SNOWSKIN SUNSCREEN ZINC - octinoxate, titanium dioxide, zinc oxide stick Skin Alive, Ltd.

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.

SnowSkin Sunscreen Zinc

Active Ingredients

Octinoxate 7.5%

Titanium Dioxide 2.35%

Zinc Oxide 4.2%

Purpose

Sunscreen

Uses

Helps prevent sunburn

Higher SPF gives more sunburn protection

Provides high protection against sunburn

Regains SPF after 80 minutes of sweating.

Warnings

For external use only

UV exposure from the sun increases the risk of skin cancer, premature skin aging and other skin damage. It is important to decrease UV exposure by limiting time in the sun, wearing protective clothing and using a sunscreen.

When using this product

Keep out of eyes. Rinse with water to remove.

Stop use and ask a doctor

if irritation occurs or rash develops and lasts.

Directions

apply generously 15mins before sun exposure and as needed

Children under 6 months of age: ask a doctor.

Reapply as needed or after towel drying, swimming or sweating.

Inactive Ingredients

Alumina, Aluminum Stearate, Avocado Oil, Beeswax, C12-15 Alkyl Bezoate, Candellia Wax, Capryllic/capric Triglyceride, Cydomethicone, Fragrance, Hydrogenerated Castor Wax, Isohexadecane, Isopropyl Myristate, Isosteric Acid, Lanolin Anhydrous, Microwax, Polyhidroxystearic Acid, Polyhydroxystearic Acid, PPG-3 Benzyl Ether Myristate, Triethylhexanoin.

Snowskin SPF 30+ Sunscreen Zinc gives great protection especially in the snow. Apply liberally to face, nose and lips.

To report a serious adverse event, contact 1-800-332-1088

Expires:

Lot Number:

ww.snowskin.co

DIST. BY Quiver Dist, Bend Oregon, USA 97701

Tested on Snowboarders - Not animals

AS/ANZ STANDARD 2604-1998

Net Wt. 0.30ox (8.5g)

SPF 30+

very water / sweat resistant

broad spectrum

UVA/UVB protection

Proven in extreme new zealand conditions

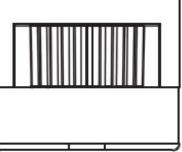
SnowSkin sunscreen Zinc

hands free

Net Wt. 0.30oz (8.5g)

contains no nanoparticles-aminobenzoic acid (PABA) free







Drug Facts

(continued).

Directions

- Apply generously 15mins before sun exposure and as needed
- Children under 6months of age:ask a doctor.
- · Reapply as needed or after towel drying, swimming, or sweating.

Inactive Ingredients

Butylated Hydroxytoluene, Castor Oil, Fragrence, Hydrogenated castor oil, Iron Oxides, Kaolin, Oleyl alcohol, Paraffin Wax, Petroleum jelly, Sunflower Oil, Talc.

To report a serious adverse event, contact 1-800-332-1088

EXPIRES: LOT NUMBER :





- sweat resistant
- broad spectrum
- UVA/UVB





www.snowskin.co

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When using this product

Keep out of eyes. Rinse with water to remove

Stop use and ask a Doctor if Irritation occurs or rash develops and lasts

Keep out of reach of children.

www.snowskin.co

DEST. BY Quiver Dist., Bend Oregon, USA 97701 AS/ANZ STANDARD 2604:1998 Tested on Snowboarders - Not An

Net Wt. .30 oz (8.5g)

Net Wt. .30 oz (8.5g) CONTAINS NO NANOPARTICLES - AMINOBENZOIC ACID (PABA) - FREE

SNOWSKIN SUNSCREEN ZINC

octinoxate, titanium dioxide, zinc oxide stick

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:75916-0634

TOPICAL Route of Administration

Active Ingredient/Active Moiety **Basis of Strength Ingredient Name** Strength OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51) OCTINOXATE 7.5 g in 100 g TITANIUM DIO XIDE (UNII: 15FIX9 V2JP) (TITANIUM DIO XIDE - UNII: 15FIX9 V2JP) TITANIUM DIO XIDE 2.35 g in 100 g ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37) ZINC OXIDE 4.2 g in 100 g

| Inactive Ingredients | | | | |
|---|----------|--|--|--|
| Ingredient Name | Strength | | | |
| BUTYLATED HYDRO XYTO LUENE (UNII: 1P9 D0 Z171K) | | | | |
| CASTOR OIL (UNII: D5340 Y2I9G) | | | | |
| HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY) | | | | |
| IRON (UNII: E1UOL152H7) | | | | |
| KAOLIN (UNII: 24H4NWX5CO) | | | | |
| OLEYL ALCOHOL (UNII: 172F2WN8DV) | | | | |
| PARAFFIN (UNII: 19 O 0 E 3 H 2 Z E) | | | | |
| PETROLATUM (UNII: 4T6H12BN9U) | | | | |

SUNFLOWER OIL (UNII: 3W1JG795YI)

TALC (UNII: 7SEV7J4R1U)

| Packaging | | | | | | |
|-----------|------------------|----------------------------------|----------------------|--------------------|--|--|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date | | |
| 1 | NDC:75916-0634-1 | 1 in 1 BOX | | | | |
| 1 | | 8.5 g in 1 TUBE, WITH APPLICATOR | | | | |

| Marketing Inform | arketing Information | | | | | | |
|-------------------------|--|----------------------|--------------------|--|--|--|--|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | | | | |
| OTC monograph not final | part352 | 0 1/27/20 11 | | | | | |
| | | | | | | | |

Labeler - Skin Alive, Ltd. (593384746)

Revised: 1/2011 Skin Alive, Ltd.