

**ALOCANE BOTANICALS- lidocaine hydrochloride spray**  
**Quest Products, LLC.**

-----  
**Alocane<sup>®</sup> Botanicals**

**Drug Facts**

**Active Ingredient**

Lidocaine HCL 4%

**Purpose**

Topical Pain Relief

**Uses**

For the temporary relief of pain and itching associated with:

- Sunburn
- Minor Skin Irritation
- Minor Burns
- Minor Cuts & Scrapes
- Insect Bites

**Warnings**

**For external use only.** Avoid contact with eyes.

**Do not use:**

- in large quantities, or over large areas of the body
- Over raw surfaces or blistered areas
- If you have an allergy to any of the ingredients

**Ask a doctor before use** if you have a severe burn or deep cut, broken or compromised skin

**Stop use and ask doctor:**

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

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

**If pregnant or breast-feeding,** ask a health care professional before use.

**Directions:**

**Adults and Children 2 years of age or older:**

Shake well. Apply top affected area not more than 3-4 times daily. To apply to face, spray on palm and carefully apply.

**Children under 2 years of age,** ask a doctor before use.

**Other Information**

Store at 15-30°C (59-86°F)

**Inactive Ingredients**

Water, Phenoxyethanol, Aloe Vera Leaf Juice, Lactobacillus Ferment, Echinacea Purpurea Root Extract, Calendula Officinalis Flower Extract, Sodium Hyaluronate, D-Panthenol, Glycerin, Hydroxyethyl Cellulose, Mannitol, Sodium Hydroxide

**PRINCIPAL DISPLAY PANEL -- Alocane Botanicals 4oz Spray Bottle**



lidocaine hydrochloride spray

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:68229-405
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

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

### Inactive Ingredients

Ingredient Name	Strength
<b>MANNITOL</b> (UNII: 3OWL53L36A)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	
<b>DEXPANTHENOL</b> (UNII: 1O6C93RI7Z)	
<b>LIMOSILACTOBACILLUS FERMENTUM</b> (UNII: 2C1F12C6AP)	
<b>HYALURONATE SODIUM</b> (UNII: YSE9PPT4TH)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>ECHINACEA PURPUREA ROOT</b> (UNII: OS64WTR4KU)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>HYDROXYETHYL CELLULOSE (5500 MPA.S AT 2%)</b> (UNII: M825OX60H9)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>CALENDULA OFFICINALIS FLOWER</b> (UNII: P0M7O4Y7YD)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68229-405-01	118 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/11/2023	
2	NDC:68229-405-02	104 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/11/2023	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	08/11/2023	

**Labeler** - Quest Products, LLC. (075402441)

**Establishment**

Name	Address	ID/FEI	Business Operations
Fill Tech USA		926433855	manufacture(68229-405)

Revised: 11/2024

Quest Products, LLC.