SOFTLIPS WATERMELON- dimethicone, octinoxate, octisalate, oxybenzone stick The Mentholatum Company

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

Dimethicone 2%

Octinoxate 7.5%

Octisalate 3%

Oxybenzone 3%

Purpose

Dimethicone - Skin protectant

Octinoxate - Sunscreen

Octisalate - Sunscreen

Oxybenzone - Sunscreen

Keep Out of Reach of Children

If swallowed, get medical help or contact a Poison Control Center right away.

Uses

- helps prevent sunburn
- temporarily protects chapped or cracked lips

Warnings

Skin Cancer/Skin Aging Alert: Spending time in the sun increased your risk of skin cancer and early skin aging. This product has been shown only to prevent sunburn, not skin cancer or early skin aging.

For external use only.

Do not use

on damaged or broken skin

When using this product

keep out of eyes. Rinse with water to remove.

Stop use and ask a doctor if

- rash occurs
- condition worsens
- symptoms last more than 7 days or clear up and occur again within a few days

Directions

- apply liberally 15 minutes before sun exposure
- use a water resistant sunscreen if swimming or sweating
- reapply at least every 2 hours
- children under 6 months: ask a doctor

Inactive ingredients

ozokerite, squalane, ethylhexyl palmitate, petrolatum, myristyl myristate, myristyl lactate, flavor, cetyl alcohol, myristyl laurate, myristyl alcohol, BHT, menthol, tocopheryl acetate [vitamin E]

Package/Label Principal Display Panel



Principal Display Panel

softlips[®]



All Softlips⁸ products are gluten-free.

Drug Facts

Active ingredients Purpose

Dimethicone 2%......Skin protectant
Octinoxate 7.5%....Sunscreen
Octisalate 3%....Sunscreen
Oxybenzone 3%....Sunscreen

Softlips® has a lip balm for your every mood and moment. Just glide it on, and it's instant happiness! With essential moisturizers plus SPF protection— it's so smart. So irresistable. So you."

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- apply liberally 15 minutes before sun exposure
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- reapply at least every 2 hours
- children under 6 months: ask a doctor

Other information

protect this product from excessive heat and direct sun

Inactive ingredients

ozokerite, squalane, ethylhexyl palmitate, petrolatum, myristyl myristate, myristyl lactate, flavor, cetyl alcohol, myristyl laurate, myristyl alcohol, BHT, menthol, tocopheryl acetate [vitamin E]

Questions?

Toll free 1-877-636-2677 MON-FRI 9 AM to 5 PM (EST) softlips.com



The Mentholatum Company Orchard Park, NY 14127



SOFTLIPS WATERMELON

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for a purchase proof of fig. to The Mentholatum Company .12 岩 SATISFACTION: IF 6 GUARANTEE

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10742-8476
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
DIMETHICONE (UNII: 92RU3N3Y10) (DIMETHICONE - UNII:92RU3N3Y10)	DIMETHICONE	20 mg in 1 g	
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 mg in 1 g	
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	30 mg in 1 g	
OXYBENZONE (UNII: 9500S7VE0Y) (OXYBENZONE - UNII:9500S7VE0Y)	OXYBENZONE	30 ug in 1 g	

Inactive Ingredients		
Ingredient Name	Strength	
CERESIN (UNII: Q1LS2UJO3A)		
SQUALANE (UNII: GW89575KF9)		
ETHYLHEXYL PALMITATE (UNII: 2865993309)		
PETROLATUM (UNII: 4T6H12BN9U)		
MYRISTYL MYRISTATE (UNII: 4042ZC00DY)		
MYRISTYL LACTATE (UNII: 1D822OC34X)		
CETYL ALCOHOL (UNII: 936JST6JCN)		
MYRISTYL LAURATE (UNII: 58U0NZN2BT)		
MYRISTYL ALCOHOL (UNII: V42034O9PU)		
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)		
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742- 8476-1	1 in 1 BLISTER PACK	04/16/2012	
1		2 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:10742- 8476-2	2 in 1 BLISTER PACK	12/01/2016	
2		2 g in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:10742- 8476-3	3 in 1 BLISTER PACK	04/06/2020	
3		2 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information			
Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date

OTC monograph final	part352	04/16/2012	

Labeler - The Mentholatum Company (002105757)

Registrant - The Mentholatum Company (002105757)

Establishment			
Name	Address	ID/FEI	Business Operations
The Mentholatum Company		002105757	manufacture(10742-8476) , label(10742-8476)

Revised: 2/2023 The Mentholatum Company