#### BIOFREEZE COLORLESS- menthol gel A-S Medication Solutions

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.

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#### **Biofreeze<sup>®</sup> Colorless**

**Drug Facts** 

## **Active ingredient**

Menthol 4%

## Purpose

Pain Relieving Gel

#### Uses

Temporarily relieves minor aches and pains of muscles and joints associated with:

- simple backache
- arthritis
- strains
- bruises
- sprains

# Warnings

For external use only.

# Flammable: Keep away from excessive heat or open flame

# When using this product

- use only as directed
- avoid contact with the eyes or on mucous membranes
- do not apply to wounds or damaged skin
- do not apply to irritated skin or if excessive irritation develops
- do not bandage tightly or use with heating pad or device

# Stop use and ask a doctor if

- you experience pain, swelling or blistering of the skin
- condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days
- arthritic pain persists for more than 10 days, or redness is present, or in conditions affecting children under 12 years of age

If pregnant or breast-feeding, ask a health professional before use.

**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: rub a thin film over affected area not more than 3 to 4 times daily
- children under 2 years of age: consult a physician
- wash hands after use with cool water

## Other information

- store at 20-25°C (68-77°F)
- store in a cool dry place away from direct sunlight

## Inactive ingredients

Aloe Barbadensis Leaf Extract, Arctium Lappa Root (Burdock) Extract, Arnica Montana Flower Extract, Boswellia Carterii Resin Extract, Calendula Officinalis Extract, Camellia Sinensis Leaf Extract, Camphor, Carbomer, Glycerin, Ilex Paraguariensis Leaf Extract, Isopropyl Alcohol, Isopropyl Myristate, Melissa Officinalis (Lemon Balm) Leaf Extract, Silica, Tocopheryl Acetate, Triethanolamine, Water

## **Questions or comments?**

## 1-800-246-3733

Dist. by: RB Health (US), Parsippany, NJ 07054-0224

## HOW SUPPLIED

Product: 50090-6361

NDC: 50090-6361-0 89 mL in a BOTTLE

## **BIOFREEZE COLORLESS**

BI	OT IOFREEZE DOL THE PAIN
ME	TVE INGREDIENTS: NTHOL USP 4% IPOSE: COOLING PAIN RELIEF
	ORE IN A COOL DRY PLACE VAY FROM DIRECT SUNLIGHT
	DLORLESS GEL FL. 0Z./89 ML
144.7	GTIN: 00350090636109
A-	S Medication Solutions ertyville, IL 60048
	URCE NDC: 59316 - 103 - 12

menthol gel								
Product Information								
Product Type	HUMAN OTC DRUG	Item Code (Sou	urce)	NDC:50090-636	1(NDC:59316-103)			
Route of Administration	TOPICAL	•	-					
Active Ingredient/Active	e Moiety							
Ingred	ient Name		Basis	of Strength	Strength			
MENTHOL (UNII: L7T10EIP3A) (MI	ENTHOL - UNII:L7T10EI	P3A) ME	ENTHOL		40 mg in 1 mL			
Inactive Ingredients	Ingredient Na	me			Strength			
Inactive Ingredients								
		Strength						
ALOE VERA LEAF (UNII: ZY81Z83H0X)								
ARCTIUM LAPPA ROOT (UNII: 597E9BI3Z3)								
ARNICA MONTANA FLOWER (UNII: OZ 0E5Y15PZ)								
FRANKINCENSE (UNII: R9XLF1R1	WM)							
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)								
GREEN TEA LEAF (UNII: W2ZU1RY8B0)								
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)								
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)								
GLYCERIN (UNII: PDC6A3C0OX)								
	NII: 10953B404F)							
ILEX PARAGUARIENSIS LEAF (U	<b>x</b> = = = 7		ISOPROPYL ALCOHOL (UNII: ND2M416302)					
ISOPROPYL ALCOHOL (UNII: ND	2M416302)							
ISOPROPYL ALCOHOL (UNII: ND ISOPROPYL MYRISTATE (UNII: C	2M416302) RE8K4LNJS)							
ISOPROPYL ALCOHOL (UNII: ND ISOPROPYL MYRISTATE (UNII: 0 MELISSA OFFICINALIS LEAF (U	2M416302) RE8K4LNJS) NII: 50D2ZE9219)							
	2M416302) RE8K4LNJS) NII: 50D2ZE9219) (BU4)							
ISOPROPYL ALCOHOL (UNII: ND ISOPROPYL MYRISTATE (UNII: C MELISSA OFFICINALIS LEAF (U SILICON DIOXIDE (UNII: ETJ7Z6)	2M416302) RE8K4LNJS) NII: 50D2ZE9219) (BU4) <b>E</b> (UNII: 9E8X80D2L0)							

Packaging						
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date		
	NDC:50090- 6361-0	89 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/08/2023			
Μ	arketing	Information				
Μ	arketing Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		

Labeler - A-S Medication Solutions (830016429)

Establishment							
Name	Address	ID/FEI	<b>Business Operations</b>				
A-S Medication Solutions		830016429	RELABEL(50090-6361)				

Revised: 2/2023

A-S Medication Solutions