## ALOE GEL PAIN RELIEVER WELL AT WALGREENS- lidocaine hydrochloride 0.5% gel Walgreens

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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#### WALGREENS BURN RELIEF ALOE GEL

#### **Active ingredient**

#### **Purpose**

Lidocaine Hydrochloride 0.5%

Pain Reliever

#### Uses for temporary relief of pain and itching due to

- sunburn
- minor burns
- insect bites
- cuts
- scrapes

#### Warnings

#### For external use only

#### When using this product

• avoid contact with eyes. Rinse with water if contact occurs.

#### Stop use and ask a doctor if

• symptoms last more than 7 days.

#### Keep out of reach of the children

#### **Directions**

- adults and children 2 years of age and older: apply to affected area, not more than 3 to 4 times a day
- children under 2 years of age: consult a physician

#### **Inactive ingredients**

Aloe Barbadensis Leaf Juice, Blue 1, Carbomer, Diazolidinyl Urea, Disodium EDTA, Glycerin, Isopropyl Alcohol, Menthol, Polysorbate 80, Propylene Glycol, Triethanolamine, Water, Yellow 5





#### ALOE GEL PAIN RELIEVER WELL AT WALGREENS

TOPICAL

lidocaine hydrochloride 0.5% gel

Route of Administration

# Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:0363-3192

#### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>LIDO CAINE HYDRO CHLO RIDE</b> (UNII: V13007Z41A) (LIDO CAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	0.5 g in 100 g

Inactive Ingredients		
Ingredient Name	Strength	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
CARBOMER HOMOPOLYMER TYPE C (UNII: 4Q93RCW27E)		
Diazolidinyl Urea (UNII: H5RIZ3MPW4)		
EDETATE DISO DIUM (UNII: 7FLD9 1C8 6 K)		
Glycerin (UNII: PDC6A3C0OX)		
Isopropyl Alcohol (UNII: ND2M416302)		
Menthol (UNII: L7T10EIP3A)		
Polysorbate 80 (UNII: 6OZP39ZG8H)		
Propylene Glycol (UNII: 6DC9Q167V3)		
TROLAMINE (UNII: 9O3K93S3TK)		
Water (UNII: 059QF0KO0R)		
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)		

l	Packaging				
l	# Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>	
l	1 NDC:0363-3192-06	227 g in 1 BOTTLE; Type 0: Not a Combination Product	03/15/2013		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part348	03/15/2013		

## Labeler - Walgreens (008965063)

### Registrant - Product Quest Mfg (927768135)

Establishment			
Name	Address	ID/FEI	Business Operations
Product Quest Mfg		927768135	manufacture(0363-3192), label(0363-3192)

Revised: 2/2018 Walgreens