SAFETY DATA SHEET

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

Material
ZEASORB AF POWDER

Synonym(s)
ZEASORB ANTI-FUNGAL POWDER * MICONAZOLE NITRATE TOPICAL POWDER *
ZEASORB-AF SUPER ABSORBANT ANTIFUNGAL POWDER (ATHLETE'S FOOT) *
ZEASORB-AF SUPER ABSORBANT ANTIFUNGAL POWDER (JOCK ITCH) *
ZEASORB-AF SUPER ABSORBANT ANTIFUNGAL POWDER (MICONAZOLE NITRATE 2%) *
FORMULATION CODE MF1048; 445A *
STIEFEL PRODUCT *
MICANZOLE NITRATE AND ALUMINUM DIHYDROXYALLANTOINATE, FORMULATED PRODUCT

Recommended Use
Cosmetic Product

Note
This safety data sheet is written to provide health, safety and environmental information for people handling this formulated product in the workplace. It is not intended to provide information relevant to medicinal use of the product. In this instance patients should consult prescribing information/package insert/product label or consult their pharmacist or physician. For health and safety information for individual ingredients used during manufacturing, refer to the appropriate safety data sheet for each ingredient.

Company Name
Stiefel Laboratories, Inc. (a GSK company)
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Brentford, Middlesex TW8 9GS UK
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5 Moore Drive
Research Triangle Park, NC 27709 USA
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EMERGENCY PHONE NUMBERS -

TRANSPORT EMERGENCIES (by country / geographic region):
Africa / EU / Israel / Middle East (English / European languages): +44 (0) 1235 239 670
Asia Pacific (except China): +65 3158 1074
China: +86 10 5100 3039
Middle East / Africa (Arabic-speaking countries): +44 (0) 1235 239 671
US: +1 703 527 3887
available 24 hrs/7 days; multi-language response

MEDICAL EMERGENCIES:
+1 612 221 3999, Ext 221
available 24 hrs/7 days; multi-language response

2. HAZARDS IDENTIFICATION

Fire and Explosion Hazards
Assume that this material is capable of producing a dust explosion if ignited as a dust cloud.
Assume that this material is capable of being ignited by an electrostatic discharge.
Assume that this material is capable of sustaining combustion.
**Health**  
May cause transient eye irritation. Health effects information is based on hazards of components. Exposure might occur via inhalation; ingestion; skin; eyes.

**Environment**  
No information is available about the potential of this product to produce adverse environmental effects.

### 3. COMPOSITION / INFORMATION ON INGREDIENTS

<table>
<thead>
<tr>
<th>Ingredients</th>
<th>CAS #</th>
<th>Percent</th>
<th>EC-No.</th>
</tr>
</thead>
<tbody>
<tr>
<td>ALUMINUM DIHYDROXYALLANTOINATE</td>
<td>5579-81-7</td>
<td>0.22</td>
<td>226-964-4</td>
</tr>
<tr>
<td>CHLOROXYLENOL</td>
<td>1321-23-9</td>
<td>0.6</td>
<td>215-316-6</td>
</tr>
<tr>
<td>MICONAZOLE NITRATE</td>
<td>22832-87-7</td>
<td>2.0</td>
<td>245-256-6</td>
</tr>
</tbody>
</table>

Other components below reportable levels  >97.0

### 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**  
Using appropriate personal protective equipment, move exposed subject to fresh air. If breathing is difficult or ceases, ensure and maintain ventilation. Give oxygen as appropriate. The exposed subject should be kept warm and at rest. Obtain medical attention in cases of known or possible over exposure, or with symptoms including chest pain, difficulty breathing, loss of consciousness or other adverse effects, which may be delayed.

**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.

### 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 Conditions Caused or Aggravated by Exposure**  
None for occupational exposure.

**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.

### 5. FIRE-FIGHTING MEASURES

**Fire and Explosion Hazards**  
The combustibility of the product is not known, however the packaging is combustible.

**Extinguishing Media**  
Water or foam extinguishers are recommended. Carbon dioxide or dry powder extinguishers may be ineffective.

**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.
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

OCCUPATIONAL EXPOSURE LIMITS

INGREDIENT  MICONAZOLE NITRATE
GSK Occupational Hazard Category 2
Other Equipment or Procedures
None required for normal handling. Wash hands and arms thoroughly after handling.

9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance
Physical Form Powder.

Dust Electrostatic Properties
Minimum Ignition Energy (Cloud) No studies have been conducted.

Dust Explosion Properties
St Class No studies have been conducted.
Train Fire No studies have been conducted.

10. STABILITY AND REACTIVITY

Stability This product is expected to be stable.
Conditions to Avoid None for normal handling of this product.

11. TOXICOLOGY INFORMATION

Pharmacological Effects
This product contains active ingredient(s) with the following activity: an antifungal agent.

Target Organ Effects
No specific target organ effects have been identified.

Routes of Exposure
Oral Toxicity Non-toxic following ingestion.
Inhalation Toxicity No studies have been conducted.
Skin Effects Irritation is not expected following direct contact.
Eye Effects Minor irritation might occur following direct contact with eyes. Assessment based on the physico-chemical properties of this material

Sensitisation Sensitisation (allergic skin reaction) is not expected.

Genetic Toxicity Not expected to be genotoxic, based on effects of individual components.

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.

Other Adverse Effects None known for occupational exposure.
12. ECOLOGICAL INFORMATION

Summary
No 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
Not regulated in transport.

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.

Classification
This product is exempt from the requirements of the OSHA Hazard Communication Standard.

Other US Regulations
TSCA Status
Exempt

16. OTHER INFORMATION

References
GSK Hazard Determination

SDS Version Number
2

SDS Sections Updated

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.