LANSINOH PAIN RELIEVING- lidocaine hcl spray Lansinoh Laboratories Inc

Lansinoh Pain Relieving Spray

DRUG FACTS

Active Ingredient	Purpose
Lidocaine 4%	Topical Analgesic

STOP USE AND ASK A DOCTOR

Stop use and ask a doctor if

- Condition worsens
- Symptoms persist for more than 7 days
- Symptoms clear up and return within a few day

KEEP OUT OF REACH OF CHILDREN

KEEP OUT OF REACH OF CHILDREN. Do not use on infants or children under 12.

WARNINGS

Warnings For extern

For external use only.

Do NOT use on breasts, nipples, in or near mouth.

KEEP OUT OF REACH OF CHILDREN. Do not use on infants or children under 12. If swallowed, get medical help or contact a Poison Control Center right away.

Stop use and ask a doctor if

- Condition worsens
- Symptoms persist for more than 7 days
- Symptoms clear up and return within a few days

Directions ■ For adult use only

■ Apply externally to the affected area not more than 3 - 4 times in a 24 hour period.

Inactive Ingredients:

Alcohol (Ethanol), Aloe Barbadensis Leaf Juice, Cellulose Gum, Eugenol, Hamamelis Virginiana (Witch Hazel) Water, Lavandula Angustifolia (Lavender) Oil, Mentha Piperita

(Peppermint) Oil, PEG-40 Hydrogenated Castor Oil, Poloxamer 188, Potassium Sorbate, Prunus Amygdalus Dulcis (Sweet Almond) Oil, Sodium Benzoate, Sodium Phytate, Water.

Distributed By: Lansinoh Laboratories, Inc

Alexandria, VA 22314, USA

Tel: +1-800-292-4794 | www.lansinoh.com

FOR EXTERNAL USE ONLY

For external use only

Topical Analgesic



Lansinoh.

SPRAY PARA ALIVIAR EL **DOLOR**

CON LIDOCAÍNA AL 4 % SPRAY POSPARTO

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Drug Facts

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Ingredients

Purpose

Uses Temporarily relieves pain

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 Condition worsens Symptoms
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 Symptoms clear up and return
 within a few days

WHEN USING THIS PRODUCT

Avoid contact with eyes
 Do not use in large quantities, particularly over raw surfaces or blistered areas.

Directions

■ For adult use only ■ Apply externally to the affected area not more than 3 - 4 times in a 24 hour

Inactive Ingredients:

Inactive Ingredients:
Alcohol (Ethanol), Aloe
Barbadensis Leaf Juice, Cellulose
Gum, Eugenol, Hamamelis
Virginiana (Witch Hazel) Water,
Lavandula Angustifolia (Lavender)
Oil, Mentha Piperita (Peppermint)
Oil, PEG-40 Hydrogenated Castor
Oil, Poloxamer 188, Potassium
Sorbate, Prunus Amygdalus Dulcis
(Sweet Almond) Oil, Sodium
Benzoate, Sodium Phytate, Water.

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care.

This collection of effective postpartum essentials is backed by research and guidance from experts. Trust the brand that moms and doctors alike

> We are with you, Mama.



Lansinoh.

PAIN RELIEF SPRAY

POSTPARTUM SPRAY

Relieves pain with blend of herbal witch hazel and aloe vera



Dermatologically & **Gynecologically Tested**

3.5oz (100 ml) €





LOT

Lansinoh.

PAIN RELIEF **SPRAY**

WITH 4% LIDOCAINE POSTPARTUM SPRAY





360° sprayer for easy,



Gynecologically Tested



LANSINOH PAIN RELIEVING

lidocaine hcl spray

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:53997-001

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient NameBasis of StrengthStrengthLIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)LIDOCAINE4 g in 100 mL

Inactive Ingredients Ingredient Name Strength PHYTATE SODIUM (UNII: 88496G1ERL) ALOE VERA WHOLE (UNII: KIZ 4X2EHYX) PEG-40 CASTOR OIL (UNII: 4ERD2076EF) EUGENOL (UNII: 3T8H1794QW) ALCOHOL (UNII: 3K9958V90M) WITCH HAZEL (UNII: 101I4J0U34) WATER (UNII: 059QF0KOOR) SODIUM BENZOATE (UNII: 0]245FE5EU)

POLOXAMER 188 (UNII: LQA7B6G8JG)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
ALMOND OIL (UNII: 66YXD4DKO9)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
CARBOXYMETHYLCELLULOSE SODIUM (0.7 CARBOXYMETHYL SUBSTITUTION PER SACCHARIDE; 2300 MPA.S AT 1%) (UNII: 0Z2R7OG99L)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	

F	Packaging						
#	t Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:53997- 001-02	1 in 1 CARTON	02/15/2021				
1	NDC:53997- 001-01	100 mL in 1 BOTTLE, PLASTIC; 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	02/15/2021			

Labeler - Lansinoh Laboratories Inc (181696907)

Establishment							
Na me	Address	ID/FEI	Business Operations				
Dhaliwal Pharmaceuticals Laboratories LLC		116933772	manufacture(53997-001)				

Revised: 3/2024 Lansinoh Laboratories Inc