BURN RELIEF ALOE DAYLOGIC- lidocaine hydrochloride 0.5% gel
Rite Aid

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.

WALGREENS BURN RELIEF ALOE GEL

Active ingredient	**Purpose**
Lidocaine Hydrochloride 0.5%	External analgesic

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

BURN RELIEF ALOE DAYLOGIC

lidocaine hydrochloride 0.5% gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11822-7782
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	0.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER HOMOPOLYMER TYPE C (UNII: 4Q93RCW27E)	
Diazolidinyl Urea (UNII: H5RIZ3MPW4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-7782-6	237 g in 1 BOTTLE; Type 0: Not a Combination Product	08/30/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	08/30/2016	

Labeler - Rite Aid (014578892)

Registrant - Product Quest Mfg (927768135)

Establishment

Name	Address	ID/FEI	Business Operations
Product Quest Mfg		927768135	manufacture(11822-7782) , label(11822-7782)

Revised: 5/2018 Rite Aid