapours of organic, inorganic, acid inorganic, alkaline compounds and toxic particles (eg. EN 14387).
Thermal hazards	Wear appropriate thermal protective clothing, when necessary.
Hygiene measures	Always observe good personal hygiene measures, such as washing after handling the material and before eating, drinking, and/or smoking. Routinely wash work clothing and protective equipment to remove contaminants.
Environmental exposure controls	
Hazard guidance and control recommendations	Contain spills and prevent releases and observe national regulations on emissions. Environmental manager must be informed of all major releases.

SECTION 9: Physical and chemical properties

9.1. Information on basic physical and chemical properties

Appearance

Physical state	Solid.
Form	Aerosol
Colour	Not available.
Odour	Not available.
Odour threshold	Not available.
pH	Not available.
Melting point/freezing point	Not available.
Initial boiling point and boiling range	Not available.
Flash point	Not available.
Evaporation rate	Not available.
Flammability (solid, gas)	Not available.
Upper/lower flammability or explosive limits	
Flammability limit - lower (%)	Not available.
Flammability limit - upper (%)	Not available.
Vapour pressure	Not available.
Vapour density	Not available.
Relative density	Not available.
Solubility(ies)	Not available.
Partition coefficient (n-octanol/water)	Not available.
Auto-ignition temperature	Not available.
Decomposition temperature	Not available.
Viscosity	Not available.
Explosive properties	Not available.
Oxidizing properties	Not available.

9.2. Other information No relevant additional information available.

SECTION 10: Stability and reactivity

10.1. Reactivity	The product is stable and non-reactive under normal conditions of use, storage and transport.
10.2. Chemical stability	Material is stable under normal conditions.
10.3. Possibility of hazardous reactions	No dangerous reaction known under conditions of normal use.
10.4. Conditions to avoid	Contact with incompatible materials.

10.5. Incompatible materials

Strong oxidising agents.

10.6. Hazardous decomposition products

Irritating and/or toxic fumes and gases may be emitted upon the products decomposition.

SECTION 11: Toxicological information**General information**

Caution - Pharmaceutical agent. Occupational exposure to the substance or mixture may cause adverse effects.

Information on likely routes of exposure**Ingestion**

Health injuries are not known or expected under normal use.

Inhalation

Prolonged inhalation may be harmful. Health injuries are not known or expected under normal use.

Skin contact

Health injuries are not known or expected under normal use.

Eye contact

Health injuries are not known or expected under normal use.

Symptoms

The following adverse effects have been noted with therapeutic use of this material: increased susceptibility to infection; headache; inflamed nasal cavity; back pain; joint pain; coughing; nausea; vomiting.

11.1. Information on toxicological effects**Acute toxicity**

Health injuries are not known or expected under normal use.

Components**Species****Test results****FLUTICASONE PROPIONATE (CAS 80474-14-2)****Acute***Oral*

LD50

Rat

> 1000 mg/kg

Subacute*Inhalation*

NOAEL

Rat

0.2 mcg/L/day, 28 Day

Subchronic*Inhalation*

LOEL

Rat

3 mcg/kg/day, 26 weeks

NOAEL

Dog

68 mcg/kg/day, 26 weeks

Rat

14 mcg/kg/day, 26 weeks

SALMETEROL XINAFOATE (CAS 94749-08-3)**Acute***Inhalation*

LC50

Rat

> 75 mg/l

Oral

LD50

Rat

> 1000 mg/kg

Subchronic*Inhalation*

LOEL

Rat

>= 0.16 mg/kg/day, 26 weeks, adrenergic effects

Oral

NOAEL

Rat

0.2 mg/kg/day, 26 weeks, adrenergic effects

* Estimates for product may be based on additional component data not shown.

Skin corrosion/irritation

Health injuries are not known or expected under normal use.

Corrosivity

FLUTICASONE PROPIONATE

OECD 404

Result: negative

SALMETEROL XINAFOATE

Result: Irritant

Species: Human

Irritation Corrosion - Skin: P.I.I. value

FLUTICASONE PROPIONATE

0

Serious eye damage/eye irritation

Direct contact with eyes may cause temporary irritation.

