MENTHOLATUM NATURAL ICE CHERRY, SPF 15 CHERRY- dimethicone, octinoxate, octisalate stick The Mentholatum Company

Drug Facts - Mentholatum Natural Ice Cherry, SPF 15

Active ingredients

Dimethicone 1% Octinoxate 7.5% Octisalate 5%

Purpose

Dimethicone - Skin protectant

Octinoxate - Sunscreen

Octisalate - Sunscreen

Uses

- helps prevent sunburn
- temporarily protects chapped or cracked lips
- protect from the drying effects of wind and cold weather

Warnings

Skin Cancer/Skin Aging Alert: Spending time in the sun increases your risk of skin cancer and early skin aging. This product has been shown only to help 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

Keep out of reach of children

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

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

Other information

• protect this product from excessive heat and direct sun

Inactive ingredients

mineral oil, ozokerite, petrolatum, lanolin, flavor, camphor, menthol

Questions?

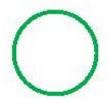
1-877-636-2677

MON-FRI 9AM to 5PM (EST) mentholatum.com

Package Label/Principal Display Panel







CHERRY .



NOT TESTED ON ANIMALS





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Active ingredients Purpose

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Octinoxate 7.5%.....Sunscreen
Octisalate 5%.....Sunscreen

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Drug Facts (continued)

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Questions? 1-877-838-2877 MON-FRI 9AM to 5PM (EST)



The Mentholstum Co. Orchard Park, NY 14127 EE738008 G2024





stain this card for complete information



MENTHOLATUM NATURAL ICE CHERRY, SPF 15 CHERRY

dimethicone, octinoxate, octisalate stick

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10742-3075
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 g	
DIMETHICONE (UNII: 92RU3N3Y10) (DIMETHICONE - UNII:92RU3N3Y10)	DIMETHICONE	10 mg in 1 g	
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 mg in 1 g	

Inactive Ingredients		
Ingredient Name	Strength	
MINERAL OIL (UNII: T5L8T28FGP)		
FRAGRANCE 13576 (UNII: 5EM498GW35)		
CERESIN (UNII: Q1LS2UJO3A)		
MENTHOL (UNII: L7T10EIP3A)		
CAMPHOR, (-)- (UNII: 213N3S8275)		
PETROLATUM (UNII: 4T6H12BN9U)		
LANOLIN (UNII: 7EV65EAW6H)		

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742- 3075-1	1 in 1 BLISTER PACK	09/01/2024	
1		4.2 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	09/01/2024	

Labeler - The Mentholatum Company (002105757)

Registrant - The Mentholatum Company (002105757)

Establishment				
Name	Address	ID/FEI	Business Operations	
The Mentholatum Company		002105757	manufacture(10742-3075)	

Revised: 6/2025 The Mentholatum Company