NERVIVE PAIN RELIEVING LIQUID ROLL-ON- lidocaine hcl and menthol liquid
The Procter & Gamble Manufacturing Company

NERVIVE™ PAIN RELIEVING Liquid Roll-on

Drug Facts

Active ingredients

Lidocaine HCl 4%

Menthol 1%

Active ingredients Purpose

Topical anesthetic

Topical analgesic

Use

temporarily relieves minor pain

Warnings

For external use only

Flammable • keep away from heat and open flame

Do not use

• If you have had an allergic reaction to lidocaine or other local anesthetics

• on large areas of the body or on cut, irritated, blistered, or swollen skin

• on puncture wounds

• for more than one week without consulting a doctor

When using this product

• use only as directed. Read and follow all directions and warnings on this carton.

• avoid contact with eyes and mucous membranes

• rare cases of serious burns have been reported with products of this type

• do not apply to wounds or damaged, broken or irritated skin

• do not bandage tightly or apply local heat (such as heating pads) or a medicated patch to the area of use

• a transient burning sensation may occur upon application but generally disappears in several days

• avoid applying into skin folds

Stop use and ask a doctor if

• condition worsens

• severe burning sensation, redness, rash or irritation develops

• symptoms persist for more than 7 days or clear up and occur again within a few days

• you experience signs of skin injury, such as pain, swelling, or blistering where the product was applied

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children and pets.

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

Directions

adults and children over 12 years:

• apply a thin layer to affected area every 6 to 8 hours
• do not exceed 3 applications in a 24 hour period
• massage into painful area until thoroughly absorbed into skin
AFTER APPLYING, WASH HANDS WITH SOAP AND WATER
children 12 years or younger:ask a doctor

Store at no greater than 25°C (77°F).

Inactive ingredients

acrylates/C10-30 alkyl acrylate crosspolymer, alcohol, aminomethylpropanol, C30-45 alkyl cetearyl dimethicone crosspolymer, caprylyl methicone, ceteth-20 phosphate, cetostearyl alcohol, dicetyl phosphate, dimethicone, edetate disodium, ethylhexylglycerin, glyceryl stearate, hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer, isohexadecane, phenoxyethanol, polyoxyl 15 hydroxystearate, polysorbate 60, sorbitan isostearate, steareth-21, tocopherol, water

Questions?

1-855-446-4345

DIST. BY: PROCTER & GAMBLE, CINCINNATI, OH 45202

NerviveHealth.com

PRINCIPAL DISPLAY PANEL - 2.5 OZ

NERVIVE™ PAIN RELIEVING LIQUID ROLL-ON

LIDOCAINE HCl & MENTHOL

TARGETS MULTIPLE NERVE PAIN RECEPTORS

MAXIMUM STRENGTH

MEDICATED ROLL-ON NET WT 2.5 OZ (70.9 g)

NERVIVE PAIN RELIEVING LIQUID ROLL-ON

lidocaine hcl and menthol liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69423-970
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	0.04 g in 1 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
CARBOMER INTERPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 132584PQMO)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
STEARETH-21 (UNII: 53J3F32P58)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%) (UNII: 86FQE96TZ4)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
CETETH-20 PHOSPHATE (UNII: 921FTA1500)	
DIMETHICONE 350 (UNII: 2Y53S6ATLU)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
ALCOHOL (UNII: 3K9958V90M)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
SORBITAN ISOSTEARATE (UNII: 01S2G2C1E4)	
CAPRYLYL TRIMETHICONE (UNII: H6HK6E4EB1)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
TOCOPHEROL (UNII: R0ZB2556P8)	
C30-45 ALKYL CETEARYL DIMETHICONE CROSSPOLYMER (UNII: 4ZK9VP326R)	
POLYOXYL 15 HYDROXYSTEARATE (UNII: 71YMM1X75O)	
ALUMINUM DICETYL PHOSPHATE (UNII: WMV3R5DS7O)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
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:69423-970-03	70.9 g in 1 CANISTER; Type 0: Not a Combination Product	02/08/2022	12/31/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	02/08/2022	12/31/2025

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 7/2024

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