

NERVIVE PAIN RELIEVING CREAM- lidocaine hcl and menthol cream
The Procter & Gamble Manufacturing Company

NERVIVE™ PAIN RELIEVING CREAM

Drug Facts

Active ingredients

Lidocaine HCl 4%

Menthol 1%

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

PRINCIPAL DISPLAY PANEL - 3.0 OZ

NERVIVE™

PAIN RELIEVING CREAM

LIDOCAINE HCl & MENTHOL

TARGETS MULTIPLE

NERVE PAIN RECEPTORS

PENETRATING CREAM

MEDICATED CREAM NET WT 3.0 OZ (85.1 g)



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QUESTIONS? 1-855-446-4345

from the global makers of
nerve care products for
over 50 years.

How to open:
1. Push down
2. Turn according to
the direction of arrow
on the top of cap

TAMPER EVIDENT.
DO NOT USE IF PRINTED
SEAL UNDER CAP IS
MISSING OR DAMAGED

SMALLER THAN
ACTUAL SIZE

**PAIN RELIEVING
CREAM**
LIDOCAINE HCl & MENTHOL
**TARGETS MULTIPLE
NERVE PAIN RECEPTORS**
PENETRATING CREAM

**NET WT
3.0 OZ (85.1 g)**

NERVIVE™
CREAM



Penetrates nerves &
relieves pain in toes,
feet, & legs



Penetrates nerves &
relieves pain in fingers,
hands, & arms

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



Patents: www.pg.com/patents



NERVIVE PAIN RELIEVING CREAM

lidocaine hcl and menthol cream

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:84126-318

Route of Administration	TOPICAL
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Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE ANHYDROUS (UNII: EC2CNF7XFP) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 g in 100 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1 g in 100 g

Inactive Ingredients	
Ingredient Name	Strength
CARBOMER INTERPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 132584PQMO)	
SORBITAN ISOSTEARATE (UNII: 01S2G2C1E4)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
TOCOPHEROL (UNII: R0ZB2556P8)	
C30-45 ALKYL CETEARYL DIMETHICONE CROSSPOLYMER (UNII: 4Z K9VP326R)	
POLYOXYL 15 HYDROXYSTEARATE (UNII: 71YMM1X75O)	
CAPRYLYL TRISILOXANE (UNII: Q95M2P1KJL)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0K00R)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
DIHEXADECYL PHOSPHATE (UNII: 2V6E5WN99N)	
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)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
ALCOHOL (UNII: 3K9958V90M)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	

Product Characteristics			
Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84126-318-03	1 in 1 CARTON	08/12/2024	

1	85.1 g in 1 TUBE; Type 0: Not a Combination Product		
Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	08/12/2024	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 10/2025

The Procter & Gamble Manufacturing Company