

CVS MAXIMUM STRENGTH LIDOCAINE PLUS- benzyl alcohol, lidocaine hydrochloride spray
CVS Pharmacy Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

CVS Health Lidocaine Plus

Active ingredients

Benzyl alcohol 10%

Lidocaine HCl 4%

Purpose

Topical anesthetic

Uses

For temporary relief of pain

Warnings

For external use only

Flammable:

Keep away from fire or flame

- Do not use near heat or flame or while smoking.
- avoid long term storage above 40 degree Celcius.
- do not puncture or incinerate. Contents under pressure
- do not store at temperatures above 49 degree Celcius

Do not use

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

When using this product

- use only as directed
- avoid contact with the eyes, mucous membranes or rashes
- avoid spraying on face
- avoid inhalation of spray

- do not bandage or apply local heat such as heating pads or a medicated patch to area of use
- do not use at the same time as other topical analgesics

Stop use and ask a doctor if

- skin reactions occur, such as rash, itching, redness, irritation, pain, swelling and blistering
- conditions worsen
- symptoms persist for more than 7 days
- symptoms clear up and occur again within a few days

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

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

Directions

Adults and children 12 years of age and over:

- spray to affected area not more than 3 to 4 times daily

Children under 12 years of age:

consult a doctor

Other information

- Avoid storing product in direct sunlight
- Protect product from excessive moisture
- Store with lid closed tightly

Inactive ingredients

alcohol denatured, arnica montana flower extract, calendula officinalis flower extract, camellia sinensis leaf extract, chamomilla recutita (matricaria) flower extract, dimethyl sulfone (msm), Echinacea angustifolia extract, ilex paraguariensis leaf extract, isopropyl myristate, juniperus communis fruit extract, purified water

Questions or comments?

Toll free **1-888-547-5492**

Principal Display Panel

Drug Facts

Active ingredients Purpose
Benzyl alcohol 10% Topical anesthetic
Lidocaine HCL 4% Topical anesthetic

Uses For temporary relief of 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 and mucous membranes avoid spraying on face avoid inhalation of spray do not bandage or apply local heat (such as heating pads) or a medicated patch to area of use do not use at the same time as other topical analgesics

Stop use and ask a doctor if

condition worsens redness is present irritation develops symptoms persist for more than 7 days or clear up and occur again within a few days

Extremely Flammable

do not use near heat or flame or while smoking avoid long term storage above 104°F (40°C) do not puncture or incinerate Contents under pressure do not store at temperatures above 120°F (49°C)

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

Keep out of reach of children.

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

Directions

Adults and children 12 years of age or over spray 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

Drug Facts (continued)

Inactive ingredients alcohol denat., arnica montana flower extract, calendula officinalis flower extract, camellia sinensis leaf extract, chamomilla recutita (matricaria) flower extract, dimethyl sulfone (msm), echinacea angustifolia extract, ilex paraguariensis leaf extract, isopropyl myristate, juniperus communis fruit extract, purified water

Questions or comments?

Call free 1-888-547-5492

Child resistant packaging.
Replace cap after each use.

Paraben & Propylene Glycol free

Not tested on animals

*Maximum strength lidocaine.

†This product is not manufactured or distributed by Hisamitsu Pharmaceutical Co., Inc., owner of the registered trademark Salonpas®.

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One CVS Drive, Woonsocket, RI 02895
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Made in the U.S.A. with U.S. and imported materials
V-36109



ACP# 91-60003-200-00



#383998

FPO 80%
UPC# 050428619155
X XXXXX XXXXXX X



Compare to the active ingredients in Salonpas® Lidocaine Plus Roll On†

MAXIMUM STRENGTH

Lidocaine Plus*

PAIN RELIEF SPRAY

**LIDOCAINE HCl 4%
BENZYL ALCOHOL 10%**

- Two powerful topical anesthetics
- Fast acting numbing relief
- Unscented



NET WT 3 OZ (85 g)

CVS MAXIMUM STRENGTH LIDOCAINE PLUS

benzyl alcohol, lidocaine hydrochloride spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69842-573
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZYL ALCOHOL (UNII: LKG8494WBH) (BENZYL ALCOHOL - UNII:LKG8494WBH)	BENZYL ALCOHOL	10 g in 100 g
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
CHAMOMILE (UNII: FGL3685T2X)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
ECHINACEA ANGUSTIFOLIA WHOLE (UNII: VB06AV5US8)	

ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)

ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)

JUNIPER BERRY (UNII: O84B5194RL)

WATER (UNII: 059QF0KO0R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-573-63	1 in 1 BOX	05/15/2020	
1		85 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	05/15/2020	

Labeler - CVS Pharmacy Inc. (062312574)

Revised: 12/2022

CVS Pharmacy Inc.