Labelling according to The Chemicals (Health and Safety) and Genetically Modified Organisms (Contained Use) (Amendment etc.) (EU Exit) Regulations 2019

Control of Substances Hazardous to Health Regulations 2002 (as amended). - The REACH etc. (Amendment etc.) (EU Exit) Regulations 2020.

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

1.1 Product identifier

Sakarat D Wax Bait : UK-2012-0370

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

Relevant identified uses: A ready-to-use liquid bait containing Difenacoum (0.005%w/w) For use against Rats and Mice Indoors and Outdoors around buildings: Rattus norvegicus (brown rat) and Mus musculus (house mice).

Uses advised against: Not to be used for Pulsed baiting or applied directly into burrows.

1.3. Details of the supplier of the safety data sheet

Address: Killgerm Chemicals Ltd, Wakefield Road, Ossett, WF5 9AJ

Tel: +44 (0)1924 268 450

Fax: +44 (0)1924 265 033 Email: technical@killgerm.com

1.4. Emergency telephone number

UK: Medical professionals should use National Poisons Information Service Tel: 0870 600 6266.

Killgerm Chemicals Ltd Tel:01924 268452 (Office hours). Emergency Number: 08701 906777

Non-medical professionals should seek information by contacting NHS by dialling 111.

IRELAND: Medical professionals should contact the national Poisons centre, Beaumont Hospital, Dublin (01-8092166)

SECTION 2: Hazards identification

2.1. Classification of the mixture according to Regulation (EC) No. 1272/2008 [CLP]

Repr. 1B; H360D May cause harm to unborn child.

STOT RE 2; H373 May cause damage to organs (blood) through prolonged or repeated exposure

2.2. Label elements



Signal Word: DANGER

Hazard statements:

H360D: May cause damage to unborn child.

H373: May cause damage to organs (blood) through prolonged or repeated exposure.

Precautionary statements:

- **P201:** Obtain special instructions before use.
- **P202:** Do not handle until all safety precautions have been read and understood.
- **P260:** Do not breathe dust/fumes/gas/mist/vapours/spray.
- **P280:** Wear protective gloves/protective clothing/eye protection/face protection.
- **P308 + P313:** IF exposed or concerned: Get medical advice/attention.
 - **P314:** Get medical advice/attention if you feel unwell.
 - P405: Store locked up.
 - P501: Dispose of contents/container in accordance with local regulation.

2.3. Other hazards

To avoid risks to human health and the environment, comply with instructions for use. Use bait containers clearly marked "poison" at all surface baiting points. Remove all remains of bait, dead rodents during and after treatment and dispose of safely. Prevent access to bait by children, domesticated animals and pets, (particularly cats, dogs, and pigs). Harmful to wildlife.

Exposure of non-target animals should be prevented.

Labelling according to The Chemicals (Health and Safety) and Genetically Modified Organisms (Contained Use) (Amendment etc.) (EU Exit) Regulations 2019 Killger

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SECTION 3: Composition/information on ingredients

3.2 Mixtures

Hazardous Components in Product

Ingredient Name	Classification	Concentration	H Phrases
Difenacoum (ISO)	Acute Tox. 1 (Oral)	0.005% w/w	H300
CAS Number:	Acute Tox. 1 (Dermal)		H310
56073-07-5	Acute Tox. 1(Inhalation)		H330
	Repr. 1B		H360D
	STOT RE 1		H372
	Aquatic Acute 1		H400
	Aquatic Chronic 1		H410
	Acute Tox. 4	0.001% w/w	H302, H332
CAS Number:	Skin Irrit. 2		H315
3734-33-6	Eye Dam. 1		H318
	Aquatic Chronic 3		H412

See section 16 for full text of R-phrases, H phrases and hazard classification of ingredients.

SECTION 4: First aid measures

4.1. Description of first aid measures

General: If during or after use/exposure you begin to feel unwell, seek medical attention bringing a copy of the product label/the SDS.

Eye contact: Flush with clean water for at least 15 minutes. Remove contact lenses, if present and easy to do. Continue rinsing. If eye irritation persists, seek medical advice.

