

**ICEQUAKE LIDOCAINE ANALGESIC- lidocaine hydrochloride liquid**  
**Southern Sales & Service, Inc.**

-----  
**IceQuake Lidocaine Analgesic**

**Drug Facts**

**Active ingredient**

Lidocaine HCl 4%

**Purpose**

Topical anesthetic

**Use**

Temporarily relieves minor pain

**Warnings**

For external use only

**Do not use**

- on large areas of the body or on cut, irritated or swollen skin
- on puncture wounds
- for more than one week without consulting a doctor

**When using this product**

- use only as directed. Read and follow all directions and warnings on this label.
- do not allow contact with the eyes
- do not bandage or apply local heat (such as heating pads) to area of use

**Do not use**

- on large areas of the body or on cut, irritated or swollen skin
- on puncture wounds
- for more than one week without consulting a doctor

**condition worsens Stop use and ask a doctor if**

- symptoms persist for more than 7 days or clear up and occur again within a few days

**Flammable**

keep away from fire or flame

**If pregnant or breast-feeding**

ask a health professional before use.

**Keep out of reach of children**

and pets. If swallowed, get medical help or contact a Poison Control Center right away.

**Directions**

- adults and children over 12 years:

apply a thin layer to affected area every 6 to 8 hours, not to exceed 3 applications in a 24 hour period

children 12 years or younger: ask a doctor

**Inactive ingredients**

Disodium EDTA, isopropyl alcohol, glycerin, methylparaben, polysorbate 20, propylparaben, purified water, xanthan gum.

**IceQuake Lidocaine Topical Analgesic Liquid 74ml**

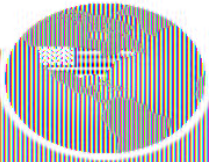
SHAKE  
WELL



Maximum Strength  
4%



Roll



DISTRIBUTED BY:  
Ice Quake Sales Service, Inc.  
10000 W. 11th Ave., Suite 100  
Denver, CO 80202  
Questions or Comments?  
Call 800.279.9570  
[icequakeonline.com](http://icequakeonline.com)

**Lidocaine**  
Topical Analgesic Liquid

2.5 fl.oz. (74 ml)

INDICATED FOR  
DRUG FREE  
ACTIVE INGREDIENT  
Lidocaine  
DRUG FREE  
LIFT LAE

<b>Drug Facts</b>	
<b>Active ingredient</b> Lidocaine HCl 4%	<b>Purpose</b> Topical anesthetic
Use Temporarily relieves minor pain	
<b>Warnings</b> For external use only	
<b>Do not use</b>	
<ul style="list-style-type: none"> <li>■ on large areas of the body or on cut, irritated or swollen skin</li> <li>■ on puncture wounds</li> <li>■ for more than one week without consulting a doctor</li> </ul>	
<b>When using this product</b>	
<ul style="list-style-type: none"> <li>■ use only as directed. Read and follow all directions and warnings on this label.</li> <li>■ do not allow contact with the eyes</li> <li>■ do not bandage or apply local heat (such as heating pads) to area of use</li> </ul>	
<b>Stop use and ask a doctor if</b>	
<ul style="list-style-type: none"> <li>■ condition worsens</li> <li>■ symptoms persist for more than 7 days or clear up and occur again within a few days</li> </ul>	
Flammable	
<ul style="list-style-type: none"> <li>■ keep away from fire or flame</li> </ul>	
<b>If pregnant or breast-feeding</b> ask a health professional before use. Keep out of reach of children and pets. If swallowed, get medical help or contact a Poison Control Center right away.	
<b>Directions</b>	

## ICEQUAKE LIDOCAINE ANALGESIC

lidocaine hydrochloride liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69822-430
<b>Route of Administration</b>	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
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	

<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>POLYSORBATE 20</b> (UNII: 7T1F30V5YH)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	

<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69822-430-03	74 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2022	

<b>Marketing Information</b>			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	08/01/2022	

**Labeler** - Southern Sales & Service, Inc. (013114906)

**Registrant** - Southern Sales & Service, Inc. (013114906)

Revised: 11/2023

Southern Sales & Service, Inc.