

Products

Safety Data Sheet

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REVISION (see box 16)

Issue: 09 18:10:2006

1 IDENT	1 IDENTIFICATION OF THE SUBSTANCE/PREPARATION AND COMPANY		
Product Name	DEADLINE LIQUID CONCENTRATE		
Description	A blue rodenticidal, liquid-concentrate with no perceptible odour and a bittering additive. For use by professional operators for the control of rats and mice, after appropriate dilution with either water or other suitable bait bases.		
Company	Rentokil Initial Supplies, Liverpool, L33 7SR. Product advice line: 0151 548 5050 Emergency line: 0800 731 6 731		

2 HAZARD IDENTIFICATION

Classification (Supply - Use): In compliance with EC Directive 1999/45.

X_n HARMFUL R20/21/22 Harmful by inhalation, in contact with skin and if swallowed.

Adverse Physical, Chemical, Significant Human Health and Environmental Effects (See also box 11):

This product contains an anticoagulant compound. If ingested symptoms may include nosebleed and bleeding gums. In severe cases there may be bruising, haematomas of the joints and blood present in the faeces and urine.

No other significant adverse effects expected under normal conditions of handling and use.

% w/w	Common*/Chemical Name, Elincs/Einecs	& CAS No. of Ingredients	EC 1999/45 Classification
0.1	Bromadiolone* / 3-[3-(4-bromobiphenyl-4-yl)-3-hydhydroxycoumarin	Iroxy-1-phenylpropyl]-4- EINECS : 249-205-9 CAS : 28772-56-7	T+ : R26/27/28 N : R50,53
>50 ≤10.0	Monopropylene glycol* / propane-1,2-diol	EINECS : 200-338-0 CAS : 57-55-6	Not classified.
≤2.5	Bitrex [®] * / denatonium benzoate	EINECS : 223-095-2 CAS : 3734-33-6	Xn: R20/22 : R38 : R41 : R52,53

FIRST-AID MEASURES (SEE ALSO "ADVERSE EFFECTS" IN BOX 2)

Inhalation Remove patient to fresh air, keep warm and at rest. Apply supportive measures if necessary and seek medical attention.

Eye Contact Rinse affected eye with clean running water, or eyewash solution, for at least 15 minutes holding eyelids

well apart. Rinsé entire surface and do not allow run-off to contaminate unaffected eye. Seek medical attention.

Skin Contact Remove and wash contaminated clothing immediately. Wash affected area thoroughly with soap and

water. If the patient feels unwell seek medical advice.

Ingestion (Swallowing) Emergency Equipment Suggested

Do NOT induce vomiting. If unconscious place in the recovery position and apply supportive measures if necessary. If conscious give patient up to ½ litre or 1 pint of water to drink. Seek medical attention. Appropriate first-aid equipment should be provided. For the UK this should be in accordance with the Health & Safety (First-Aid) Regulations 1981. See also the Approved Code of Practice "First-aid at

Work". **Note To Doctor**

Further information on all Rentokil Initial formulations is lodged with the National Poisons Information Service in the UK. Bromadiolone is an indirect anticoagulant. Phytomenadione, Vitamin K1, is antidotal. Determine prothrombin times not less than eighteen hours after consumption. If elevated, administer Vitamin K1 until prothrombin time normalises. Continue determination of prothrombin time for two weeks after withdrawal of antidote and resume treatment if elevation occurs in that time.

5 FIRE FIGHTING MEASURES

Fire Extinguisher Type

Use carbon dioxide, foam, water, or dry powder extinguishers.

Special Fire-Fighting

Wear suitable personal protective equipment.

Procedures Special Exposure Hazards

Combustion or thermal decomposition may evolve toxic and irritant vapours.

ACCIDENTAL RELEASE MEASURES

Personal Precautions (See also box 8)

Wear suitable personal protective equipment.

Environmental Precautions

Keep away from drains, surface and ground water, and soil.

Clean-up Procedure (See also box 13)

Absorb spill with an inert material such as sand, earth or sawdust. Transfer to a suitable container for subsequent disposal. DO NOT contaminate watercourses or ground.

HANDLING AND STORAGE (SEE ALSO BOX 8)

Handling Avoid all contact by mouth.

