LIDOPLUS PAIN RELIEF- lidocaine hydrochloride cream
Centura Pharmaceuticals, Inc

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.

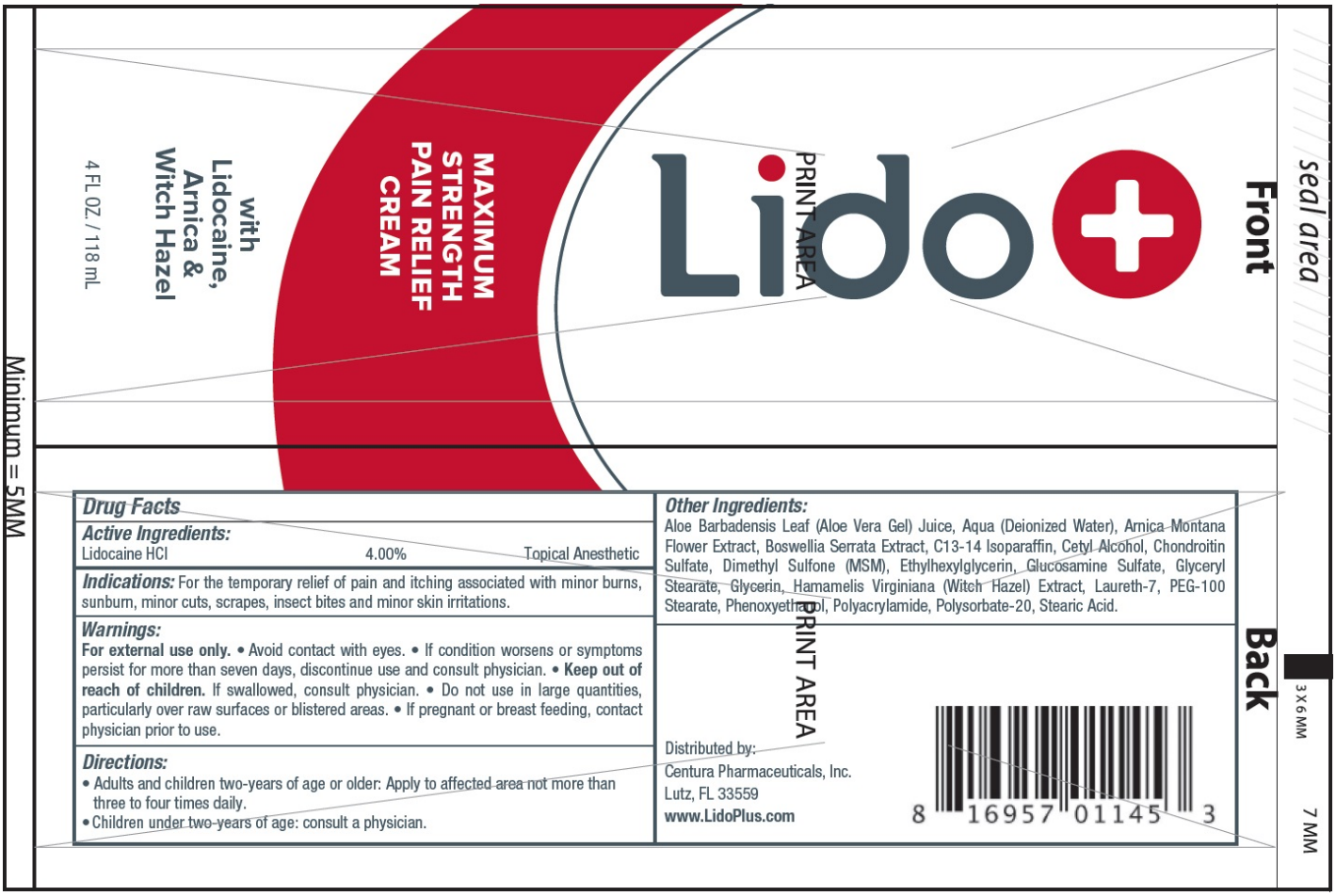
• Children under two-years of age: consult a physician.

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: 1/2024 Centura Pharmaceuticals, Inc