Eye		
	SALMETEROL XINAFOATE	OECD 405 Result: Severe Species: Rabbit
	FLUTICASONE PROPIONATE	OECD 405 Result: negative Species: Rabbit
Respiratory sensitisation	Based on available data, the classification criteria are not met.	
Skin sensitisation	This product is not expected to cause skin sensitisation.	
Maximisation assay (Magnusson and Kligman)		
	SALMETEROL XINAFOATE	Result: negative Species: Guinea pig
Sensitisation		
	FLUTICASONE PROPIONATE	0 % OECD 406 Result: negative Species: Guinea pig
Germ cell mutagenicity	Based on available data, the classification criteria are not met.	
Germ cell mutagenicity		
Mutagenicity		
	FLUTICASONE PROPIONATE	Ames Result: negative
	SALMETEROL XINAFOATE	Ames - Screen Result: negative
	FLUTICASONE PROPIONATE	Bacterial High Throughput Fluctuation Test Result: negative Chinese Hamster Ovarian Cell Test Result: negative Chromosomal aberration assay Result: negative GreenScreen Assay Result: negative HPRT gene mutation in human lymphocytes Result: negative High throughput fluctuation test (HTFT) Result: negative In vitro cytogenetic Assay Result: negative L5178Y mouse lymphoma thymidine kinase locus assay Result: negative
	FLUTICASONE PROPIONATE	Micronucleus Assay Result: negative Species: Mouse Micronucleus Test Result: negative Species: Mouse
	SALMETEROL XINAFOATE	Rat Micronucleus Assay Result: negative
	FLUTICASONE PROPIONATE	SOS/umu Assay Result: negative Yeast Result: negative
Carcinogenicity	This product is not considered to be a carcinogen by IARC, ACGIH, NTP, or OSHA. Not classifiable as to carcinogenicity to humans.	
	SALMETEROL XINAFOATE	>= 0.15 mg/kg/day, Species-specific Result: positive Species: Rat Organ: Pituitary/ Uterus >= 1.4 mg/kg/day, Species-specific Result: positive Species: Mouse Organ: uterus
	FLUTICASONE PROPIONATE	Inhalation Result: negative Species: Rat dermal Result: negative Species: Mouse oral Result: negative Species: Mouse

Reproductive toxicity

Components in this product have been shown to cause birth defects and reproductive disorders in laboratory animals.

Reproductive toxicity**Reproductivity**

SALMETEROL XINAFOATE

0.1 mg/kg/day Reproductive performance and development of two untreated generations, NOEL

Species: Rat

Notes: GR33343X

1 mg/kg/day Reproductive performance and development of two untreated generations

Species: Rat

Organ: Skeletal effects

Notes: GR33343X

FLUTICASONE PROPIONATE

100 mcg/kg/day Embryofetal Development

Result: reduced foetal bodyweight, minor skeletal variations

Species: Rat

100 mcg/kg/day Female fertility (Segment I)

Result: reduced foetal bodyweight, minor skeletal variations

Species: Rat

SALMETEROL XINAFOATE

2 mg/kg/day Reproductive performance and development of two untreated generations, NOAEL

Species: Rat

Notes: GR33343G

FLUTICASONE PROPIONATE

50 mcg/kg/day Pre- and Post-natal development

Result: maternal toxicity

Species: Rat

SALMETEROL XINAFOATE

>= 1 mg/kg/day Embryo-foetal development- Oral,

Species-specific

Species: Rabbit

Organ: Skeletal effects, open eye, cleft palate

Notes: GR33343G

FLUTICASONE PROPIONATE

>= 25.7 mcg/kg/day Embryofetal Development

Result: maternal toxicity, reduced foetal body weight; no malformations or other variations

Species: Rat

>= 45 mcg/kg/day Embryofetal Development

Result: cleft palate

Species: Mouse

>= 50 mcg/kg/day Embryofetal Development

Result: maternal toxicity; reduced foetal weight; foetal resorptions

Species: Rabbit

SAR / QSAR, Glucocorticoid

Specific target organ toxicity - single exposure

Heart. Based on available data, the classification criteria are not met.

Specific target organ toxicity - repeated exposure

Adrenal glands. Bone tissue. Immune system. May cause damage to organs through prolonged or repeated exposure.

Aspiration hazard

Not likely, due to the form of the product.

Mixture versus substance information

No information available.

Other information

Caution - Pharmaceutical agent.

SECTION 12: Ecological information**12.1. Toxicity**

The product contains a substance which may cause long-term adverse effects in the environment. No information is available about the potential of this product to produce adverse environmental effects.

