LIP BALM MINT FLAVORED- spf 15 lipstick Webb Business Promotions

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.

SPF 15 Lip Balm Mint Flavored

Purpose

Purpose

Sun Screen

Drug Facts

Octinoxate 3.6% Octisalate 2.5% Benzophenone 1.5%

Inactive Ingredients

Hydrogenated Soybean Oil, Sunflower Oil, White Beeswax Pastilles, Cocoa Butter, Shea Butter, Vitamin E, Aloe Vera Gel, Mint Flavoring.

Warnings

- **Skin Cancer/Skin Aging Alert:** Spending too much time in the sun increases your risk of skin cancer and early skin aging. This product has been shown to only help prevent sunburn, not skin cancer or early skin aging.
- For external use only.
- **Do not use** on damaged or broken skin.
- **Stop use and ask doctor** if rash occurs.
- When using this product keep out of eyes.
- **Keep out of reach of children.** If swallowed get medical help or contact posion control center right away.

Directions

- Apply liberally 15 minutes before sun exposure
- Use a water resistant sunscreen if swimming or sweating
- Reapply every 2 hours
- Children under 6 months : Ask a doctor

Storage and Handling Information

Protect this product from excessive heat and direct sun

MINT FLAVORED SPF 15 NATURAL BEESWAX LIP BALM

MINT FLAVORED SPF 15 NATURAL BEESWAX LIP BALM

| DRUG FACTS | Net Wt. : 0.15 oz / 4.2 g | | | |
|--|---------------------------|----------------|---|--|
| Active Ingredien Octinoxate Octyl Salicylate Oxybenzone | 3.6% 2.5% 1.5% | Sunscreen | Inactive Ingredients: Hydrogenated Soybean Oil, Sunflower Oil, White Bees Wax Pastilles, Cocoa Butter, Shea Butter, Vitamin E, Aloe Vera Gel, Mint Flavoring. | |
| Uses Helps prev | ent sundi | Irn | Gei, Mini Flavoring. | |
| Warnings | Aging Ale | rt. Sponding | time in the sun increases your | |
| | | | This product has been shown | |
| | | | ancer or early skin aging. | |
| For external use | | rn, not skin t | ancer or early skin aging. | |
| Do not use on dat | | brokon skin | | |
| | - | | | |
| Stop use and ask When using this is | | | | |
| | | | ves. Rinse with water to remove. | |
| Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. | | | | |
| | Lenter rig | nt away. | | |
| Directions | | | | |
| Apply liberally 1 | .5 minute | es before si | in exposure | |
| Use a water resistant sunscreen if swimming or sweating | | | | |
| Reapply at least | every 2 | hours | | |
| Children under | 6 month | s: Ask a doo | tor | |
| Other Informed | tion: Dra | | | |
| direct sun | ion: Pro | tect this pro | duct from excessive heat and | |

LIP BALM MINT FLAVORED

spf 15 lipstick

| Product Information | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:70445-243 |
| Route of Administration | TOPICAL | | |

Strength

3.6 g in 1 g

2.5 g in 1 g

1.5 g in 1 g

OXYBENZONE

Active Ingredient/Active Moiety Ingredient Name Basis of Strength OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51) OCTINOXATE OCTISALATE (UNII: 4X49 Y0596W) (OCTISALATE - UNII:4X49 Y0596W) OCTISALATE

OXYBENZONE (UNII: 9500S7VE0Y) (OXYBENZONE - UNII:9500S7VE0Y)

| Strength | T 11 . N T |
|----------|---|
| Ŭ | Ingredient Name |
| | IYDRO GENATED SO YBEAN OIL (UNII: A2M91M918C) |
| | COCOA BUTTER (UNII: 5120 YT1CRR) |
| | HEA BUTTER (UNII: K49155WL9Y) |
| | LOE VERA LEAF (UNII: ZY81Z83H0X) |
| | |

| WHITE WAX (UNII: 7G1J5DA97F) | | | | | | | |
|----------------------------------|--|-----------------|----------------------|----------------------|-----------|----------|--|
| SUNFLOWER OIL (UNII: 3W1JG795YI) | | | | | | | |
| | | | | | | | |
| | | | | | | | |
| Product Characteristics | | | | | | | |
| Color | | hite | Score | | | | |
| Shape | | | Size | | | | |
| Flavor | M | IINT | Imprint Code | | | | |
| Contains | | | | | | | |
| | | | | | | | |
| | | | | | | | |
| Packaging | | | | | | | |
| # Item Code | Package Description | | Marketing Start Date | Marketing End Date | | | |
| 1 NDC:70445-243-01 5 g | 5 g in 1 TUBE; Type 0: Not a Combination Product 09/11/200 | | | 09/11/2009 | | | |
| | | | | | | | |
| | | | | | | | |
| Marketing Information | | | | | | | |
| Marketing Category | Application N | Number or Monog | aph Citation | Marketing Start Date | Marketing | End Date | |
| OTC monograph final pa | rt352 | | | 0 1/0 1/20 0 9 | | | |
| | | | | | | | |

Labeler - Webb Business Promotions (154445647)

| Establishment | | | | | | |
|--------------------------|---------|-----------|------------------------|--|--|--|
| Name | Address | ID/FEI | Business Operations | | | |
| Webb Business Promotions | | 154445647 | manufacture(70445-243) | | | |

Revised: 3/2017

Webb Business Promotions