NIX DERMAL CREAM

# **SAFETY DATA SHEET**



# 1. IDENTIFICATION OF THE SUBSTANCE/PREPARATION AND OF THE COMPANY/UNDERTAKING

Material

**NIX DERMAL CREAM** 

**Synonyms** 

NIX DERMAL CREAM 5% (CANADA) \* NIX DERMAL CREME POUR LA PEAU 5% \* NIX CREAM \* LOTRIX CREAM \* LOXAZOL HYDROFIELE CREME \* LYCLEAR CREAM \* LYCLEAR DERMAL CREAM \* ZALVOR CREAM \*

PERMETHRIN, FORMULATED PRODUCT

Company Name

GlaxoSmithKline, Corporate Environment, Health & Safety

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TW8 9GS UK

UK General Information:

+44-20-8047-5000 +44-1865-407333

Transport Emergency (EU)

+1-612-221-3999. Ext 221

Medical Emergency
Information and Advice:

US number, available 24 hours

Multi-language response

GlaxoSmithKline, Corporate Environment, Health & Safety

One Franklin Plaza, 200 N 16th Street

Philadelphia, PA

19102-1225 US

US General Information:

+1-888-825-5249

Transport Emergency (non EU)

+1-703-527-3887

US number, available 24 hours Multi-language response

## 2. COMPOSITION / INFORMATION ON INGREDIENTS

Ingredients	CAS RN	Percentage
PERMETHRIN	52645-53-1	5
NON-HAZARDOUS INGREDIENTS	Unassigned	95

## 3. HAZARDS IDENTIFICATION

Fire and Explosion

Expected to be non-combustible.

Health

Exposure might occur via eyes; skin; ingestion.

Eye irritant.

Health effects information is based on hazards of components.

\* Environment

Harmful to aquatic organisms. May cause long-term adverse effects in the

aquatic environment.

## 4. FIRST-AID MEASURES

SDS Number 127995 Approved/Revised 09-Aug-2006

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Ingestion Never attempt to induce vomiting. Do not attempt to give any solid or liquid

by mouth if the exposed subject is unconscious or semi-conscious. Wash out the mouth with water. If the exposed subject is fully conscious, give

plenty of water to drink. Obtain medical attention.

Inhalation Physical form suggests that risk of inhalation exposure is negligible.

Skin Contact Using appropriate personal protective equipment, remove contaminated

clothing and flush exposed area with large amounts of water. Obtain medical attention if skin reaction occurs, which may be immediate or

delayed.

Eye Contact Wash immediately with clean and gently flowing water. Continue for at least

15 minutes. Obtain medical attention.

None for occupational exposure.

NOTES TO HEALTH PROFESSIONALS

Medical Treatment Treat according to locally accepted protocols. For additional guidance, refer

to the local poison control information centre.

Medical Conditions
Caused or Aggravated

by Exposure

Antidotes No specific antidotes are recommended.

5. FIRE-FIGHTING MEASURES

Fire and Explosion Hazards Not expected for the product, although the packaging is combustible.

Extinguishing Media

Water is recommended for fires involving packaging.

Special Firefighting Procedures

For single units (packages): No special requirements needed.
For larger amounts (multiple packages/pallets) of product: Since toxic,

corrosive or flammable vapours might be evolved from fires involving this product and associated packaging, self contained breathing apparatus and

full protective equipment are recommended for firefighters.

If possible, contain and collect firefighting water for later disposal.

Hazardous Combustion Products

Toxic, corrosive or flammable thermal decomposition products are expected

when the product is exposed to fire.

6. ACCIDENTAL RELEASE MEASURES

Personal Precautions Wear protective clothing and equipment consistent with the degree of

hazard.

Environmental Precautions Prevent entry into waterways, sewers, surface drainage systems and poorly

ventilated areas.

Clean-up Methods Collect and place it in a suitable, properly labelled container for recovery or

disposal.

Decontamination

Procedures

Detergent solutions can be used for clean-up and decontamination

operations.

7. HANDLING AND STORAGE

HANDLING

General Requirements No special control measures required for the normal handling of this

product. Normal room ventilation is expected to be adequate for routine

handling of this product.

STORAGE No storage requirements necessary for occupational hazards. Follow

product information storage instructions to maintain efficacy.

8. EXPOSURE CONTROLS/PERSONAL PROTECTION

PERSONAL PROTECTIVE EQUIPMENT

Version 4

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**Eye Protection** 

Wear approved safety glasses with side shields if eye contact is possible.

