

LIDONEX- lidocaine hcl liquid
Newpharma, Inc.

Active Ingredient

Lidocaine HCL, USP 4%

Purpose

Topical Anesthetic

Use

For temporarily relief of pain

Warnings

For external use only

Flammable: Keep away from fire or flame

Do not use

- on wounds or damaged skin
- in large quantities
- with a heating pad
- if you are allergic to any ingredients of this product

When using this product

- use only as directed.
- avoid contact with the eyes, mucous membranes or rashes
- do not bandage tightly

Stop use and ask doctor if

- skin reactions occur, such as rash, itching, redness, irritation, pain, swelling and blistering
- condition worsens
- symptoms persist for more than 7 days
- symptoms clear up and occur again within a few days

If pregnant or breast-feeding. Ask a health professional before use.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Adults and children over 12 years of age and over

- clean and dry affected area
- apply to affected area not more than 3 to 4 times daily

Children under 12 years of age: Consult a doctor

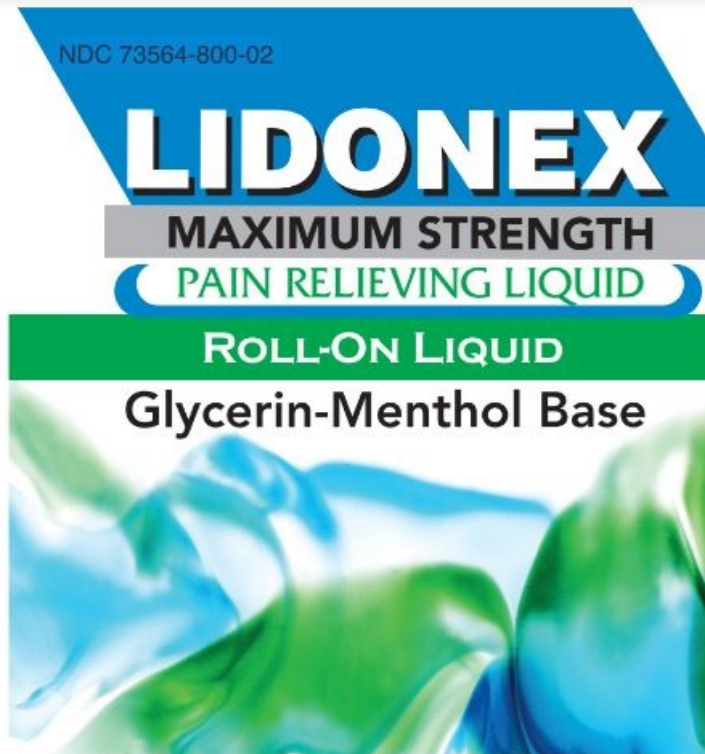
Inactive ingredients

glycerin, isopropyl alcohol, menthol, methylparaben, propylparaben, purified water,

xanthan gum

- avoid storing product in direct sunlight
- protect product from excessive moisture
- store with lid closed tightly

Questions or comments? (305) 592 - 9216



NDC 73564-800-02

LIDONEX

MAXIMUM STRENGTH

PAIN RELIEVING LIQUID

ROLL-ON LIQUID

Glycerin-Menthol Base

Contents: 2 fl. oz (60mL)

LOT#/Exp. Date:



DIST. BY: NEWPHARMA, INC. MIAMI, FL 33122

DRUG FACTS	
Active Ingredients Lidocaine HCl, USP 4%.....	Purpose Topical Anesthetic
Uses For temporary relief of pain	
Warnings For external use only	
Flammable: Keep away from fire or flame	
Do not use <ul style="list-style-type: none"> ■ on wounds or damaged skin ■ in large quantities ■ with a heating pad ■ if you are allergic to any ingredients of this product 	
When using this product <ul style="list-style-type: none"> ■ use only as directed ■ avoid contact with the eyes, mucous membranes or rashes ■ do not use bandage tightly 	
Stop use and ask a doctor if <ul style="list-style-type: none"> ■ skin reactions occur, such as rash, itching, redness, irritation, pain, swelling and blistering ■ condition worsens ■ symptoms persist for more than 7 days ■ symptoms clear up and occur again within a few days 	
If pregnant or breast-feeding. Ask a health professional before use. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	
Directions Adults and children 12 years of age and over <ul style="list-style-type: none"> ■ clean and dry affected area ■ apply to affected area not more than 3 to 4 times daily 	
Children under 12 years of age: Consult a doctor	
Other information <ul style="list-style-type: none"> ■ Avoid storing product in direct sunlight ■ Protect product from excessive moisture ■ Store with lid closed tightly 	
Inactive Ingredients glycerin, isopropyl alcohol, menthol, methylparaben, propylparaben, purified water, xanthan gum	
Questions or comments? (305)592-9216	

CODE L-003

Rev: 09/22

LIDONEX

lidocaine hcl liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73564-800
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
MENTHOL (UNII: L7T10EIP3A)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73564-800-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/04/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	10/04/2022	

Labeler - Newpharma, Inc. (019170437)

Revised: 2/2024

Newpharma, Inc.