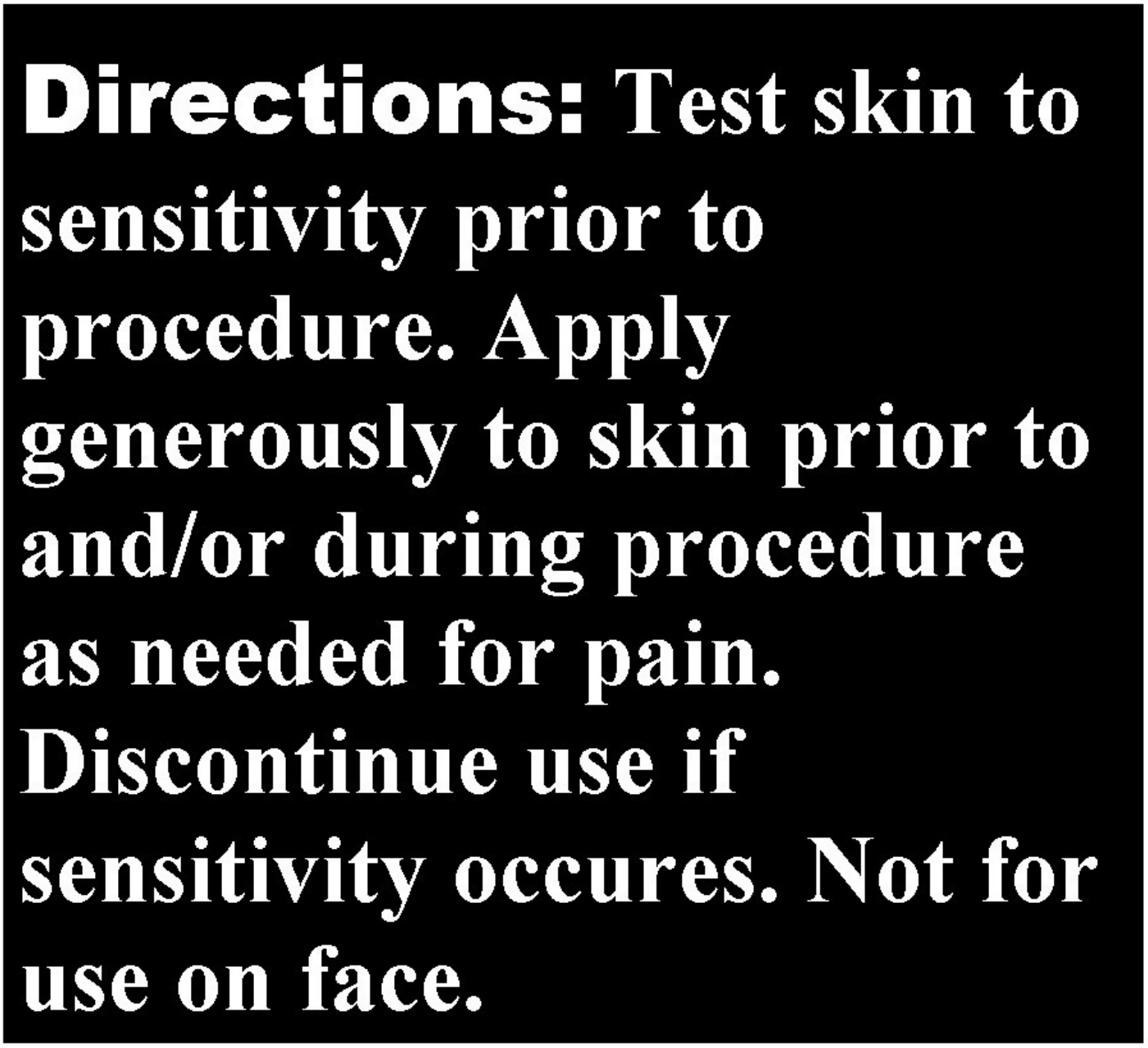
EXTRA STRENGTH LIQUID BEDSORE RELIEF GEL- lidocaine hcl gel
Ridge Properties

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.



Warning - Keep out of reach of children - For external use only - Avoid contact with the eyes

Warning - Keep out of reach of children - For external use only - Avoid contact with the eyes

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

Need some more? Pick some up at WW.NATURALLYHL.COM

FDA Registered
NDC # 69804-022-15

Active ingredients:
Lidocaine HCL 4%
Purpose:
Topical Anesthetic

Warning - Keep out of reach of children - For external use only Avoid contact with the eyes

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

Drug Facts:
For professional use only

Uses: Temporarily relieves pain

Extra Strength
Naturally
HL
Liquid Numbing Gel With Added Lidocaine

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

7 37278 91769 2

2 Oz
Liquid Gel

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.

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

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

Shake well before each use

EXTRA STRENGTH LIQUID BEDSORE RELIEF GEL

lidocaine hcl gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69804-023
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**
ALOE VERA WHOLE (UNII: KIZ4X2EHYX)	310 mg in 1000 mg
WITCH HAZEL (UNII: 101I4J0U34)	200 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
COPAIBA OIL (UNII: 64VX45Y68N)	75 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:69804-023-13	14200 mg in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	01/13/2017	
2	NDC:69804-023-14	28500 mg in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	01/13/2017	
3	NDC:69804-023-15	56700 mg in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	01/13/2017	
4	NDC:69804-023-16	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(69804-023)