RAPID PAIN-RELIEF TOPICAL- lidocaine hydrochloride, menthol cream NORDIC HEALTHY LIVING INC.

RAPID PAIN-RELIEF TOPICAL CREAM

Drug Facts

Active ingredients

Lidocaine HCL 4.00% Menthol 1.00%

Purpose

Topical Analgesic

Uses:

• For the temporary relief of pain.

Warnings:

not intended for ingestion. For external use only

Do not use

• in large quantities, particularly raw surfaces or blistered areas.

When using this product

Avoid contact with eyes

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.

Keep out of reach of children.

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

If pregnant or breast-feeding,

ask a health professional before use.

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 years of age: consult a doctor.

Other information:

• Store at 20-25°C (68-77°F) and protect from moisture.

Inactive ingredients:

Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Arnica Montana Flower Extract, Ascorbic Acid (Vitamin C), Boswellia Serrata Extract, Caprylic/Capric Triglyceride, Carbomer, Ceteareth-20, Cetearyl Alcohol, Cetyl Alcohol, Ethylhexylglycerin, Glycerin, Ilex Paraguariensis (Yerba Mate') Extract, Magnesium Sulfate, Methylsulfonylmethane (MSM), Methyl salicylate, Phenoxyethanol, Potassium Hydroxide.

Questions?

(800) 564-1408

Package Labeling: RAPID PAIN-RELIEF TOPICAL CREAM, 2oz/60ml





RAPID PAIN-RELIEF TOPICAL

lidocaine hydrochloride, menthol cream

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:82798-295

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	40 mg in 1 mL
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	10 mg in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
WATER (UNII: 059QF0KO0R)		
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)		
ASCORBIC ACID (UNII: PQ6CK8PD0R)		
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)		
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)		
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)		
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)		
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)		
CETYL ALCOHOL (UNII: 936JST6JCN)		
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)		
GLYCERIN (UNII: PDC6A3C0OX)		
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)		
MAGNESIUM SULFATE, UNSPECIFIED FORM (UNII: DE08037SAB)		
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)		
METHYL SALICYLATE (UNII: LAV5U5022Y)		
PHENOXYETHANOL (UNII: HIE492ZZ3T)		
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)		

F	Packaging					
#	tem Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:82798-295- 02	60 mL in 1 TUBE; Type 0: Not a Combination Product	07/01/2022			
2	NDC:82798-295- 06	175 mL in 1 TUBE; Type 0: Not a Combination Product	07/01/2022			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M017	07/01/2022		

Labeler - NORDIC HEALTHY LIVING INC. (118546272)