

**EXTRA STRENGTH SCAR PREVENTION- 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 # 69804-064-01

Drug Facts:
 For professional use only

Active ingredients:
 Lidocaine HCL 4%
Purpose:
 Topical Anesthetic

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

Shake Well Before Each Use

EXTRA STRENGTH SCAR PREVENTION

lidocaine hcl spray

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:69804-064 |
| 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 |
|---|-------------------|
| WITCH HAZEL (UNII: 10114J0U34) | 510 mg in 1000 mg |
| PIPER METHYSTICUM WHOLE (UNII: 3P306S300W) | 200 mg in 1000 mg |
| ACHILLEA MILLEFOLIUM OIL (UNII: 97P5D0WG43) | 75 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:69804-064-08 | 14200 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 05/01/2017 | |
| 2 | NDC:69804-064-07 | 28500 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 05/01/2017 | |
| 3 | NDC:69804-064-01 | 56700 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 05/01/2017 | |
| 4 | NDC:69804-064-04 | 113400 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 05/01/2017 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|-------------------------|--|----------------------|--------------------|
| OTC monograph not final | part348 | 05/01/2017 | |

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

| Name | Address | ID/FEI | Business Operations |
|------------------|---------|-----------|------------------------|
| ridge properties | | 029478762 | manufacture(69804-064) |

Revised: 4/2017

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