EXTRA STRENGTH NATURALLY HL ITCH RELIEF- lidocaine hcl cream 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.

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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, Propolis, Yarrow, Nutmeg, Copaiba Balsam, BTMS 50 (A natural extract of the Colza Seed)

Topical Anesthetic





This product was manufactured by Pain Relief Naturally. For

Water, Witch Hazel

Organic Alcohol, Kav Kava, Yarrow, Nutmeg Propolis, (BTMS 50 -A Natural Extract of The

EXTRA STRENGTH NATURALLY HL ITCH RELIEF

lidocaine hcl cream

Product Information

HUMAN OTC DRUG NDC:69804-033 Product Type Item Code (Source)

Route	οf	Αd	ministration	ı
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TOPICAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
LIDO CAINE HYDRO CHLO RIDE (UNII: V13007Z41A) (LIDO CAINE - UNII:98 PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	40 mg in 1000 mg		

Inactive Ingredients			
Ingredient Name	Strength		
PROPOLIS WAX (UNII: 6 Y8 XYV2NOF)	40 mg in 1000 mg		
COPAIBA OIL (UNII: 64VX45Y68N)	90 mg in 1000 mg		
NUTMEG OIL (UNII: Z1CLM48948)	60 mg in 1000 mg		
BEHENTRIMO NIUM METHO SULFATE (UNII: 5SHP745C61)	120 mg in 1000 mg		
PIPER METHYSTICUM WHO LE (UNII: 3P306S300W)	190 mg in 1000 mg		
WITCH HAZEL (UNII: 10 1I4J0 U34)	400 mg in 1000 mg		
ACHILLEA MILLEFOLIUM O IL (UNII: 97P5D0WG43)	60 mg in 1000 mg		

I	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:69804-033-06	14200 mg in 1 JAR; Type 0: Not a Combination Product	0 2/0 1/20 17		
2	NDC:69804-033-07	28500 mg in 1 JAR; Type 0: Not a Combination Product	0 2/0 1/20 17		
3	NDC:69804-033-02	56700 mg in 1 JAR; Type 0: Not a Combination Product	0 2/0 1/20 17		
4	NDC:69804-033-03	113400 mg in 1 JAR; Type 0: Not a Combination Product	0 2/0 1/20 17		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part348	02/01/2017		

Labeler - ridge properties (029478762)

Establishment					
Name	Address	ID/FEI	Business Operations		
ridge properties		029478762	manufacture(69804-033)		

Revised: 1/2017 ridge properties