Wash hands and exposed skin before meals and after use.

Storage Store in original container in a cool, dry, ventilated place out of the reach of children and away from food,

drink and animal feeding stuffs.

EXPOSURE CONTROLS/PERSONAL PROTECTION

Exposure Standard -Directive EC/98/24 (1st **IOELV Directive)**

Workplace Exposure Limit (WEL) for monopropylene glycol is 474 mg/m³ long-term exposure (8 hour Time Weighted Average) and for particulates is 10 mg/m³ long-term exposure (8 hour Time Weighted

Average).

Engineering Controls

Monopropylene glycol is referred to as propane-1,2-diol in Directive EC/98/24 (1St IOELV Directive). Where exposure may occur, engineering controls, rather than the provision of Personal Protective Equipment (PPE) should be employed. On completion of a risk assessment, the following PPE may be

Eye Protection Hand Protection None necessary during normal handling and use. Suitable hand protection such as gloves.

Skin Protection Breathing Protection

Suitable skin protection such as coveralls.

Environmental Exposure Controls None necessary during normal handling and use. Use only in accordance with instructions given.

PHYSICAL AND CHEMICAL PROPERTIES

Appearance and Odour A blue rodenticidal, liquid-concentrate with no perceptible odour and a bittering additive.

Solubility in Water Not determined. Miscible. Solubility in Other Solvents **Specific Gravity** 1.1 Miscible.

Flash Point Flammability Boiling Point/Range Vapour Density Vapour Pressure

Viscosity

Not applicable. Non-flammable. Starts at ca. 150°C Not applicable. Not applicable. Not determined. Other Data

Explosive Properties Combustibility **Oxidising Properties Evaporation Rate Partition Coefficient**

Not determined. Combustible. Not determined. Not determined. Not applicable. None known.

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10 STABILITY AND REACTIVITY

Conditions to avoid Materials to avoid Hazardous Breakdown Avoid extremes of temperature, e.g. below 0°C and above 40°C.

None known.

Combustion or thermal decomposition may evolve toxic and irritant vapours.

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11 **TOXICOLOGICAL INFORMATION (SEE ALSO BOX 2)**

Acute Toxicity Oral

LD₅₀ (rat): Males: 979 mg/kg

Females: 65 mg/kg

Inhalation Dermal

Harmful by inhalation.

LD₅₀ (rat): Males: 1421 mg/kg Females: >2000 mg/kg

Corrosivity/ Irritation

Skin Eyes No skin irritation expected. No eye irritation expected.

Respiratory tract

No respiratory tract irritation expected.

Sensitisation

Skin

Not expected to be a sensitiser.

Respiratory

Contains no known respiratory sensitisers.

Repeat-Dose Toxicity

Product does not contain any components known to have any effects relating to repeated-dose

Mutagenicity Carcinogenicity Reproductive Toxicity

Product does not contain any components known to have a mutagenic effect. Product does not contain any components known to have a carcinogenic effect. Product does not contain any components known to have effects on fertility.

Product does not contain any components known to be toxic to the reproductive system.

Other Information None known.

12 **ECOLOGICAL INFORMATION**

General Information

The bromadiolone in this product is classified as very toxic to aquatic organisms and may cause long-term adverse effects in the aquatic environment. The Bitrex in this product is classified as harmful to aquatic organisms and may cause long-term adverse effects in the aquatic environment. However, when used in accordance with instructions given, controlled release of this product is not expected to cause environmental contamination.

Ecotoxicity Data

LC₅₀ (96h) (Bluegill sunfish): 3.0 mg/L LC₅₀ (96h) (Rainbow trout): 1.4 mg/L

EC₅₀ (48h) (*Daphnia*): 2.0 mg/L

E_bC₅₀ (73h) (Algae: Scenédesmus subspicatus): 0.17 mg/L

For Bitrex

 LC_{50} (96h) (Rainbow trout): >1000 mg/L EC_{50} (48h) (*Daphnia magna*): 13 mg/L LC_{50} (96h) (Shrimp): 400 mg/L

Mobility

For Bitrex[®]: Water solubility: 45 g/L.

