# EXTRA STRENGTH NATURALLY HL SUNBURN RELIEF- lidocaine hcl liquid 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 Affected area as needed for pain. Discontinue use if sensitivity occurs. 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:

Water, Witch Hazel, Kava kava, Organic Alcohol, Yarrow, Nutmeg, Copaiba Balsam

### FDA Registered NDC # 69804-040-01

#### **Drug Facts:**

For professional use only

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Lidocaine HCL 4%
Purpose:
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Uses: Temporarily relieves pain

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2 Oz Spray



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

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

Inactive Ingredients: Water, Witch Hazel, Kava kava, Organic Alcohol, Yarrow, Nutmeg, Copaiba Balsam

#### Other information:

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

#### EXTRA STRENGTH NATURALLY HL SUNBURN RELIEF

lidocaine hcl liquid

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69804-040

Route of Administration TOPICAL

#### **Active Ingredient/Active Moiety**

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

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P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:69804-040- 08	14200 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	0 2/10/20 17				
2	NDC:69804-040- 07	28500 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	0 2/10/20 17				
3	NDC:69804-040- 04	113400 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	0 2/10/20 17				
4	NDC:69804-040- 01	56700 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	02/10/2017				

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part348	02/10/2017				

## **Labeler -** ridge properties (029478762)

Establishment						
Name	Address	ID/FEI	Business Operations			
ridge properties		029478762	manufacture(69804-040)			

Revised: 2/2017 ridge properties