Skin contact: Remove all contaminated clothing. Wash with water and then with soap and water.

Ingestion: DO NOT induce vomiting. Rinse mouth carefully with water. Never give anything by mouth to an unconscious person. If swallowed, seek medical advice immediately and show the product's container or label.

Inhalation: Remove from exposure. Get medical attention if any symptoms persist.

Contact a veterinary surgeon in case of ingestion by a pet.

DO NOT LEAVE POISONED PERSON ALONE UNDER ANY CIRCUMSTANCE.

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

This product contains Difenacoum, an indirect anticoagulant. Any signs of poisoning are unlikely to occur until 12-18 hours after ingestion. Thereafter, they will develop progressively and may rapidly appear. If ingested, symptom, which may be delayed, may include nosebleed and bleeding gums. In severe cases, there may be bruising, and blood present in the faeces or urine.

Clinical signs result from an increased bleeding tendency and include: an increase in prothrombin time, bruising easily with occasional gum bleeding, blood in the stool or urine, excessive bleeding from minor cuts and abrasions, pale mouth and cold gums, and general weakness. More severe cases of poisoning include haemorrhage (usually internal) and shock. and consequent

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

ADVICE FOR DOCTORS: Difenacoum is an indirect anti-coagulant. Phytomenadione and Vitamin K1 are antidotal. Determine prothrombin time not less than 18 hours after consumption. If elevated, administer Vitamin K1 40mg/day for adults and 20mg/day for children in divided doses. Continue until prothrombin times normalise. Continue determination of prothrombin time for two weeks after withdrawal of antidote and resume treatment if elevation occurs in that time.

N.B. Vitamin K3 is not effective.

For comprehensive medical advice on the treatment of poisoning, contact the nearest Poisons Information centre.

Labelling according to The Chemicals (Health and Safety) and Genetically Modified Organisms (Contained Use) (Amendment etc.) (EU Exit) Regulations 2019



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SECTION 5: Firefighting measures

5.1. Extinguishing media

Use water spray, foam, dry chemical or carbon dioxide. Cool the smouldering material with water spray to minimise the possibility of re-ignition. Keep containers and surroundings cool with water spray.

5.2. Special hazards arising from the substance or mixture

This product is non-flammable, but combustible. May produce toxic fumes of carbon monoxide if involved in a fire.

5.3. Advice for firefighters

Wear self-contained breathing apparatus.

SECTION 6: Accidental release measures

6.1 Personal precautions, protective equipment and emergency procedures

Use personal protective equipment (Section 8). Refer to protective measures listed in sections 7 and 8.

6.2. Environmental precautions

Ensure the product does not reach any water channel or drains. If this occurs, notify the relevant authorities immediately.

6.3. Methods and material for containment and cleaning up

Scrape up material. Place in marked receptacle ready for disposal. Contact supplier for advice on disposal. See section 13

6.4. Reference to other sections

See also sections 8 and 13

SECTION 7: Handling and storage

7.1. Precautions for safe handling

The product must be used and stored only in accordance with the product label. Refer also to the section "Exposure Controls/Personal Protection". Avoid all contact by mouth. Wash hands and exposed skin before meals and after use. Empty container completely and dispose of safely.

7.2. Conditions for safe storage, including any incompatibilities

Store in the original container under cool and dry conditions in a secure, well ventilated place, inaccessible to children, and away from foodstuffs and animal feed. Store and transport away from products which may have an odour

7.3. Specific end use(s)

Before using this product read the instructions for use. The product is intended for use as a rodenticide.

SECTION 8: Exposure controls/personal protection

8.1. Control parameters

No specific national limit values have been established.

8.2. Exposure controls

Where exposure may occur engineering controls should be employed. A risk assessment should be carried out and the following PPE may be appropriate /required.

PPE	Item In Use	Spillage	
Respirators		Use half-mask (EN140) with a particle filter P (EN 143) to required (nominal) protection factor (Minimum).	
Gloves	Protective Gloves (EN 374). Nitrile or PVC.	Protective Gloves (EN 374). Nitrile or PVC.	
	Basic type e.g. Heavy duty polycotton or coverall type 5/6.	coverall type 5/6.	
Goggles / Face shield		Safety glasses to EN 166 3459B.	

