

**LIDOCAINE PAIN RELIEF ROLL-ON- lidocaine 4% liquid**  
**Humco Holding Group, Inc.**

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**Private Label Lidocaine 4% 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

**CVS Label**



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

CVS Health

Drug Facts	
<b>Active ingredient</b>	<b>Purpose</b>
Lidocaine 4%	Topical analgesic
<b>Uses</b> For temporary relief of pain and itching	
<b>Warnings</b> 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.	
<b>Stop use and ask a doctor if:</b> condition worsens; symptoms persist for more than 7 days or clear up and occur again within a few days.	
<b>Keep out of the reach of children.</b> If swallowed, get medical help or contact a Poison Control Center immediately.	
<b>Directions</b>	
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.	
<b>Inactive ingredients</b>	
Acrylates/C10-30 Alkyl Crosspolymer, Aloe Barbadensis Leaf Extract, Aminomethyl Propanol, C30-45 Alkyl Cetearyl Dimethicone Crosspolymer, Caprylyl Methicone, Cetearyl Alcohol, Cetyl-20 Phosphate, Dicalyl Phosphate, Dimethicone, Disodium EDTA, Erythexylglycerin, Glyceryl Stearate, Methylparaben, SD Alcohol 40, Stearalk-21, Purified Water.	

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



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



**MAXIMUM STRENGTH**  
**Lidocaine Pain Relief Roll-On**

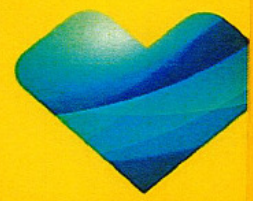
**LIDOCAINE 4%**  
**Odor Free**

- 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)**

**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

Distributed by: CVS Pharmacy, Inc.  
 One CVS Drive, Woonsocket, RI 02895  
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 CVS.com 1-800-6SHOP-CVS  
 9-39138









**Equate Label**



## LIDOCAINE PAIN RELIEF ROLL-ON

lidocaine 4% liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0395-9130
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 g in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>AMINOMETHYLPROPANOL</b> (UNII: LU49E6626Q)	
<b>WATER</b> (UNII: 059QF0KO0R)	

<b>STEARETH-21</b> (UNII: 53J3F32P58)
<b>ALCOHOL</b> (UNII: 3K9958V90M)
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)
<b>CAPRYLYL TRISILOXANE</b> (UNII: Q95M2P1KJL)
<b>CARBOMER INTERPOLYMER TYPE A (55000 CPS)</b> (UNII: 59TL3WG5CO)
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)
<b>EDETATE DISODIUM ANHYDROUS</b> (UNII: 8NLQ36F6MM)
<b>DIHEXADECYL PHOSPHATE</b> (UNII: 2V6E5VN99N)
<b>CETETH-20 PHOSPHATE</b> (UNII: 921FTA1500)
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)
<b>CETOSTEARYL ALCOHOL</b> (UNII: 2DMT128M1S)
<b>HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%)</b> (UNII: 86FQE96TZ4)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0395-9130-97	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** - Humco Holding Group, Inc. (825672884)

**Registrant** - Pharma Nobis, LLC (118564114)

### Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	manufacture(0395-9130) , analysis(0395-9130) , label(0395-9130) , pack(0395-9130)