BLISTEX MEDICATED- dimethicone, camphor (synthetic), menthol, and phenol ointment
Blistex Inc.

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.

Blistex®
Medicated Lip ointment

Drug Facts

Active ingredients	***Purpose***
Camphor 0.5% (w/w)	External analgesic
Dimethicone 1.1% (w/w)	Lip protectant
Menthol 0.625% (w/w)	External analgesic
Phenol 0.5% (w/w)	External analgesic

Uses
- for the temporary relief of pain and itching associated with minor lip irritation or cold sores
- temporarily protects and helps relieve chapped or cracked lips

Warnings

For external use only

Do not use on
- deep or puncture wounds
- animal bites
- serious burns

When using this product
- do not get into eyes
- do not apply over large areas of the body or bandage

Stop use and ask a doctor if
- 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

- adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily
- children under 2 years of age: Consult a doctor

Inactive ingredients

allantoin, ammonium hydroxide, beeswax, calcium disodium EDTA, calcium hydroxide, cetyl alcohol, flavors, glycerin, hydrated silica, lanolin, lauric acid, mineral oil, myristic acid, oleic acid, palmitic acid, paraffin, petrolatum, polyglyceryl-3 diisostearate, potassium hydroxide, purified water, SD alcohol 36, sodium hydroxide, sodium saccharin, stearyl alcohol

PRINCIPAL DISPLAY PANEL - 6 g Tube Carton

Penetrating
Medication
And Moisture

Blistex®
MEDICATED LIP OINTMENT

Blistex®

LIP PROTECTANT/
EXTERNAL ANALGESIC

MEDICATED LIP OINTMENT

Helps Heal Severe Dryness
And Relieve Cold Sores

Clinically Shown
To Help Heal Dry, Chapped Lips

Net Wt. 0.21 oz. (6 g)

SATISFACTION GUARANTEED
Blistex®

Penetrating
Medication
And Moisture

0 41388 21021 9

dilsostearate, potassium hydroxide, purified water, SD alcohol 36, sodium hydroxide, sodium saccharin, stearyl alcohol

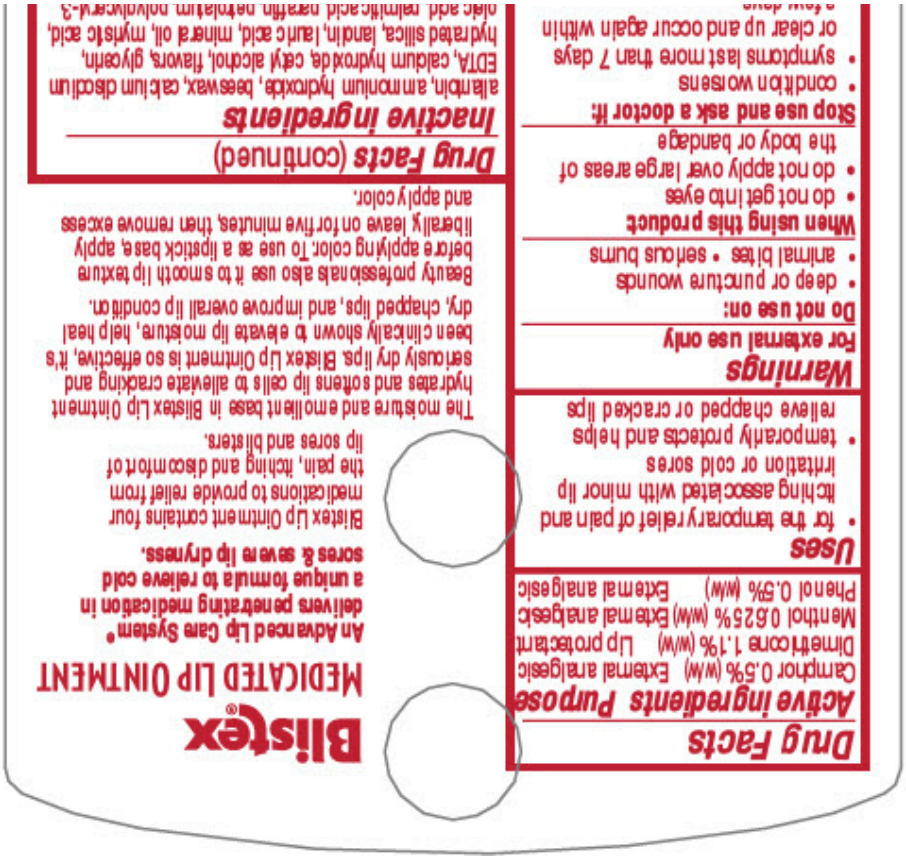
P.O. Box 5392, Oak Brook, IL 60522-5392 #C030261

