## EXTRA STRENGTH NATURALLY HL WARMING ICE- lidocaine hcl gel ridge properties

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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Directions: Test skin to sensitivity prior to procedure. Apply generously to skin prior to and/or during procedure as needed for pain. Discontinue use if sensitivity occures. Not for use on face.

Warning - Keep out of reach of children - For external use only - Avoid contact with the eyes

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**Purpose:**Topical Anesthetic

**Uses:** Temporarily relieves pain

# Active ingredients: Lidocaine HCL

4%

# **Inactive Ingredients: Aloe Vera, Flax** Seed, Oils Of Grape seed, Olive, Black Pepper, Wintergreen, & Eucalyptus

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FDA Registered NDC # 69804-038-15

Active ingredients: Lidocaine HCL Purpose:

**Topical Anesthetic** 

Warning - Keep out of reach of children - For external use only Avoid contact with the eyes

#### Other information:

This product was manufactured by Pain Relief Naturally. For contact info please visit

WWW.NATURALLYHL.COM

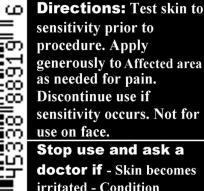
#### **Drug Facts:**

For professional use only

**Uses:** Temporarily relieves pain



**Inactive Ingredients: Aloe Vera, Flax** Seed, Oils Of Grape seed, Olive, Black Pepper, Wintergreen, & Eucalyptus



Stop use and ask a doctor if - Skin becomes irritated - Condition

worsens or symptoms last longer than 7 days -Symptoms clear up then reoccur within a few days

**Do Not Use** if you have any known allergy to any of the ingredients in this product. Discontinue use and seek medical attention should any occur

Shake well before each use

lidocaine hcl gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69804-038	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
LIDO CAINE HYDRO CHLO RIDE (UNII: V13007Z41A) (LIDO CAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	40 mg in 1000 mg	

Inactive Ingredients			
Ingredient Name	Strength		
ALOE VERA WHOLE (UNII: KIZ4X2EHYX)	540 mg in 1000 mg		
EUCALYPTUS OIL (UNII: 2R04ONI662)	40 mg in 1000 mg		
BLACK PEPPER OIL (UNII: U17J84S19Z)	40 mg in 1000 mg		
GRAPE SEED OIL (UNII: 930 MLC8 XGG)	120 mg in 1000 mg		
OLIVE OIL (UNII: 6 UYK2W1W1E)	120 mg in 1000 mg		
FLAX SEED (UNII: 4110 YT348 C)	100 mg in 1000 mg		

Packaging				
# Iten	ı Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:6		200 mg in 1 BOTTLE, DISPENSING; Type 0: Not a Combination oduct	02/09/2017	
2 NDC:6		500 mg in 1 BOTTLE, DISPENSING; Type 0: Not a Combination oduct	02/09/2017	
3 NDC:6	9804-038- Pro	5700 mg in 1 BOTTLE, DISPENSING; Type 0: Not a Combination oduct	02/09/2017	
4 NDC:6		3400 mg in 1 BOTTLE, DISPENSING; Type 0: Not a Combination oduct	02/09/2017	
16	Pro	oduct	32,33,231.	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	02/09/2017	

### Labeler - ridge properties (029478762)

Establishment				
Name	Address	ID/FEI	Business Operations	
ridge properties		029478762	manufacture(69804-038)	

Revised: 2/2017 ridge properties