

LIDOCAINE PAIN RELIEF ROLL-ON- lidocaine 4% liquid
CVS Pharmacy

CVS Lido Roll-On

Adults and children 2 years of age and older: apply to the affected area no more than 3 to 4 times daily.

Children under 2 years of age: consult a doctor.

acrylates/C10-30 alkyl acrylate crosspolymer, aloe barbadensis leaf juice, aminomethyl propanol, C30-45 alkyl cetearyl dimethicone crosspolymer, caprylyl methicone, cetearyl alcohol, ceteth-20 phosphate, dicetyl phosphate, dimethicone, disodium EDTA, ethylhexylglycerin, glyceryl stearate, hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer, methylparaben, polysorbate 60, SD alcohol 40 (15%), steareth-2, steareth-21, water

Lidocaine 4%

For External Use Only.

When using this product:

use only as directed

do not bandage tightly or use with a heating pad

avoid contact with eyes

do not apply to wounds or damaged skin

do not use in large quantities

partifcularly over raw surfaces or blistered areas

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

For temporarily relief of pain and itching.

Topical analgesic



MAXIMUM STRENGTH
Lidocaine Pain
Relief Roll-On
LIDOCAINE 4%
Odor Free

Drug Facts

Active ingredient Purpose
Lidocaine 4% Topical anesthetic

Uses For temporary relief of pain and
itching

Warnings

For external use only.

When using this product • use only
as directed • do not bandage tightly
• avoid contact with eyes • do not
apply to wounds or damaged skin • do
not use in large quantities, particularly
over raw surfaces or blistered areas.

Stop use and ask a doctor if •
condition worsens • symptoms persist
for more than 7 days or clear up and
occur again within a few days.

Keep out of the reach of children. If
swallowed, get medical help or contact
a Poison Control Center immediately.

Directions

Adults and children 2 years of age
and older: apply to the affected area
no more than 3 to 4 times daily.

Children under 2 years of age:
consult a doctor.

Inactive ingredients

Acrylates/C10-30 Alkyl
Crosspolymer, Aloe Barbadensis Leaf
Extract, Aminomethyl Propanol,
C30-45 Alkyl Cetearyl Dimethicone
Crosspolymer, Caprylyl Methicone,
Cetearyl Alcohol, Ceteleth-20
Phosphate, Dicapyl Phosphate,
Dimethicone, Disodium EDTA,
Ethyhexylglycerin, Glyceryl Stearate,
Methylparaben, SD Alcohol 40,
Steareth-21, Purified Water.

*This product is not manufactured or distributed
by Glaxo, Inc., the distributor of Aspercreme®.

#256563



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Compare to the active ingredient
in Aspercreme® Odor Free
with 4% Lidocaine*



MAXIMUM STRENGTH Lidocaine Pain Relief Roll-On

LIDOCAINE 4%

Odor Free

- Temporary relief of pain
- Helps numb pain-affected
areas without irritation
- No-touch applicator



Package Contains One Bottle

Actual Size



Actual Bottle
Size on
Side Panel

2.5 FL OZ (74 mL)

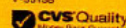
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Distributed by: CVS Pharmacy, Inc.
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CVS.com 1-800-SHOP-CVS
V-39136





LIDOCAINE PAIN RELIEF ROLL-ON

lidocaine 4% liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69842-029
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z 41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
WATER (UNII: 059QF0KO0R)	
STEARETH-21 (UNII: 53J3F32P58)	
ALCOHOL (UNII: 3K9958V90M)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
CAPRYLYL TRISILOXANE (UNII: Q95M2P1KJL)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
DIHEXADECYL PHOSPHATE (UNII: 2V6E5WN99N)	
CETETH-20 PHOSPHATE (UNII: 921FTA1500)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%) (UNII: 86FQE96TZ4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-029-97	1 in 1 CARTON	06/05/2019	
1		74 mL in 1 BOTTLE, WITH APPLICATOR; 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	06/05/2019	

Labeler - CVS Pharmacy (062312574)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	manufacture(69842-029) , analysis(69842-029) , label(69842-029) , pack(69842-029)