PAIN RELIVING ROLL-ON- menthol gel KAREWAY PRODUCT, 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.

Pure-Aid Pain Relievng Gel Roll-On

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, Camellia Sinensis (Green Tea) Leaf Extract, Camphor, Carbomer, Centella Asiatica Extract, Ethylhexylglycerin, FD&C Blue No.1, FD&C Yellow No.5, IIex Paraguariensis Leaf Extract, Isopropyl Alcohol, Isopropyl Myristate, Phenoxyethanol, Propylene Glycol, Tocopheryl Acetate, Trolamine, Water

Exclusively distributed by:

Kareway Product, Inc.

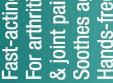
Compton, CA 90220

www.kareway.com

PRINCIPAL DISPLAY PANEL







Drug Facts

Active Ingredient Menthol 4%Pain Relieving Gel

Purpose

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Warnings

For external use only

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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.

*This product is not manufactured or distributed by RB Health(US) LLC, owner of the registered trademark Biofreeze[®].

Drug Facts (continued)

Directions

VANISHIN

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)

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Exclusively distributed by: Kareway Product Inc. Compton, CA 90220 www.kareway.com

MADE IN CHINA



PAIN RELIVING ROLL-ON

menthol gel

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Ρ	roduct Info	ormation						
Product Type			HUMAN OTC DRUG Item Code (Sou			rce) NDC:67510-0673		
Route of Administration TOPICAL								
Active Ingredient/Active Moiety								
Ingredient Name Basis of St							rength	Strengt
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL FORM - UNII:L7T10EIP3A)					ECIFIED	D MENTHOL, 41 UNSPECIFIED FORM in		
Ir	active Ing	redients						
Ingredient Name								ength
ALOE VERA LEAF (UNII: ZY81Z83H0X)								J
ARCTIUM LAPPA ROOT (UNII: 597E9BI3Z3)								
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)								
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)								
CAMPHOR OIL (UNII: 75IZZ8Y727)								
GREEN TEA LEAF (UNII: W2ZU1RY8B0)								
CARBOMER 940 (UNII: 4Q93RCW27E)								
CENTELLA ASIATICA (UNII: 7M867G6T1U)								
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)								
ISOPROPYL ALCOHOL (UNII: ND2M416302)								
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F) FD&C BLUE NO. 1 (UNII: H3R47K3TBD)								
FD&C BLUE NO. 1 (UNII: H3R47K31BD) FD&C YELLOW NO. 5 (UNII: I753WB2F1M)								
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)								
TROLAMINE (UNII: 903K93S3TK)								
WATER (UNII: 059QF0KO0R)								
PHENOXYETHANOL (UNII: HIE492ZZ3T)								
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)								
Packaging								
#	Item Code	P	ackage Description		Mai	keting Star Date		eting End Date
1	NDC:67510- 0673-2	74 mL in 1 BOTT Combination Pro	LE, WTH APPLICATOR; Type duct	e 0: Not a	01/24	4/2023		
Marketing Information								
	Marketing Category	Applica	tion Number or Mono Citation	graph		ting Start Date		eting End Date
OTC monograph not final		not part348	part348		01/24/2023			

Labeler - KAREWAY PRODUCT, INC. (121840057)

Revised: 1/2023

KAREWAY PRODUCT, INC.