

BLUESTOP MAX- external analgesic cream
Clavel Corporation

BlueStop Max Cream

Active Ingredients

Lidocaine HCl 4%

Menthol 1%

Purpose

Topical Analgesic

Use

Temporary relief of pain

Warnings

For external use only

When using this product: avoid contact with the eyes

Stop use and ask a doctor if: condition worsens, or symptoms persist for 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

Other Information

- Store at room temperature
- Avoid excessive heat

Inactive Ingredients

Aloe Vera Leaf, Alpha-Tocopherol Acetate, Ammonium Acryloyldimethyltaurate, Dimethylacrylamide, Lauryl Methacrylate and Laureth-4 Methacrylate Copolymer, Trimethylolpropane Triacrylate Crosslinked (45000 Mpa.S), Cetyl Alcohol, Coconut Oil,

Dimethyl Sulfone, Ethylhexylglycerin, Ethylhexyl Palmitate, Fragrance, Garcinia Indica Seed Butter, Glucosamine Sulfate, Glycerin, Melaleuca Alternifolia Leaf, Phenoxyethanol, Sodium Acrylate/Sodium Acryloyldimethyltaurate Copolymer (4000000 MW), Vitamin A Palmitate, Water, Witch Hazel

Questions or Comments?

1-800-432-5464

Tube and Carton Labels

4% LIDOCAINE + 1% MENTHOL

BLUESTOP[®] MAX

DUAL ACTION PAIN RELIEF CREAM

NET WT 3 OZ (85 G)

Drug Facts

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Manufactured exclusively for and distributed by:

CLAVÉL CORPORATION

4150 E. OVERLAND TRAIL
ABILENE, TEXAS 79601

1.800.HEALING
BLUESTOPMAX.COM



WOMEN
OWNED



Blue Stop Max
Lidocaine + Menthol
Crepe Tube & Analgesics
Roll on STE
2.594 x 1.562 x 0.188
Ver 1 Rev 2.018 CIB SBS
11.672 Grain Dir

0.875

1.947

0.969

1.563

2.594

1.563

2.563

0.656

1.508

2.422

0.719

10.933

6.186

1.516

0.672



BLUESTOP MAX

external analgesic cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82714-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z 41A) (LIDOCAINE -	LIDOCAINE HYDROCHLORIDE	40 mg

UNII:98PI200987)	ANHYDROUS	in 1 g
MENTHOL, (+)- (UNII: C6B1OE8P3W) (MENTHOL, (+)- - UNII:C6B1OE8P3W)	MENTHOL, (+)-	10 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
COCONUT OIL (UNII: Q9L0O73W7L)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)	
ETHYLHEXYL PALMITATE (UNII: 2865993309)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
MELALEUCA ALTERNIFOLIA LEAF (UNII: G43C57162K)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE, DIMETHYLACRYLAMIDE, LAURYL METHACRYLATE AND LAURETH-4 METHACRYLATE COPOLYMER, TRIMETHYLOLPROPANE TRIACRYLATE CROSSLINKED (45000 MPA.S) (UNII: Q7UI015FF9)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0K00R)	
GARCINIA INDICA SEED BUTTER (UNII: US2H3D7800)	
WITCH HAZEL (UNII: 101I4J0U34)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM ACRYLATE/SODIUM ACRYLOYLDIMETHYLTAURATE COPOLYMER (4000000 MW) (UNII: 1DXE3F3OZX)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82714-002-01	1 in 1 CARTON	01/10/2023	
1		85 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	M017	01/10/2023	

Labeler - Clavel Corporation (927855643)

Establishment

Name	Address	ID/FEI	Business Operations
TAKA USA dba Cosmetic Innovations		802860515	manufacture(82714-002)

