

SKIN SO SOFT BUG GUARD PLUSITCH RELIEF ITCH RELIEF - menthol spray
Avon Products, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Drug Facts

Active ingredient

Menthol 0.5 %.....

Purpose

.....External Analgesic

Uses

For the temporary relief of pain and/or itching associated with minor burns, sunburn, minor cuts, scrapes, insect bites or minor skin irritations.

Warnings

For external use only

When using this product

- Avoid contact with eyes. If contact occurs, rinse with water to remove

Stop use and ask a doctor if

- If condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away

Directions

Adults and children 2 years of age and older:

- Apply to affected area not more than 3 to 4 times daily

Children under 2 years of age:

- Consult a doctor before use

Inactive Ingredients:

Water/Eau, C12-15 Alkyl Benzoate, Ethylhexyl Palmitate, Glycerin, SD Alcohol 40-B, Butylene Glycol, Glyceryl Stearate, Cetareth-20, Avena Sativa (Oat) Meal Extract, Citric Acid, Cetareth-12, Cetearyl Alcohol, Cetyl Palmitate, Cholesterol, Chlorphenesin, Sodium Hydroxide, Sodium Dehydroacetate.

Questions?

Call **1-800-FOR-AVON**



SKIN SO SOFT BUG GUARD PLUS ITCH RELIEF ITCH RELIEF

menthol spray

Product Information

| | | | |
|-------------------------|----------------|--------------------|----------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:10096-0282 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|--------------|
| Menthol (UNII: L7T10EIP3A) (Menthol - UNII:L7T10EIP3A) | Menthol | 5 mg in 1 mL |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|--------------------------|----------------------|--------------------|
| 1 | NDC:10096-0282-1 | 59 mL in 1 BOTTLE, SPRAY | | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|-------------------------|--|----------------------|--------------------|
| OTC monograph not final | part348 | 06/18/2012 | |

Labeler - Avon Products, Inc. (001468693)

Establishment

| Name | Address | ID/FEI | Business Operations |
|---------------------|---------|-----------|-------------------------|
| Avon Products, Inc. | | 005149471 | manufacture(10096-0282) |