Labelling according to The Chemicals (Health and Safety) and Genetically Modified Organisms (Contained Use) (Amendment etc.) (EU Exit) Regulations 2019



Control of Substances Hazardous to Health Regulations 2002 (as amended) The REACH etc. (Amendment etc.) (EU Exit) Regulation			
SECTION 9: Physical and chemical properties			
9.1. Information on basic physical and chemical properties			
Appearance:	Blue liquid		
Odour:	odourless.		
Odour threshold:	No available data.		
pH:	7		
Melting point/freezing point:	Not applicable.		
Initial boiling point/boiling range:	Not applicable.		
Flash point:	No available data		
Evaporation rate:	No available data		
Flammability:	not applicable.		
Upper/lower flammability	No available data.		
Vapour pressure:	Not applicable.		
Vapour density:	Not applicable.		
Relative Density:	1.06		
Solubility in water:	No available data		
Solubility in other solvents:	No available data.		
Partition coefficient:	No available data.		
Auto-ignition temp:	371°C		

No available data. No available data.

Product is not explosive

No oxidising properties.

SECTION 10: Stability and reactivity

10.1. Reactivity

Decomposition temp:

Explosive properties:

Oxidising properties:

Viscosity:

Stable under recommended transport or storage conditions.

10.2. Chemical stability

Product is stable under normal conditions according to handling and storage.

10.3. Possibility of hazardous reactions

None anticipated.

10.4. Conditions to avoid

Avoid extremes of temperature.

10.5. Incompatible materials

Store away from strong oxidising agents.

10.6. Hazardous decomposition products

Carbon monoxide and oxides of nitrogen, Toxic and irritant fumes released if mixture is involved in a fire.

Labelling according to The Chemicals (Health and Safety) and Genetically Modified Organisms (Contained Use) (Amendment etc.) (EU Exit) Regulations 2019



SECTION 11: Toxicological information

11.1. Information on toxicological effects

a) acute toxicity; Information has been derived from the properties of the individual ingredients. Oral LD50

(rat) >2000mg/kg. Inhalation- Not an anticipated route of exposure.

b) irritation; Skin eyes, respiratory tract – no irritation potential expected. Information derived from the properties of the individual ingredients

c) corrosivity; The product is not classified as corrosive

d) sensitisation; Contains no known skin or respiratory sensitizers.

e) repeated dose toxicity; The product has not been tested. Repeated exposure to small quantities may affect certain organs. Damages the coagulation system

f) carcinogenicity/mutagenicity; Product does not contain any ingredients known to have such effects.

g) toxicity for reproduction; may cause harm to the unborn child.

11.2. Other Data

See section 2.3

SECTION 12: Ecological information

12.1. Toxicity

Difenacoum (ISO) (56073-07-5) in this product is classified as very toxic to aquatic organisms and may cause long term adverse effects in the aquatic environment. However, when used in accordance with instructions, controlled release of this product is not expected to cause environmental contamination.

LC50 Fish (Oncorhynchus mykiss) 0.064 mg/L - (Directive 92/69/EEC, C.1)

LC50 Crustacea (48hr Daphnia magna) 0.52mg/L - (Directive 92/69/EEC, C.2)

ErC50 Algae No observed effect concentration (72 h) 0.25 mg/l (growth rate), Pseudokirchneriella subcapitata

12.2. Persistence and degradability

Degradation in soil is slow. Difenacoum, soil half-life of 439 days.

12.3. Bioaccumulative potential

Difenacoum: Because of the n-Octanol/Water distribution coefficient (log Pow) accumulation in organisms is possible.

12.4. Mobility in soil

Difenacoum mobility is very low and will depend principally on the soil type. Difenacoum and any potential degradation products, even if released indirectly into the soil in small quantities are not likely to move through the soil profile and are unlikely to reach groundwater in significant quantities.

