INSTANT BURN RELIEF ALOE EXTRA CVS- lidocaine 0.50% spray CVS

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.

Actives

Lidocaine USP .50%

Purpose

External analgesic

Uses

Temporarily relieves pain and itching due to:

- sunburn
- minor burns
- scrapes
- insect bites
- minor irritations

Warnings

For external use only.

Flammable: Do not use while smoking or near heat or flame

Do not use

in large quantities, particularly over raw surfaces or blistered areas

When using this product

- keep out of eyes
- use only as directed
- do not puncture or incinerate. Contents under pressure. Do not store at temperatures above 120F

Stop use and ask a doctor if

- conditions gets worse
- symptoms last more than 7 days
- symptoms clear up and occur again in a few days

Keep out of reach of children.

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

Directions

- shake well
- adults and children 2 years of age and older: apply to affected area not more than 3-4 times daily

- children under 2 years of age: ask a doctor
- to apply to face, spray into palm of hand and gently apply

Inactive ingredients

Aloe Barbadensis Leaf Juice, SD Alcohol 40, Propylene Glycol, Glycerin, Simethicone, Tocopheryl Acetate (Vitamin E Acetate), Triethanolamine, Carbomer, Diazolidinyl Urea, Methylparaben, Propylparaben, Disodium Cocoamphodipropionate, Disodium EDTA.

CVS

pharmacy

INSTANT

BURN RELIEF

SPRAY

ALOE EXTRA

- instant pain relief with Lidocaine
- · Cooling, moisturizing itch relief

Net WT 4.5 OZ (127g)



lidocaine 0.50% spray

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59779-230	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
LIDO CAINE (UNII: 98 PI200987) (LIDO CAINE - UNII:98 PI200987)	LIDOCAINE	.50 g in 100 g		

Inactive Ingredients	
Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
ALCOHOL (UNII: 3K9958V90M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6 A3C0 OX)	
.ALPHATO CO PHERO L ACETATE (UNII: 9E8X80D2L0)	
TROLAMINE (UNII: 9O3K93S3TK)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
METHYLPARABEN (UNII: A2I8 C7HI9 T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
DISO DIUM CO CO AMPHO DIPRO PIO NATE (UNII: 6 K8 PRP39 7 M)	
EDETATE DISO DIUM (UNII: 7FLD9 1C86K)	

	Packaging			
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:59779-230-04	127 g in 1 CAN; Type 0: Not a Combination Product	11/11/20 14	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part348	11/11/2014		

Labeler - CVS (062312574)

Registrant - Product Quest Mfg, LLC (927768135)

Establishment				
Name	Address	ID/FEI	Business Operations	
Product Quest Mfg, LLC		927768135	manufacture(59779-230), label(59779-230)	

Revised: 11/2017 CVS