EXTRA STRENGTH NATURALLY HL HAND CARE- lidocaine hcl spray 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.

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 # 69804-031-01

Drug Facts:

For professional use only

Active ingredients:
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 HAND CARE

lidocaine hcl spray

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69804-031

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength LIDO CAINE HYDRO CHLO RIDE (UNII: V13007Z41A) (LIDO CAINE UNII:98 PI200987) LIDO CAINE HYDRO CHLO RIDE ANHYDRO US in 1000 mg

Packaging							
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:69804-031- 07	28500 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	0 1/25/20 17				
2	NDC:69804-031- 08	14200 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	0 1/25/20 17				
3	NDC:69804-031- 01	56700 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	0 1/25/20 17				
4	NDC:69804-031- 04	113400 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	0 1/25/20 17				

Marketing Information								
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date					
OTC monograph not final	part348	0 1/25/20 17						

Labeler - ridge properties (029478762)

Establishment							
Name	Address	ID/FEI	Business Operations				
ridge properties		029478762	manufacture(69804-031)				

Revised: 1/2017 ridge properties