**LIDOPLUS PAIN RELIEF- lidocaine hydrochloride cream**
**Centura Pharmaceuticals, Inc**

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**LidoPlus Pain Relief Cream**

***Drug Facts***

***Active Ingredients:***

Lidocaine HCl 4.00%

Topical Anesthetic

***Indications:***

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

***Warnings:***

**For external use only.**

• Avoid contact with eyes. • If condition worsens or symptoms persist for more than seven days, discontinue use and consult physician.

**Keep out of reach of children.**

If swallowed, consult physician.

**Do not use**

in large quantities, particularly over raw surfaces or blistered areas.

**If pregnant or breast feeding,**

contact physician prior to use.

***Directions:***

• Adults and children two-years of age or older: Apply to affected area not more than three to four times daily.

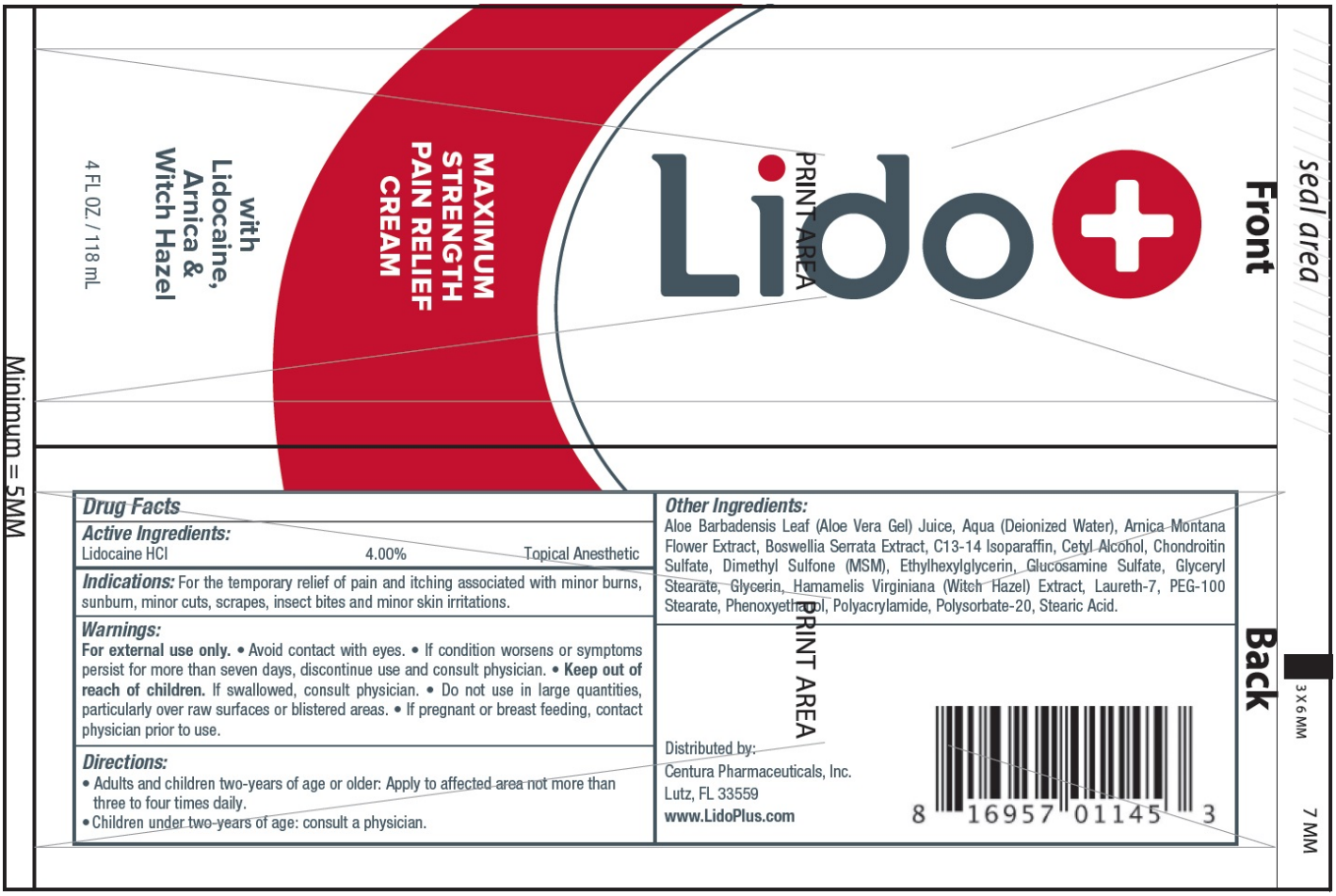
• Children under two-years of age: consult a physician.

***Other Ingredients:***

Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Arnica Montana Flower Extract, Boswellia Serrata Extract, C13-14 Isoparaffin, Cetyl Alcohol, Chondroitin

Sulfate, Dimethyl Sulfone (MSM), Ethylhexylglycerin, Glucosamine Sulfate, Glyceryl Stearate, Glycerin, Hamamelis Virginiana (Witch Hazel) Extract, Laureth-7, PEG-100 Stearate, Phenoxyethanol, Polyacrylamide, Polysorbate-20, Stearic Acid.

**Package Labeling:**




# LIDOPLUS PAIN RELIEF

lidocaine hydrochloride cream

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:70372-723 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **LIDOCAINE HYDROCHLORIDE** (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987) | LIDOCAINE HYDROCHLORIDE | 40 mg in 1 mL |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |

| | |
|---|---|
| **ALOE VERA LEAF** (UNII: ZY81Z83H0X) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **ARNICA MONTANA FLOWER** (UNII: OZ0E5Y15PZ) | |
| **INDIAN FRANKINCENSE** (UNII: 4PW41QCO2M) | |
| **C13-14 ISOPARAFFIN** (UNII: E4F12ROE70) | |
| **CETYL ALCOHOL** (UNII: 936JST6JCN) | |
| **ETHYLHEXYLGLYCERIN** (UNII: 147D247K3P) | |
| **GLUCOSAMINE SULFATE** (UNII: 1FW7WLR731) | |
| **GLYCERYL MONOSTEARATE** (UNII: 230OU9XXE4) | |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **HAMAMELIS VIRGINIANA TOP** (UNII: UDA30A2JJY) | |
| **LAURETH-7** (UNII: Z95S6G8201) | |
| **PEG-100 STEARATE** (UNII: YD01N1999R) | |
| **PHENOXYETHANOL** (UNII: HIE492ZZ3T) | |
| **POLYSORBATE 20** (UNII: 7T1F30V5YH) | |
| **STEARIC ACID** (UNII: 4ELV7Z65AP) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:70372-723-01 | 1 in 1 BOX | 02/07/2017 | |
| **1** | | 118 mL in 1 TUBE; Type 0: Not a Combination Product | | |
| **2** | NDC:70372-723-02 | 5 mL in 1 PACKET; Type 0: Not a Combination Product | 03/03/2024 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | M017 | 12/25/2015 | |

## Labeler - Centura Pharmaceuticals, Inc (084921637)

## Registrant - Centura Pharmaceuticals, Inc (084921637)

Revised: 1/2026 Centura Pharmaceuticals, Inc