

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

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**Drug Facts - Mentholatum Natural Ice**

**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

**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 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

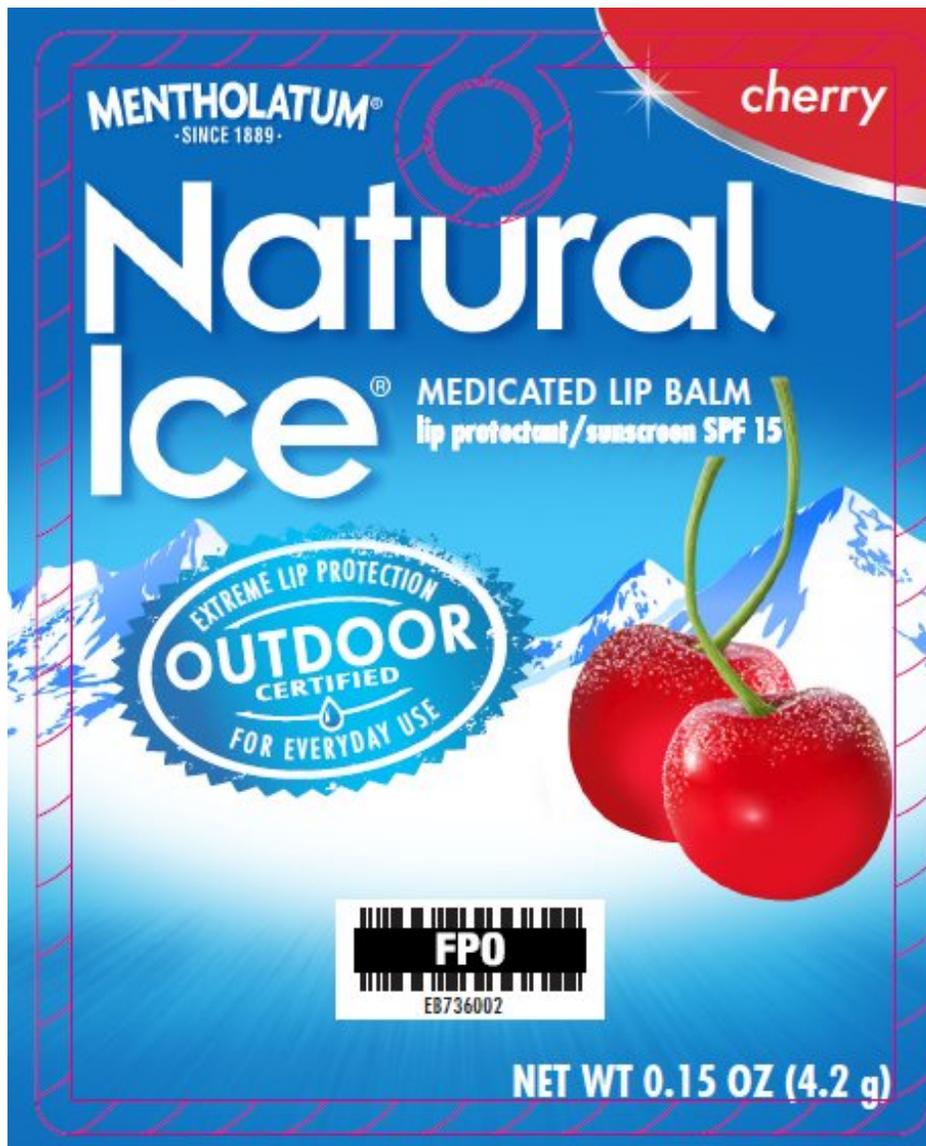
**Questions?**

**1-877-636-2677**

MON-FRI 9AM to 5PM (EST)

mentholatum.com

**Package/Label Principal Display Panel**



**Package/Label Principal Display Panel**

# Natural Ice<sup>®</sup> CHERRY

Medicated Lip Protectant/Sunscreen SPF 15



Manufactured By:  
The Mentholatum Co.  
Orchard Park, NY 14127



## Drug Facts

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Octinoxate 7.5%	Sunscreen
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Retain this card for complete information.

## Drug Facts (continued)

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## MENTHOLATUM NATURAL ICE CHERRY

dimethicone, octinoxate, octisalate stick

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10742-1048
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE	10 mg in 1 g
<b>OCTINOXATE</b> (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 mg in 1 g
<b>OCTISALATE</b> (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 g

### Inactive Ingredients

Ingredient Name	Strength
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<b>MINERAL OIL</b> (UNII: T5L8T28FGP)	
<b>CERESIN</b> (UNII: Q1LS2UJO3A)	
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)	
<b>LANOLIN</b> (UNII: 7EV65EAW6H)	
<b>CAMPHOR (SYNTHETIC)</b> (UNII: 5TJD82A1ET)	
<b>LEVOMENTHOL</b> (UNII: BZ1R15MTK7)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742-1048-1	1 in 1 BLISTER PACK	08/19/1999	04/01/2019
1		4.5 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:10742-1048-9	4.5 g in 1 TUBE; Type 0: Not a Combination Product	08/19/1999	04/01/2019
3	NDC:10742-1048-2	1 in 1 BLISTER PACK	04/02/2019	
3		4.2 g in 1 TUBE; Type 0: Not a Combination Product		
4	NDC:10742-1048-8	4.2 g in 1 TUBE; Type 0: Not a Combination Product	04/02/2019	
5	NDC:10742-1048-3	1 in 1 BLISTER PACK	08/19/1999	
5		4.5 g in 1 TUBE; Type 0: Not a Combination Product		
6	NDC:10742-1048-4	4.5 g in 1 TUBE; Type 0: Not a Combination Product	08/19/1999	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	08/19/1999	

**Labeler** - The Mentholatum Company (002105757)

**Registrant** - The Mentholatum Company (002105757)

### Establishment

Name	Address	ID/FEI	Business Operations
The Mentholatum Company		002105757	manufacture(10742-1048)