

EXTRA STRENGTH POSTPARTUM RELIEF- lidocaine hcl spray
ridge properteis

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 # 69804-043-01

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 POSTPARTUM RELIEF

lidocaine hcl spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69804-043
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:69804-043-08	14200 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/03/2017	
2	NDC:69804-043-07	28500 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/03/2017	
3	NDC:69804-043-01	56700 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/03/2017	
4	NDC:69804-043-04	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 properteis (029478762)**Establishment**

Name	Address	ID/FEI	Business Operations
ridge properties		029478762	manufacture(69804-043)

Revised: 3/2017

ridge properteis