

**MAXOCAINE PAIN RELIEVING ROLL-ON- lidocaine hydrochloride,  
menthol liquid  
Prime Commerce, LLC**

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**MAXOCAINE PAIN RELIEVING ROLL-ON**

***Drug Facts***

***Active Ingredient***

Lydocaine HCL 4%

Menthol 1%

***Purpose***

Topical anesthetic

***Use:***

For the temporary relief of pain and itching associated with minor burns, sunburn, minor cuts, scrapes, insect bites or minor skin irritations.

***Warnings:***

- **For external use only**
- Avoid contact with eyes.
- If symptoms persist for more than seven days, discontinue use and consult physician.

**Keep out of reach of children.**

If swallowed, consult physician.

**Do not apply**

- to wounds or damaged skin.
- Do not bandage tightly.

**If pregnant or breast feeding,**

contact physician prior to use.

***Directions:***

Apply to affected area not more than three to four times daily. consult a physician.

**Adults and children two-years of age or older:Children under two-years of age:**

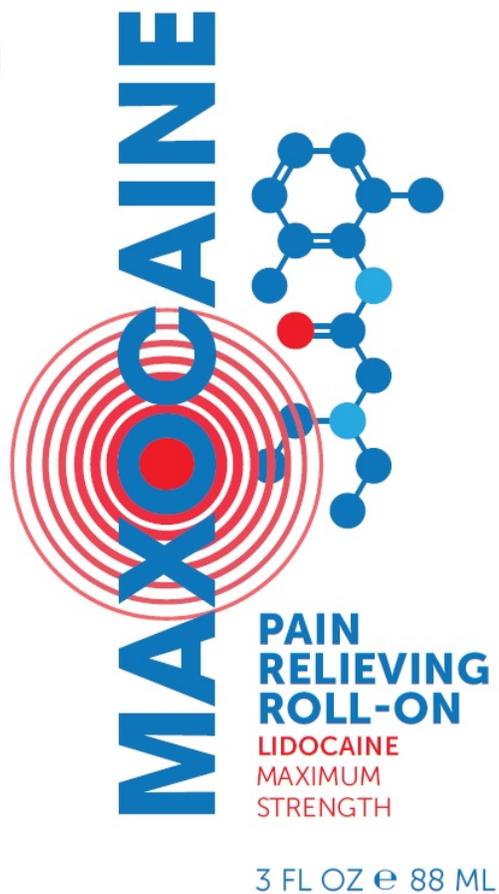
**Additional Information:**

Store at room temperature.

**Inactive Information:**

Acrylates C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Arnica Montana Extract, Boswellia Serrata Extract, Cetyl Alcohol, Ethylhexylglycerin, Ilex Paraguayensis (Yerba Mate') Extract, Isopropyl Alcohol, Isopropyl Myristate, Methylsulfonylmethane (MSM), Phenoxyethanol, Polysorbate-20, Sodium Lauryl Sulfate, Stearic Acid, Triethanolamine

**Package Labeling:**



**Drug Facts**

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|------------------------|--------------------|
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| Menthol 1% .....       | Topical anesthetic |

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Mfg. for Prime Commerce  
Irving, TX 75063 MADE IN USA  
For more information visit: [www.PrimeCommerce.net](http://www.PrimeCommerce.net)

**MAXOCAINE PAIN RELIEVING ROLL-ON**

lidocaine hydrochloride, menthol liquid

**Product Information**

|                                |                |                           |               |
|--------------------------------|----------------|---------------------------|---------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:72188-170 |
| <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 | 40 mg in 1 mL |
| <b>MENTHOL</b> (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)                   | MENTHOL                 | 10 mg in 1 mL |

### Inactive Ingredients

| Ingredient Name                                    | Strength |
|--|----------|
| <b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)           |          |
| <b>WATER</b> (UNII: 059QF0KO0R)                    |          |
| <b>ARNICA MONTANA</b> (UNII: O80TY208ZW)           |          |
| <b>INDIAN FRANKINCENSE</b> (UNII: 4PW41QCO2M)      |          |
| <b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)            |          |
| <b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)       |          |
| <b>ILEX PARAGUARIENSIS LEAF</b> (UNII: 1Q953B4O4F) |          |
| <b>ISOPROPYL ALCOHOL</b> (UNII: ND2M416302)        |          |
| <b>ISOPROPYL MYRISTATE</b> (UNII: 0RE8K4LNJS)      |          |
| <b>DIMETHYL SULFONE</b> (UNII: 9H4PO4Z4FT)         |          |
| <b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)           |          |
| <b>POLYSORBATE 20</b> (UNII: 7T1F30V5YH)           |          |
| <b>SODIUM LAURYL SULFATE</b> (UNII: 368GB5141J)    |          |
| <b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)             |          |
| <b>TROLAMINE</b> (UNII: 9O3K93S3TK)                |          |

### Packaging

| # | Item Code        | Package Description   | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:72188-170-03 | 88 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product | 03/01/2018           |                    |

### Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M017                                     | 03/01/2018           |                    |

**Labeler** - Prime Commerce, LLC (016785608)