Assessment transport between environmental compartments:

Following exposure to soil, adsorption to solid soil particles is probable, therefore contamination of ground water is not expected.

12.5. Results of PBT and vPvB assessment

Does not meet requirement for assessment.

12.6. Other adverse effects

None known.

SECTION 13: Disposal considerations

13.1. Waste treatment methods

Disposal of unconsumed product, empty containers and contaminated packaging must be made in accordance with the local law. For information on disposal in the UK contact the environment agency (www.environment-agency.gov.uk) or SEPA (www.SEPA.org.uk). Dispose of unused product in the original container as hazardous waste.

Empty containers and contaminated PPE should be considered hazardous and disposed of appropriately.

Suggested European waste code 20 01 19.

Labelling according to The Chemicals (Health and Safety) and Genetically Modified

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SECTION 14: Transport information

14.1. UN number

Not applicable.

14.2. UN proper shipping name

Not applicable.

14.3. Transport hazard class(es).

Not applicable.

14.4. Environmental hazards

No

14.5. Special precautions for user

Not applicable.

14.6. Transport in bulk according to Annex II of Marpol and the IBC Code.

Not applicable.

SECTION 15: Regulatory information

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

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The REACH etc. (Amendment etc.) (EU Exit) Regulations 2020.

• Refer to other relevant measures such as the Health and Safety at Work etc Act 1974 and the

COSHH regulations and guidance.

• The information contained in this data sheet does not constitute the user's own assessment of workplace risks as required by legislation.

15.2. Chemical safety assessment

Advice on product handling can be found in sections 7 and 8.

SECTION 16: Other information

Use only in accordance with label instructions. Operatives using this product should be trained in its use.

The information in this data sheet should be considered when undertaking a risk assessment under the COSHH regulations.

Ingredient classification data:	
Acute Toxicity Category 1 (Oral)	H300 Fatal if swallowed.
Acute Toxicity Category 3 (Oral)	H302 Harmful if swallowed.
Acute Toxicity Category 1 (Dermal)	H310 Fatal in contact with skin.
Skin irritant category 2	H315 Causes skin irritation.
Eye damage category 1	H318 Causes serious eye damage.
Acute toxicity Category 1 (Inhalation)	H330 Fatal if inhaled.
Acute toxicity Category 3 Inhalation)	H332 Harmful if inhaled.
Reproductive Toxicity Category 1	H360D May damage the unborn child.
Single target organ toxicity – Repeat exposure Category 1	H372 Causes damage to organs through prolonged or repeated exposure. (Blood)
Single target organ toxicity – Repeat exposure Category 2	H372 May Cause damage to organs through prolonged or repeated exposure. (Blood)
Aquatic Toxicity Acute Category 1	H400 Very toxic to aquatic life.
Aquatic Toxicity Chronic Category 1	H410 Very toxic to aquatic life with long lasting effects.
Aquatic Toxicity Chronic Category 3	H412 Harmful to aquatic life with long lasting effects.

Labelling according to The Chemicals (Health and Safety) and Genetically Modified



Organisms (Contained Use) (Amendment etc.) (EU Exit) Regulations 2019 Control of Substances Hazardous to Health Regulations 2002 (as amended). - The REACH etc. (Amendment etc.) (EU Exit) Regulations 2020.

Addition of emergency number.

Issue number (date)	Section amended
Version (Dec 2017)	SDS was first produced
Version (Jan 2020)	Minor updates to text and layout.
Version (Sep 2021)	Updates to regulatory information.

This safety data sheet does not constitute a COSHH assessment.

Version (DEC 2021)

The information contained within this data sheet is strictly for general guidance only and should not be relied upon over and above this. This data sheet is intended to provide general health and safety guidance on the handling, storage, and transportation of the preparation. The information provided in this data sheet is accurate at the date of publication and will be updated as and when appropriate. No liability will be accepted by Killgerm Chemicals Limited for any loss, injury or damage arising from any failure to comply with the information and advice contained within this data sheet and/or failure to comply with the manufacturer's guidelines, product label data and any associated technical usage literature.