a few days

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily
- children under 2 years of age: Consult a doctor



BLISTEX MEDICATED

dimethicone, camphor (synthetic), menthol, and phenol ointment

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10157-9951
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Dimethicone (UNII: 92RU3N3Y1O) (Dimethicone - UNII:92RU3N3Y1O)	Dimethicone	1.1 g in 100 g
Camphor (synthetic) (UNII: 5TJD82A1ET) (Camphor (synthetic) - UNII:5TJD82A1ET)	Camphor (synthetic)	0.5 g in 100 g
Menthol, unspecified form (UNII: L7T10EIP3A) (Menthol, unspecified form - UNII:L7T10EIP3A)	Menthol, unspecified form	0.625 g in 100 g
Phenol (UNII: 339NCG44TV) (Phenol - UNII:339NCG44TV)	Phenol	0.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
allantoin (UNII: 344S277G0Z)	
ammonia (UNII: 5138Q19F1X)	
yellow wax (UNII: 2ZA36H0S2V)	

edetate calcium disodium anhydrous (UNII: 8U5D034955)	
calcium hydroxide (UNII: PF5DZW74VN)	
cetyl alcohol (UNII: 936JST6JCN)	
glycerin (UNII: PDC6A3C0OX)	
hydrated silica (UNII: Y6O7T4G8P9)	
lanolin (UNII: 7EV65EAW6H)	
lauric acid (UNII: 1160N9NU9U)	
mineral oil (UNII: T5L8T28FGP)	
myristic acid (UNII: 0I3V7S25AW)	
oleic acid (UNII: 2UMI9U37CP)	
palmitic acid (UNII: 2V16EO95H1)	
paraffin (UNII: I9O0E3H2ZE)	
petrolatum (UNII: 4T6H12BN9U)	
polyglyceryl-3 diisostearate (UNII: 46P231IQV8)	
potassium hydroxide (UNII: WZH3C48M4T)	
water (UNII: 059QF0KO0R)	
sodium hydroxide (UNII: 55X04QC32I)	
saccharin sodium (UNII: SB8ZUX40TY)	
stearyl alcohol (UNII: 2KR89I4H1Y)	

Product Characteristics

Color	WHITE	**Score**	
Shape		**Size**	
Flavor	MINT	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10157-9951-1	6 g in 1 TUBE; Type 0: Not a Combination Product	12/01/2000	
2	NDC:10157-9951-3	10 g in 1 TUBE; Type 0: Not a Combination Product	12/01/2000	
3	NDC:10157-9951-4	2.25 g in 1 TUBE; Type 0: Not a Combination Product	12/01/2000	
4	NDC:10157-9951-5	8 g in 1 TUBE; Type 0: Not a Combination Product	12/01/2000	
5	NDC:10157-9951-2	1 in 1 CARTON	12/01/2000	
5	NDC:10157-9951-6	6 g in 1 TUBE; Type 0: Not a Combination Product		
6	NDC:10157-9951-7	3 in 1 CARTON	12/01/2000	
6		6 g in 1 TUBE; Type 0: Not a Combination Product		
7	NDC:10157-9951-8	4 in 1 CARTON	12/01/2000	
7		6 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 NOT FINAL	part348	12/01/2000	

Labeler - Blistex Inc. (005126354)

Establishment

Name	Address	ID/FEI	Business Operations
Blistex Inc.		005126354	MANUFACTURE(10157-9951)

Revised: 12/2021 Blistex Inc.