

SALONPAS LIDOCAINE PLUS- benzyl alcohol, lidocaine hydrochloride liquid
Hisamitsu America, Inc.

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

- on wounds or damaged skin
- in large quantities
- with a heating pad
- if you are allergic to any ingredients of this product

When using this product

- use only as directed
- avoid contact with the eyes, mucous membranes or rashes
- do not bandage tightly

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:

- clean and dry affected area
- apply 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

aloe barbadensis leaf juice, aminomethyl propanol, carbomer, SD alcohol 40-B, water

Questions or comments?

Toll free **1-800-826-8861** www.salonpas.us

Principal Display Panel

Hisamitsu

NDC#55328-901-03

No-Mess Roll On

Salonpas LIDOCAINE PLUS Pain Relieving Liquid

TOPICAL ANALGESIC

4% LIDOCAINE MAXIMUM STRENGTH *1
plus 10% Benzyl alcohol

2 Powerful Anesthetics

Fast Acting

Numbing Relief

Unscented

MADE IN USA

3 FL OZ (88 mL)

The logo for Hisamitsu, featuring a stylized 'H' symbol followed by the word 'Hisamitsu' in a bold, sans-serif font, with a registered trademark symbol (®) to the right.

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benzyl alcohol, lidocaine hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55328-901
Route of Administration	TOPICAL, PERCUTANEOUS, TRANSDERMAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZYL ALCOHOL (UNII: LKG8494WBH) (BENZYL ALCOHOL - UNII:LKG8494WBH)	BENZYL ALCOHOL	10.0000 g in 100 g
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4.0000 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
alcohol (UNII: 3K9958V90M)	
TERT-BUTYL ALCOHOL (UNII: MD83SFE959)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
water (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55328-901-03	1 in 1 BOX	09/01/2017	
1		80 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 Drug	M017	09/01/2017	
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Labeler - Hisamitsu America, Inc. (191877802)

Revised: 12/2024

Hisamitsu America, Inc.