DZUL ARTHRITIS PAIN RELIEF- menthol cream
Caball Sales, Inc.

83037-015-04

Menthol 1.5%

Topical Analgesic

USE

For the temporary relief of minor aches and pains of muscles and joints associated arthritis

•For external use only.

•Ask a doctor before use if you have redness over affected area.

Avoid contact with the eyes or mucous membranes.
Do not apply to wounds or damaged skin.
Do not apply to the irritated skin or if excessive irritation develops.
Do not bandage tightly.
Do not use with heating pad or device.

Stop use and ask a doctor if condition worsens, or if symptoms persist for more than 7 days, or clear up and occur again within a few days.

If pregnant or breast – feeding

Ask a health professional before use.

If swallowed, get medical help, or contact a Poison Control Center right away.

DIRECTIONS

Adults and Children over 12 years

Apply a small amount on the affected area.
Massage in circular motion, let set for a few seconds.
Repeat as necessary, but no more than 3 to 4 times daily.

Children under 12 years of age: Do not use, consult a doctor.

OTHER INFORMATION

Store tightly closed in a dry place at controlled room temperature between 59-86 F (15°-30° C).

Water (Aqua), Paraffinum Liquidum, Alcohol Denat, Cetearyl Alcohol, Cetyl Alcohol, Stearic Acid, Dimethicone, Glyceryl Stearate, Eucalyptus Globulus Leaf Oil, Glycereth-26, Polysorbate 60, Methylsulfonylmethane (MSM), Sodium Chondroitin Sulfate, Glucosamine Sulfate, Sodium Hyaluronate, Sodium PCA, Wheat Amino Acids, Panthenol, Symphytum Officinale (Comfrey) Extract, Hydroxyproline, Acrylamide/Sodium Acrylate Copolymer,

Trideceth-6, Stearyl Alcohol, Polysorbate 20, Caprylyl Glycol, Sodium Hydroxide, Phenoxyethanol, Hexylene Glycol.

DZUL ARTHRITIS PAIN RELIEF CREAM

DZUL ARTHRITIS PAIN RELIEF

menthol cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83037-015
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	1.5 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
EUCALYPTUS OIL (UNII: 2R04ONI662)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
MINERAL OIL (UNII: T5L8T28FGP)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
ACRYLIC ACID/SODIUM ACRYLATE COPOLYMER (1:1; 600 MPA.S AT 0.2%) (UNII: M4PPW69Y4H)	
AMINO ACIDS, WHEAT (UNII: 0370GZL32F)	
PANTHENOL (UNII: WV9CM0O67Z)	
ALCOHOL (UNII: 3K9958V90M)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
GLYCERETH-26 (UNII: NNE56F2N14)	

SODIUM CHONDROITIN SULFATE (PORCINE; 5500 MW) (UNII: H5BJH23Z9A)	
GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SYMPHYTUM OFFICINALE WHOLE (UNII: H8FJJ6KX5Y)	
HYDROXYPROLINE (UNII: RMB44WO89X)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
TRIDECETH-6 (UNII: 3T5PCR2H0C)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83037-015-04	114 g in 1 JAR; Type 0: Not a Combination Product	04/20/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	04/20/2024	

Labeler - Caball Sales, Inc. (047422138)

Revised: 1/2026 Caball Sales, Inc.