

RR ROLL-ON FOR PAIN- lidocaine hcl gel

Revolutionary Relief dba Medical XLabs

Active Ingredient

Lidocaine HCL 4%

Purpose

Topical Analgesic

Uses

Temporarily relieves pain and itching

Warnings

- For external use only Use only as directed
- Do not allow contact with the eyes and mucous membranes.
- Do not apply to wounds or damaged skin
- Do not use at the same time as other topical analgesics
- **Stop use and ask a doctor if:** allergic reaction occurs
- condition worsen or does not improve within 7 days
- redness, irritation, swelling, pain or other symptoms begin or increase.
- **If pregnant or Breastfeeding**

Ask a health professional before use.

Keep out of reach of children

If swallowed, get medical help or Contact a Poison Control right away. Emergency number : 1-800-222-1222

Direction

Adults and children 12 years of age or older:

- Apply to affected area not more than 3 to 4 times daily
- Children under two years of age, consult a physician

Additional information

Store in a cool, dry place

Inactive ingredients

Aqua, Alcohol Denat., Butylene Glycol, Steareth-21, Steareth-2, Cetyl Alcohol, Hemp

Seed Oil, Glycerin, Caprylyl Glycol, Kannopia-Active™ Proprietary Hemp Extract, Tocopherol, Arnica Montana Flower Extract, Boswellia Serrata (Frankincense) Extract, Plantago Major (Plantain) Leaf Extract, Phenoxyethanol, Hexelene Glycol, Xanthan Gum, Hexyl Cinnamic Aldehyde, Coumarin, Limonene, Peppermint

Product label

Drug Facts	
Active Ingredient: Lidocaine HCL 4%	Purpose: Topical Analgesic
Uses: Temporarily relieves pain and itching.	
Warnings: <ul style="list-style-type: none"> For external use only. Use only as directed. Do not allow contact with the eyes and mucous membranes. Do not apply to wounds or damaged skin. Do not use at the same time as other topical analgesics. Stop use and ask a doctor if: allergic reaction occurs - condition worsens or does not improve within 7 days - redness, irritation, swelling, pain or other symptoms begin or increase 	
IF PREGNANT OR BREASTFEEDING: Ask a health professional before use.	
KEEP OUT OF REACH OF CHILDREN. If swallowed, get medical help or contact a Poison Control right away. Emergency Number: 1-800-222-1222	
Directions: <ul style="list-style-type: none"> Adults and children 12 years of age or older: Apply to affected area not more than 3 to 4 times daily. Children under two years of age, consult a physician. 	
Additional Information: Store in a cool, dry place.	
Inactive Ingredients: Aqua, Alcohol Denat., Butylene Glycol, Steareth-21, Steareth-2, Cetyl Alcohol, Hemp Seed Oil, Glycerin, Caprylyl Glycol, Kannopia-Active™ Proprietary Hemp Extract, Tocopherol, Arnica Montana Flower Extract, Boswellia Serrata (Frankincense) Extract, Plantago Major (Plantain) Leaf Extract, Phenoxyethanol, Hexelene Glycol, Xanthan Gum, Hexyl Cinnamic Aldehyde, Coumarin, Limonene, Peppermint.	



REVOLUTIONARY
RELIEF™

from outside

**ROLL ON
FOR PAIN**

**300mg
Kannopia-Active™**
2 fl oz / 60mL



Manufactured For:
Medical XLabs
333 Las Olas Way, Suite 100
Ft. Lauderdale, FL 33301
1 (800) 757-4137



WWW.REVRELIEF.COM

RR ROLL-ON FOR PAIN

lidocaine hcl gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83048-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z 41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
STEARETH-21 (UNII: 53J3F32P58)	
STEARETH-2 (UNII: V56DFE46J5)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
GLYCERIN (UNII: PDC6A3C0OX)	

CAPRYLYL GLYCOL (UNII: 00YIU5438U)
HEMP (UNII: TD1MUT01Q7)
TOCOPHEROL (UNII: R0ZB2556P8)
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)
PLANTAGO MAJOR LEAF (UNII: 7DC28K241X)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
HEXYLENE GLYCOL (UNII: KEH0A3F75J)
XANTHAN GUM (UNII: TTV12P4NEE)
.ALPHA.-HEXYLCINNAMALDEHYDE (UNII: 7X6O37OK2I)
COUMARIN (UNII: A4VZ22K1WT)
LIMONENE, (+)- (UNII: GFD7C86Q1W)
PEPPERMINT (UNII: V95R5KMY2B)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83048-001-01	60 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	11/03/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	11/03/2022	

Labeler - Revolutionary Relief dba Medical XLabs (118810950)

Revised: 11/2024

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