Persistence and Degradability

For bromadiolone: Bromadiolone is not considered volatile and is not expected to volatilise to air in significant

quantities, For Bitrex

Abiotic degradation: 10% after 5 days at 50°C at all pHs. Abiotic degradation: 10% after 30 days at 25°C at all pHs.

Bioaccumulative Potential

For bromadiolone:

Log Pow is greater than 3, which indicates a potential to bioaccumulate.

Other Adverse Effects

For Bitrex

If this substance is discharged at low concentrations into an adapted biological effluent treatment plant, the degrading action of the activated sludge will not be affected

13 **DISPOSAL CONSIDERATIONS**

Disposal of Waste / Containers

Ensure this product is disposed of as hazardous waste, in accordance with the appropriate

regulations.

Classification

Hazard Code: H5 - Harmful

(Council Directive 91/689/EC. Commission Decision 2000/532/EC (amended) Commission Decision

Components making the waste hazardous **Bromadiolone**

Concentrations (%): 0.1

2001/118/EC))

Note for Disposal

For further advice about disposal, in the UK, contact the local office of the Environment Agency

(England and Wales) or Scottish Environment Protection Agency. Local rate from anywhere in the UK: +44 (0) 870 850 6506.

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14 TRANSPOR	T INFORMATION	(INTERNATION	AL UNLESS OTHERWISE IN	IDICATED)
UN No.	Not classified.	Tremcard Reference No.	Not required.	RIS Code
Transport Category	Not required.	UK Hazchem EAC	Not required.	PSD56
ADR 2005	Class Not required.	ADR HIN	Not required.	Labels
(International Road)				
Proper Shipping Name	Not required.			
Limited Quantity Exemptions	Not required.			Not required.
Special Requirements	Not required.		Packing Group Not required.	Not required.
IMDG 2004 (Sea)	Class Not required.	IMDG EMS	Not required.	
Proper Shipping Name	Not required.			
Limited Quantity Exemptions	Not required.			
Special Requirements	Not required.		Packing Group Not required.	
Note for Transport	Local, State or National	requirements may apply	y to the carriage of this product.	

15 REGULAT	ORY INFORMATION (HEALTH AND SAFETY INFORMATION (SEE ALSO BOX 2))		
Safety Phrases	S35 This material and its container must be disposed of in a safe way. S36/37 Wear suitable protective clothing and gloves.		
Additional Label Phrases	To avoid risks to man and the environment, comply with the instructions for use.		
Legislation	Labelling is in accordance with UK regulations implementing the EC Directive 1999/45. Additional labelling requirements may be necessary in accordance with other National legislation. Outside the UK, the registration of this product may be necessary before use and any additional local requirements must be observed at all times.		
	The information given on this Safety Data Sheet (SDS) does not constitute an assessment in accordance with Control of Substances Hazardous to Health (COSHH) Regulations 2002, in the UK. Other National measures or guidance should be followed where appropriate.		

Bitrex® is a registered trade	mark of Macfarlan Smith Ltd.
Packaging Information	150ml high density polyethylene bottle with a screw top lid supplied in packs of 12.
Revisions	Changes have been made to the content of boxes 1, 2, 3, 4, 5, 8, 9, 11, 12, 13, 14, 15 & 16 (as indicated by the thick lines on the left-hand side of the boxes) compared with issue 08.
Risk phrase text (From box 3 - These refer to the ingredients only. See box 2 for the product risk phrases)	R20/22 : Harmful in contact with skin and if swallowed. R26/27/28 : Very toxic by inhalation, in contact with skin and if swallowed. R38 : Irritating to skin. R41 : Risk of serious damage to eyes. R50,53 : Very 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.

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Before using any product, ensure that you read and understand its label.

The information contained in this safety data sheet is, to the best of our knowledge and belief, accurate and reliable at the time of publication. The information relates only to the specific material designated in this safety data sheet and may not be valid for such material if it is used in combination with any other material(s) or any other use than that specified herein. Rentokil Initial UK Ltd is not liable for the use of this product for any other purpose than that described in this safety data sheet. This does not affect your statutory rights. It is the user's responsibility to satisfy him/herself as to the suitability in completeness of such information for his/her own particular use.

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