

**EXTRA STRENGTH LIQUID BURN RELIEF GEL- 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.

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 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: Aloe Vera, Witch Hazel, Organic Alcohol, Kava Kava, Yarrow, Nutmeg, Copaiba Balsam, Flax Seed Extract

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FDA Registered NDC # 69804-026-15	Drug Facts: For professional use only		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.
Active ingredients: Lidocaine HCL 4%	Uses: Temporarily relieves pain		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
Purpose: Topical Anesthetic		2 Oz Liquid Gel	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
Warning - Keep out of reach of children - For external use only Avoid contact with the eyes			Inactive Ingredients: Aloe Vera, Witch Hazel, Organic Alcohol, Kava Kava, Yarrow, Nutmeg, Copaiba Balsam, Flax Seed Extract

EXTRA STRENGTH LIQUID BURN RELIEF GEL

lidocaine hcl gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69804-026
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
COPAIBA OIL (UNII: 64VX45Y68N)	75 mg in 1000 mg
ALOE VERA WHOLE (UNII: KIZ4X2EHYX)	310 mg in 1000 mg
WITCH HAZEL (UNII: 101I4J0U34)	200 mg in 1000 mg
NUTMEG OIL (UNII: Z1CLM48948)	75 mg in 1000 mg
ACHILLEA MILLEFOLIUM OIL (UNII: 97P5D0WG43)	75 mg in 1000 mg
PIPER METHYSTICUM WHOLE (UNII: 3P306S300W)	150 mg in 1000 mg
FLAX SEED (UNII: 4110YT348C)	75 mg in 1000 mg

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69804-026-13	14200 mg in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	01/13/2017	
2	NDC:69804-026-14	28500 mg in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	01/13/2017	
3	NDC:69804-026-15	56700 mg in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	01/13/2017	
4	NDC:69804-026-16	113400 mg in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	01/13/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	01/13/2017	

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

Name	Address	ID/FEI	Business Operations
ridge properties		029478762	manufacture(69804-026)