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 yeas of age: Consult a doctor

Other Information

- Store at room temperature
- Avoidexcessive heat

Inactive Ingredients

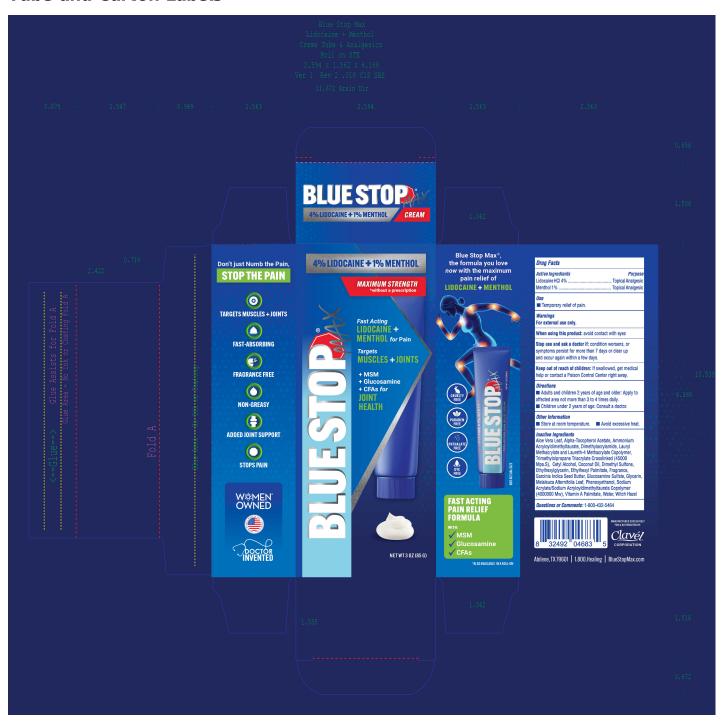
Aloe Vera Leaf, Alpha-Tocopherol Acetate, Ammonium Acryloyldimethyltaurate, Dimethylacrylamide, Lauryl Methacrylate and Laureth-4 Methacrylate Copolymer, Trimethylolpropane Triacrylate Crosslinked (45000 Mpa.S), Cetyl Alcohol, Coconut Oil,

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



% LIDOCAINE + 1% MENTHO

NET WT 3 0Z (85 G)

CREAM Щ ~ ACTION PAIN DUAL

Drug Facts

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Questions or Comments: 1-800-432-5464

Manufactured exclusively for and distributed by:

- CLAVÉL CORPORATION -

4150 E. OVERLAND TRAIL ABILENE, TEXAS 79601

1.800.HEALING BLUESTOPMAX.COM













BLUESTOP MAX

external analgesic cream

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Product I	nform	ation

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: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	40 mg in 1 g	
MENTHOL, (+)- (UNII: C6B10E8P3W) (MENTHOL, (+) UNII:C6B10E8P3W)	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: 059QF0KO0R)	
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)	

Revised: 1/2024 Clavel Corporation