

**DERMATOOL NERVE CARE- lidocaine hcl, menthol cream**  
**Papalo Group, LLC**

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**Dermatool Nerve Care Cream**

Lidocaine HCl 4%, Menthol 1%

Topical Analgesic

For the temporary relief of pain.

**For external use only.**

**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** only use 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** the condition worsens, 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.

Ask a health professional before use.

In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

**Adults and children 12 years and older:** apply to the affected area, not more than 6 to 8 hours. Do not exceed 4 applications in a 24-hour period. Massage into painful area until thoroughly absorbed into the skin. **AFTER APPLYING WASH HANDS WITH SOAP AND WATER.** **Children under 12 years of age:** ask a doctor.

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf Juice, Aminomethyl Propanol, C30-45 Alkyl Cetearyl Dimethicone Crosspolymer, Caprylyl Methicone, Cetearyl Alcohol, Ceteth-20 Phosphate, Dicyetyl Phosphate, Dimethicone, Disodium EDTA, Ethylhexylglycerin, Glyceryl Stearate, Hydroxyethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Isohexadecane, Methylparaben, Polysorbate 60, SD Alcohol 40 (15%), Steareth-21, Water.

NDC# 0000-0000-00

**Drug Facts**

**Active ingredients**  
 Lidocaine HCl 4%  
 Menthol 1%

**Purpose**  
 Topical anesthetic  
 Topical analgesic

**Use** For the temporary relief of pain.

**Warnings**

For external use only.

**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**

- only use 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 breastfeeding, 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 over 12 years:**

- apply thin layer to affected area every 6 to 8 hours
- do not exceed 4 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.

**Other information** Store at 15°-30° C (59°-86° F).

**Inactive ingredients** acrylates/C10-30 alkyl acrylate crosspolymer, aloe barbadensis leaf juice, aminomethyl propanol, C30-45 alkyl cetearyl dimethicone crosspolymer, caprylyl methicone, cetearyl alcohol, ceteth-20 phosphate, dicetyl phosphate, dimethicone, disodium EDTA, ethylhexylglycerin, glyceryl stearate, hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer, isohexadecane, methylparaben, polysorbate 60, SD alcohol 40 (15%), steareth-21, water

Child-resistant packaging. Replace cap after each use.

❓ **QUESTIONS?**

support@dermatoolcare.com



📍 **DISTRIBUTED BY:** Dermatool®  
 Jersey City, NJ 07307

**MADE IN THE U.S.A.**  
 with domestic &  
 imported components

🌐 www.dermatoolcare.com

Helps Calm Nerve  
 Pain Signals



Nerve Relief

Application Areas



Feet, Hands, Legs,  
 and Arms

Deep-Absorbing  
 Cream



Localized Topical Relief  
 for Affected Areas

DERMATOOL®

Nerve  
 Care Cream

RELIEF FOR PAIN  
 & DISCOMFORT

TARGETS NERVE PAIN SIGNALS

LIDOCAINE HCl & MENTHOL

Net Wt 5.5 oz (156 g)

**HOW TO USE**

Apply a thin layer to affected areas.

Massage gently until absorbed.

Apply a small amount first to sensitive skin.

For external use only.

**WHAT TO EXPECT**

Provides topical relief for nerve-related pain and discomfort.

Response may vary by person and area of use.

**FOR REGULAR USE**

Intended for repeat application as part of a nerve care routine.

**IMPORTANT**

Not intended to replace medical care or prescription treatment.

**DERMATOOL NERVE CARE**

lidocaine hcl, menthol cream

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:84728-359
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>MENTHOL</b> (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1 g in 100 g
<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 g in 100 g

**Inactive Ingredients**

Ingredient Name	Strength
ISOHEXADECANE (UNII: 918X1OUF1E)	
CETETH-20 PHOSPHATE (UNII: 921FTA1500)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%) (UNII: 86FQE96TZ4)	
STEARETH-21 (UNII: 53J3F32P58)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
CARBOMER COPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 809Y72KV36)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
DIHEXADECYL PHOSPHATE (UNII: 2V6E5VN99N)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
CAPRYLYL TRISILOXANE (UNII: Q95M2P1KJL)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
C30-45 ALKYL CETEARYL DIMETHICONE CROSSPOLYMER (UNII: 4ZK9VP326R)	
DIMETHICONE 200 (UNII: RGS4T2AS00)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84728-359-05	1 in 1 CARTON	05/01/2026	
1		159 g in 1 BOTTLE; 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	05/01/2026	

**Labeler** - Papalo Group, LLC (119324672)

**Registrant** - Derma Care Research Labs, LLC (116817470)

### Establishment

Name	Address	ID/FEI	Business Operations
Derma Care Research Labs, LLC		116817470	manufacture(84728-359)