\* Other Equipment or **Procedures** 

Follow all local regulations if personal protective equipment (PPE) is used in the workplace. Wear appropriate clothing to avoid skin contact. Wash hands

and arms thoroughly after handling.

# 9. PHYSICAL AND CHEMICAL PROPERTIES

**Appearance** 

**Physical Form** 

Cream.

# 10. STABILITY AND REACTIVITY

**Stability** 

This product is expected to be stable.

**Conditions to Avoid** 

None for normal handling of this product.

## TOXICOLOGICAL INFORMATION

**Oral Toxicity** 

Not expected to be toxic following ingestion.

Inhalation Toxicity

No studies have been conducted.

Skin Effects

Irritation is not expected following direct contact.

**Eye Effects** 

Irritation might occur following direct contact with eyes.

Sensitisation

Sensitisation (allergic skin reaction) is not expected.

**Genetic Toxicity** 

Not expected to be genotoxic under occupational exposure conditions.

Carcinogenicity

No components are listed as carcinogens by GSK, IARC, NTP or US OSHA.

Reproductive Effects

Insufficient information available to classify for reproductive toxicity.

Other Adverse Effects

Overexposure in the workplace might have the following effects: abnormal

nervous system sensations.

# 12. ECOLOGICAL INFORMATION

Summary

This product contains an active ingredient that has been tested and which may be harmful if released directly to the environment. Consult the MSDS of the active ingredient for specific information about potential environmental effects. Appropriate precautions should be taken to limit release of this mixture to the environment. Local regulations and procedures should be consulted prior to environmental release.

**ECOTOXICITY** 

Aquatic

**Microtox** 

Microtox is a general toxicity test which utilizes a sensitive marine photo bacteria as the test species. This material contains an active pharmaceutical

ingredient that is toxic to these microorganisms.

Daphnid

This mixture contains an active pharmaceutical ingredient that is very toxic

to daphnids.

Fish

This mixture contains an active pharmaceutical ingredient that is very toxic

to fish.

**MOBILITY** 

**Partitioning** 

This mixture contains an active pharmaceutical ingredient with octanol/water partition coefficient data that suggests that for environmental fate predictions the active pharmaceutical ingredient may have the tendency to distribute into fats.

# PERSISTENCE/DEGRADATION

Biodegradation

This mixture contains an active pharmaceutical ingredient that has been tested and is expected to be biodegradable.

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\* BIOACCUMULATION

This material contains an active pharmaceutical ingredient that will have a tendency to bioaccumulate in the food chain.

## 13. DISPOSAL CONSIDERATIONS

Disposal Recommendations Collect for recycling or recovery if possible. The disposal method for rejected products/returned goods must ensure that they cannot be re-sold or

re-used.

**Regulatory Requirements** 

Observe all local and national regulations when disposing of this product.

#### 14. TRANSPORT INFORMATION

The SDS should accompany all shipments for reference in the event of spillage or accidental release. Only authorised persons trained and competent in accordance with appropriate national and international regulatory requirements may prepare dangerous goods for transport.

## **UN Classification and Labelling**

Transport Information

Transportation and shipping of this product is not restricted. It has no known, significant hazards requiring special packaging or labelling for air, maritime, US or European ground transport purposes.

#### 15. REGULATORY INFORMATION

The information included below is an overview of the major regulatory requirements. It should not be considered to be an exhaustive summary. Local regulations should be consulted for additional requirements.

#### EU Classification and Labelling

Exempt from requirements of EU Dangerous Preparations directive - product regulated as a medicinal product, cosmetic product or medical device.

#### US OSHA Standard (29 CFR Part 1910.1200)

Classification

This product is classified as hazardous according to the OSHA Hazard

Communication Standard.

Other US Regulations

**TSCA Status** 

Exempt

#### 16. OTHER INFORMATION

References

**GSK Hazard Determination** 

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### **SDS Sections Updated**

**Sections** 

Subsections

COMPOSITION / INFORMATION ON INGREDIENTS

**EXPOSURE CONTROLS / PERSONAL PROTECTION** 

HAZARDS IDENTIFICATION

Other Equipment or Procedures

Environment

IDENTIFICATION OF SUBSTANCE / PREPARATION AND OF

**COMPANY** 

REGULATORY INFORMATION

US Environmental (EPA) Requirements

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.