SAFETY DATA SHEET



IDENTIFICATION OF THE SUBSTANCE/PREPARATION AND OF THE COMPANY/UNDERTAKING

Material VASOXYL INJECTION

Synonyms VASOXYL INJECTION 20MG * VASOXINE INJECTION 20MG/ML *

METHOXAMINE HYDROCHLORIDE, FORMULATED PRODUCT

Company Name GlaxoSmithKline, Corporate Environment, Health & Safety

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Medical Emergency +1-612-221-3999, Ext 221
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Multi-language response

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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
METHOXAMINE HYDROCHLORIDE	61-16-5	2
NON-HAZARDOUS INGREDIENTS	Unassigned	98

3. HAZARDS IDENTIFICATION

Fire and Explosion This product is expected to be non-combustible.

Health Handling this product in its final form presents minimal risk from

occupational exposure. Health effects information is based on hazards of

components.

Possible effects of overexposure in the workplace include: increased blood

pressure; headache; vomiting.

Environment No information is available about the potential of this product to produce

adverse environmental effects.

4. FIRST-AID MEASURES

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.

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

15 minutes. Obtain medical attention.

NOTES TO HEALTH PROFESSIONALS

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

to the current prescribing information or to the local poison control

information centre. Medical treatment in cases of overexposure should be

treated as an overdose of alpha-adrenergic receptor blocker.

Medical Conditions Caused or Aggravated by Exposure

Refer to prescribing information for detailed description of medical conditions caused by or aggravated by overexposure to this product.

Health Surveillance Procedures

Pre-placement and periodic health surveillance is not usually indicated. The final determination of the need for health surveillance should be determined

by local risk assessment.

Antidotes No specific antidotes are recommended. For the latest information, refer to

the local poison control information centres.

FIRE-FIGHTING MEASURES

Fire and Explosion

Hazards

This product is non-combustible, 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.

ACCIDENTAL RELEASE MEASURES

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

hazard.

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

ventilated areas.

Spread an inert absorbent on the spill and place in a suitable, properly **Clean-up Methods**

labelled container for recovery or disposal.

Decontamination Procedures

Water 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.

EXPOSURE CONTROLS/PERSONAL PROTECTION

ENGINEERING CONTROLS

Exposure Controls An internal GSK Occupational Exposure Level (OEL) of 0.2 ppm (15 min

TWA) has been set for methoxamine hydrochloride, the active substance in

this product.

PERSONAL PROTECTIVE EQUIPMENT

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

Other Equipment or

None required for normal handling. Wash hands and arms thoroughly after

Procedures

handling.

9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance

Physical Form Aqueous solution.

Packaging Vial.

pH of Aqueous Solutions 3 to 5

10. STABILITY AND REACTIVITY

Stability DO NOT FREEZE - dispose of properly if frozen.

Conditions to AvoidNone for normal handling of this product.

11. TOXICOLOGICAL INFORMATION

Oral Toxicity

Not expected to be toxic following ingestion.

Skin Effects

Irritation might occur following direct contact.

Eye Effects Minor irritation might occur following direct contact with eyes.

Target Organ Effects Adverse effects might occur in the following organ(s) following

overexposure: cardiovascular system.

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 Not expected to produce adverse effects on fertility or development under

occupational exposure conditions.

Pharmacological Effects This material is a selective adrenergic alpha 1A agonist It is an agent

intended for the treatment of decrease in blood pressure.

Other Adverse Effects Overexposure in the workplace might have the following effects: increased

blood pressure; headache; vomiting.

12. ECOLOGICAL INFORMATION

SummaryNo information is available about the potential of this product to produce

adverse environmental effects. Local regulations and procedures should be

consulted prior to environmental release.

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

None.

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

Date Approved/Revised 23-Aug-2003

SDS Version Number 5

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.