Material Data Safety Sheet

Product Name: Nyloxin™ Oral Spray 4x (140 mcg/mL)
Category: Analgesic

This is a pharmaceutical product manufactured in accordance with FDA requirements. The information contained on this MSDS is based primarily on a review of the active pharmaceutical ingredient(s).

I. Manufacturer Information

Manufacturer
Nutra Pharma Corporation
2776 University Drive
Coral Springs, FL 33065
www.NutraPharma.com

Emergency Number: (954) 509-0911
MSDS Prepared by: Nutra Pharma Corporation
MSDS Date: 07/21/10
MSDS Version: 01

II. Information on Primary Ingredients

Active Ingredient:
Asian Cobra Venom 140 mcg/mL

Inactive Ingredients:
Flavoring
Purified water
Sodium benzoate
Sodium phosphate
Xylitol

III. Physical Data

Appearance: Clear, Colorless liquid

Specific Gravity (H2O = 1): 1
Vapor Pressure (mm Hg 20°C): Not available.
Vapor Density (air = 1): Not available.
Molecular Weight: Not available.
pH: 3.0 – 6.0
Viscosity @ 25ºC: Not available.
% Volatile by Volume: 0

Melting Point: Not available.
Boiling Point: 100°C
Evaporation Rate: Not available.
Solubility in Water: Totally soluble.
Solubility in Alcohol: Not available.
Solubility in Oil: Not available.

IV. Fire and Explosion Hazard Data

Flash Point (and method): Not established**.**
Flammability: Not considered to be a fire hazard.
Conditions of flammability: Not established.
Flammable Limits: Not established.
Auto-ignition temperature: Not established.
Extinguishing Media: Not established.
Special Fire Fighting Procedures: Not established.
Sensitivity to static discharge & Explosive power: Not established.

V. Reactivity Data

Stability: Stable for 24 months at ambient temperatures (15-27 °C / 65-80 °F).
Incompatibility (material to avoid): Not established.
Conditions to Avoid: Avoid direct UV light. Do not freeze.
Hazardous decomposition products: Not established.
Hazardous Polymerization: Will not occur.

VI. Health Hazard Assessment

Routes of Entry: Oral medication. Handling this product in its final form presents minimal risk from occupational exposure.
Skin: Minimal risk from occupational exposure.
Eyes: May cause irritation.
Inhalation: Minimal risk from occupational exposure.
Ingestion: May induce headache, nausea, sore throat, allergic rhinitis, gastrointestinal discomfort, itchiness or mild rash.

VII. Emergency Procedures

Skin: Wash skin with mild soap and water. If irritation or other symptoms develop, use an over-the-counter antihistamine to control symptoms. If symptoms persist, seek medical attention.
Eyes: Flush with water. If irritation or other symptoms develop, use an over-the-counter antihistamine to control symptoms. If symptoms persist, seek medical attention.
Inhalation: If irritation or other symptoms develop, use an over-the-counter antihistamine to control symptoms. If symptoms persist, seek medical attention.
Ingestion If irritation or other symptoms develop, use an over-the-counter antihistamine to control symptoms. If symptoms persist, seek medical attention. Never give anything by mouth to an unconscious person. Do not induce vomiting unless directed by medical personnel.
Additional information: If a rare sensitivity reaction occurs, the drug should be stopped.

VIII. Toxicological Information

This product is not toxic in any amounts.

IX. Precaution Information

Steps to be taken in case material is released or spilled: Contain spill and soak up liquid with absorbent material.
Waste Disposal Method: Dispose of waste in respect with all regulatory agencies and laws.
Precautions to be taken in Handling and Storing: Store between 15 and 27 °C. Avoid direct UV light. Do not freeze.

X. Control Measures

Respiratory Protection (Specify Type): Not necessary.
Ventilation: Local Exhaust & Mechanical (General): Not necessary.
Protective Gloves: Not necessary.
Eye Protection: Not necessary.
Other Protective Clothing or Equipment: Not necessary.

XI. Regulatory Information

OSHA Regulations: Not OSHA regulated.
DOT Regulations: Not DOT regulated.
EPA Regulations: Not EPA regulated.
FDA Regulations: Consult CPG 7132.15, "Conditions under which homeopathic drugs may be marketed".

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