# PAIN RELIEF EQUATE- lidocaine hcl 4% cream Walmart

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.

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### **Drug Facts**

## Active ingredient Purpose

Lidocaine HCl 4%.....Topical analgesic

#### 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 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 affected area not more than 3 to 4 times daily. Children under 2 years of age: consult a doctor.

#### **Inactive Ingredients**

Acrylates/C10-30 Alkyl Acrylate Crosspolymer

Aloe Barbadensis Leaf Extract

Aminomethyl Propanol

C30-45 Alkyl Cetearyl Dimethicone Crosspolymer

Caprylyl Methicone

Cetearyl Alcohol

Ceteth-20 Phosphate

Dicetyl Phosphate

Dimethicone

Disodium EDTA

Ethylhexylglycerin

Glyceryl Stearate

Methylparaben

SD Alcohol 40

Steareth-21

Water



# PAIN RELIEF EQUATE

lidocaine hcl 4% cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-287
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
LIDO CAINE HYDRO CHLO RIDE (UNII: V13007Z41A) (LIDO CAINE - UNII:98PI200987)	LIDOCAINE	4 g in 100 mL	

Inactive Ingredients	
Ingredient Name	Strength

ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
CAPRYLYL TRISILO XANE (UNII: Q95M2P1KJL)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETETH-20 PHO SPHATE (UNII: 921FTA1500)	
DIHEXADECYL PHO SPHATE (UNII: 2V6E5WN99N)	
DIMETHICO NE (UNII: 92RU3N3Y1O)	
EDETATE DISO DIUM (UNII: 7FLD91C86K)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERYL MONOSTEARATE (UNII: 230 O U9 XXE4)	
METHYLPARABEN (UNII: A2I8 C7HI9 T)	
STEARETH-21 (UNII: 53J3F32P58)	
WATER (UNII: 059QF0KO0R)	

Packaging			
# Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date
1 NDC:49035-287-03	1 in 1 CARTON	05/29/2017	
1	74 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	05/29/2017	

# Labeler - Walmart (051957769)

# Registrant - Product Quest Mfg (927768135)

Establishment			
Name	Address	ID/FEI	Business Operations
Product Quest Mfg		927768135	manufacture(49035-287), label(49035-287)

Revised: 12/2017 Walmart