MENTHOLATUM NATURAL ICE ORIGINAL- dimethicone, octinoxate, octisalate ointment
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 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, fragrance, menthol

Questions?

1-877-636-2677

MON-FRI 9AM to 5PM (EST)

mentholatum.com

Principal Display Panel

Principal Display Panel

MENTHOLATUM NATURAL ICE ORIGINAL

dimethicone, octinoxate, octisalate ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10742-3004
Route of Administration	TOPICAL		

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

Inactive Ingredients	
Ingredient Name	**Strength**

MINERAL OIL (UNII: T5L8T28FGP)	
CERESIN (UNII: Q1LS2UJO3A)	
PETROLATUM (UNII: 4T6H12BN9U)	
LANOLIN (UNII: 7EV65EAW6H)	
LEVOMENTHOL (UNII: BZ1R15MTK7)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742-3004-1	1 in 1 BLISTER PACK	01/19/1999	04/01/2019
1		4.5 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:10742-3004-9	4.5 g in 1 TUBE; Type 0: Not a Combination Product	01/19/1999	04/01/2019
3	NDC:10742-3004-2	1 in 1 BLISTER PACK	04/02/2019	
3		4.2 g in 1 TUBE; Type 0: Not a Combination Product		
4	NDC:10742-3004-8	4.2 g in 1 TUBE; Type 0: Not a Combination Product	04/02/2019	
5	NDC:10742-3004-3	1 in 1 BLISTER PACK	01/19/1999	
5		4.5 g in 1 TUBE; Type 0: Not a Combination Product		
6	NDC:10742-3004-4	4.5 g in 1 TUBE; Type 0: Not a Combination Product	01/19/1999	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	01/19/1999	

Labeler - The Mentholatum Company (002105757)

Registrant - The Mentholatum Company (002105757)

Establishment

Name	Address	ID/FEI	Business Operations
The Mentholatum Company		002105757	manufacture(10742-3004)

Revised: 2/2023 The Mentholatum Company