

**EXTRA STRENGTH SKIN REPAIR- 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

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FDA Registered
NDC # 69804-049-01

Drug Facts:
 For professional use only

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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

Shake Well Before Each Use

EXTRA STRENGTH SKIN REPAIR

lidocaine hcl spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69804-049
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98P200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	40 mg in 1000 mg

Inactive Ingredients

Ingredient Name	Strength
PIPER METHYSTICUM WHOLE (UNII: 3P306S300W)	200 mg in 1000 mg
ACHILLEA MILLEFOLIUM OIL (UNII: 97P5D0WG43)	75 mg in 1000 mg
WITCH HAZEL (UNII: 101I4J0U34)	510 mg in 1000 mg
COPAIBA OIL (UNII: 64VX45Y68N)	100 mg in 1000 mg
NUTMEG OIL (UNII: Z1CLM48948)	75 mg in 1000 mg

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69804-049-08	14200 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/03/2017	
2	NDC:69804-049-07	28500 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/03/2017	
3	NDC:69804-049-01	56700 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/03/2017	
4	NDC:69804-049-04	113400 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/03/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	04/03/2017	

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

Name	Address	ID/FEI	Business Operations
ridge properties		029478762	manufacture(69804-049)

Revised: 3/2017

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