

NATURAL RELIEF 1222- menthol lotion
RENU LABORATORIES, INC.

Natural Relief 1222 Menthol

Menthol. . . . 1.25 percent

For external use only Avoid contact with eyes and sensitive areas. 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.

Stop use and ask a doctor if condition persists for more than 3 days or worsens.

Keep out of reach of children.

Topical Analgesic

Temporary relief of minor aches and pains associated with

- arthritis
- simple backache
- muscle strains
- sprains
- bruises

Directions

adults and children 12 years of age and older - apply liberally to affected area

- massage into painful area until thoroughly absorbed into skin
- repeat as necessary
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Store at 68 ° to 77° F.

Natural Relief 1222 HOMEOPATHIC JOINT & MUSCLE PAIN RELIEF

Allium Sativum (Garlic) Bulb Extract, Aloe Barbadensis (Aloe) Leaf Extract, Benzoic Acid, Collagen, Dehydroacetic Acid, Deionized Water, Emulsifying Wax NF, Fragrance, Glycine Soja (Soybean) Oil, HPUS Bryonia 10x, HPUS Rhus Toxicodendron 10x, Hydolyzed Elastin (Powder), Mineral Oil, Myristyl Myristate, Phenoxyethanol, Propylene Glycol, USP, Prunus Amygdalus Dulcis (Sweet Almond Oil), Squalane, Stearic Acid, Triethanolamine, Vitamin E dl Alpha Tocopherol

When using this product:

- Do not bandage tightly -
- avoid contact with eyes
- do not apply to wounds or damaged skin
- do not use heating pads or other heating devices

Keep out of reach of children

If swallowed, seek medical help or contact a poison control center right away

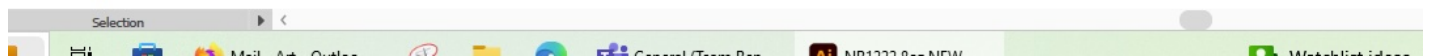
Natural Relief 1222

Soothing Joint & Muscle

Pain Relief

Natural Relief 1222 Label

LABEL ART



NATURAL RELIEF 1222			
menthol lotion			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76348-711
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	2.8 g in 224 g

Inactive Ingredients	
Ingredient Name	Strength
STEARIC ACID (UNII: 4ELV7Z65AP)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
GARLIC (UNII: V1V998DC17)	
TOXICODENDRON PUBESCENS LEAF (UNII: 6IO182RP7A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
MYRISTYL MYRISTATE (UNII: 4042ZC00DY)	
BRYONIA ALBA ROOT (UNII: T7J046YI2B)	
SQUALANE (UNII: GW89575KF9)	
WHITE WAX (UNII: 7G1J5DA97F)	
WATER (UNII: 059QF0KO0R)	
TROLAMINE (UNII: 9O3K93S3TK)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
HYDROLYZED BOVINE ELASTIN (BASE; 1000 MW) (UNII: ZR28QKN0WT)	
MARINE COLLAGEN, SOLUBLE (UNII: 8JC99XGU4W)	
ALMOND OIL (UNII: 66YXD4DKO9)	
SOYBEAN OIL (UNII: 241ATL177A)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
TETRASODIUM EDTATE DIHYDRATE (UNII: 3JGX4KKZ4A)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76348-711-08	224 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/22/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/22/2024	

Labeler - RENU LABORATORIES, INC. (945739449)

Establishment			
Name	Address	ID/FEI	Business Operations
Renu Laboratories, Inc.		945739449	manufacture(76348-711)