Components	Species		Test results
FLUTICASONE PROPIONATE (CAS 80474-14-2)			
Acute	IC50	Activated sludge	> 1000 mg/l, 3 hours
Aquatic			
Acute			
Crustacea	EC50	Water flea (Daphnia magna)	> 0.55 mg/l, 48 hours, Static test
Terrestrial			
Acute			
Earthworm	EC50	Manure worm (Eisenia foetida)	> 1000 mg/kg, 28 days

Components		Species	Test results
SALMETEROL XINAFOATE (CAS 94749-08-3)			
Aquatic			
<i>Acute</i>			
Activated Sludge Respiration	IC50	Residential sludge	> 998 mg/l, 3 hours
Algae	EC50	Green algae (Scenedesmus subspicatus)	4 mg/l, 72 hours, Measured
	NOEC	Green algae (Scenedesmus subspicatus)	1.9 mg/l
Crustacea	EC50	Water flea (Daphnia pulex)	20 mg/l, 48 hours
	NOEC	Water flea (Daphnia pulex)	6.7 mg/l, 48 hours
Fish	EC50	Rainbow trout (Juvenile Oncorhyncus mykiss)	35 mg/l, 96 hours, Static renewal test
	NOEC	Rainbow trout (Juvenile Oncorhyncus mykiss)	7.5 mg/l
<i>Chronic</i>			
Crustacea	LOEC	Water flea (Ceriodaphnia dubia)	5 mg/l, 8 days, Static renewal test
	NOEC	Water flea (Ceriodaphnia dubia)	1.6 mg/l, 8 days
Terrestrial			
<i>Acute</i>			
Earthworm	EC50	Manure worm (Eisenia foetida)	334 mg/kg, 28 days
	NOEC	Manure worm (Eisenia foetida)	209 mg/kg, 28 days

* Estimates for product may be based on additional component data not shown.

12.2. Persistence and degradability

Persistence and degradability

Photolysis

UV/visible spectrum wavelength

SALMETEROL XINAFOATE 338 nm

Hydrolysis

Half-life (Hydrolysis-neutral)

FLUTICASONE PROPIONATE > 1 years Measured

SALMETEROL XINAFOATE > 1 years Measured

Biodegradability

Percent degradation (Aerobic biodegradation-inherent)

SALMETEROL XINAFOATE 50 %, 12.8 days Modified Zahn-Wellens, primary biodegradation, loss of parent.

Percent degradation (Aerobic biodegradation-ready)

FLUTICASONE PROPIONATE < 44 %, 28 days

Percent degradation (Aerobic biodegradation-soil)

FLUTICASONE PROPIONATE 9 - 50 %, 64 days

SALMETEROL XINAFOATE 29.9 - 49.9 %, 64 days

12.3. Bioaccumulative potential

Partition coefficient

n-octanol/water (log Kow)

1,1,1,2-TETRAFLUOROETHANE 1.274

FLUTICASONE PROPIONATE 2.78

SALMETEROL XINAFOATE 2.1 (Measured).

12.4. Mobility in soil

Adsorption

Sludge/biomass distribution coefficient - log Kd

FLUTICASONE PROPIONATE 3.13 - 3.55 Estimated

Soil/sediment sorption - log Koc

FLUTICASONE PROPIONATE 3.41 - 3.83 Measured

SALMETEROL XINAFOATE 3.84 - 4.52

Mobility in general

Distribution

Octanol/water distribution coefficient log DOW

SALMETEROL XINAFOATE

1.32, pH 9

1.71, pH 7

2.06, pH 5

12.5. Results of PBT and vPvB assessment Not available.

12.6. Other adverse effects Not available.

SECTION 13: Disposal considerations

13.1. Waste treatment methods

Residual waste	Dispose of in accordance with local regulations. Empty containers or liners may retain some product residues. This material and its container must be disposed of in a safe manner (see: Disposal instructions).
Contaminated packaging	Empty containers should be taken to an approved waste handling site for recycling or disposal. Since emptied containers may retain product residue, follow label warnings even after container is emptied.
EU waste code	The Waste code should be assigned in discussion between the user, the producer and the waste disposal company.
Disposal methods/information	Collect and reclaim or dispose in sealed containers at licensed waste disposal site. This material and its container must be disposed of as hazardous waste. Do not allow this material to drain into sewers/water supplies. Do not contaminate ponds, waterways or ditches with chemical or used container. Dispose of contents/container in accordance with local/regional/national/international regulations.
Special precautions	Dispose in accordance with all applicable regulations.

SECTION 14: Transport information

ADR

14.1. UN number	UN1950
14.2. UN proper shipping name	AEROSOLS
14.3. Transport hazard class(es)	2.2
Subsidiary class(es)	-
14.4. Packing group	Not available.
14.5. Environmental hazards	No
Tunnel code	E
Labels required	2.2
Additional information:	
LTD QTY index	LQ2
Special Provisions	190, 327, 601, 625

IATA

14.1. UN number	UN1950
14.2. UN proper shipping name	Aerosols, non-flammable
14.3. Transport hazard class(es)	2.2
Subsidiary class(es)	-
14.4. Packing group	Not available.
Labels required	2
Additional Information:	
Passenger & cargo	Allowed.
Packaging Instruction	203
Pkg Inst cargo only	203
Pkg Inst pasenger & cargo	Y203
LQ	
SP See 44	A98,A145,A167
Max net qty pkg	75 kg
Max net qty pkg cargo only	150 kg
Max net qty pkg LQ	30 kg G

IMDG

14.1. UN number	UN1950
14.2. UN proper shipping name	AEROSOLS

14.3. Transport hazard class(es)	2
Subsidiary class(es)	5A
14.4. Packing group	Not available.
14.5. Environmental hazards	
Marine pollutant	No
Labels required	2
14.6. Special precautions for user	Not available.

14.7. Transport in bulk according to Annex II of MARPOL73/78 and the IBC Code MARPOL Annex II applies to liquids used in a ship's operation that pose a threat to the marine environment. These materials may not be transported in bulk.

ADR; IATA



SECTION 15: Regulatory information

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

EU regulations

Regulation (EC) No. 1005/2009 on substances that deplete the ozone layer, Annex I

Not listed.

Regulation (EC) No. 1005/2009 on substances that deplete the ozone layer, Annex II

Not listed.

Regulation (EC) No. 850/2004 On persistent organic pollutants, Annex I as amended

Not listed.

Regulation (EC) No. 689/2008 concerning the export and import of dangerous chemicals, Annex I, part 1 as amended

Not listed.

Regulation (EC) No. 689/2008 concerning the export and import of dangerous chemicals, Annex I, part 2 as amended

Not listed.

Regulation (EC) No. 689/2008 concerning the export and import of dangerous chemicals, Annex I, part 3 as amended

Not listed.

Regulation (EC) No. 689/2008 concerning the export and import of dangerous chemicals, Annex V as amended

Not listed.

Regulation (EC) No. 166/2006 Annex II Pollutant Release and Transfer Registry

Not listed.

Regulation (EC) No. 1907/2006, REACH Article 59(1) Candidate List as currently published by ECHA

Not listed.

Authorisations

Regulation (EC) No. 1907/2006, REACH Annex XIV Substances subject to authorization, as amended

Not listed.

Restrictions on use

Regulation (EC) No. 1907/2006, REACH Annex XVII Substances subject to restriction on marketing and use as amended

Not listed.

Directive 2004/37/EC: on the protection of workers from the risks related to exposure to carcinogens and mutagens at work

Not listed.

Directive 92/85/EEC: on the safety and health of pregnant workers and workers who have recently given birth or are breastfeeding

Not listed.

Other EU regulations

Directive 96/82/EC (Seveso II) on the control of major-accident hazards involving dangerous substances

Not listed.

Not listed.

Not listed.

Other regulations

The product is classified and labelled in accordance with EC directives or respective national laws. This Safety Data Sheet complies with the requirements of Regulation (EC) No 1907/2006.

National regulations

Follow national regulation for work with chemical agents.

15.2. Chemical safety assessment

No Chemical Safety Assessment has been carried out.

SECTION 16: Other information

List of abbreviations

Not available.

References

GSK Hazard Determination

Information on evaluation method leading to the classification of mixture

The classification for health and environmental hazards is derived by a combination of calculation methods and test data, if available.

Full text of any statements or R-phrases and H-statements under Sections 2 to 15

R36/38 Irritating to eyes and skin.
R48/20/21 Harmful: danger of serious damage to health by prolonged exposure through inhalation and in contact with skin.
R51/53 Toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment.
R52/53 Harmful to aquatic organisms, may cause long-term adverse effects in the aquatic environment.
R53 May cause long term adverse effects in the aquatic environment.
R61 May cause harm to the unborn child.
R62 Possible risk of impaired fertility.
H315 Causes skin irritation.
H319 Causes serious eye irritation.
H360D May damage the unborn child.
H361f Suspected of damaging fertility.

Revision information

Product and Company Identification: Business Units
Composition / Information on Ingredients: Ingredients
Physical & Chemical Properties:
Ecological Information: Mobility
Transport Information: Product Shipping Name/Packing Group
Regulatory Information: United States
GHS: Classification

Training information

Follow training instructions when handling this material.

Disclaimer

The information and recommendations in this safety data sheet are, to the best of our knowledge, accurate as of the date of issue. Nothing herein shall be deemed to create any warranty, express or implied. It is the responsibility of the user to determine the applicability of this information and the suitability of the material or product for any particular purpose.