

**EQUATE BURN RELIEF- lidocaine hydrochloride gel**  
**Walmart Inc**

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**Walmart 79903-237-08**

**Active ingredient**

Lidocaine HCl      0.5%

**Purpose**

External analgesic

**Uses**

for the temporary relief of pain and itching associated with

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

**Warnings**

**For external use only.**

**Do not use** in large quantities, particularly over raw surfaces or blistered areas

**When using this product** avoid contact with eyes

**Stop use and ask a doctor** if condition worsens, or if 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 right away.

**Directions**

- adults and children 2 years of age and older: apply to affected area no more than 3 to 4 times daily
- children under 2 years of age: ask a doctor

**Inactive ingredients**

Aloe Barbadensis Leaf Juice, Blue 1, Carbomer, Disodium EDTA, Ethylhexylglycerin, Glycerin, Maltodextrin, Menthol, Phenoxyethanol, Polysorbate 20, Propanediol, SD Alcohol, Sodium Hydroxide, Water, Yellow 5



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associated with ■ minor burns ■ sunburn ■ minor cuts ■ scrapes ■ insect bites ■ minor skin irritations

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***Questions?*** 1-888-287-1915

**DISTRIBUTED BY: Walmart Inc., Bentonville, AR 72716**  
**PRODUCT OF CHINA**

\*This product is not manufactured or distributed by Bayer Healthcare, LLC, owner of the registered trademark Solarcaine® Cool Aloe Burn Relief Formula.  
Satisfaction guaranteed - For questions or comments please call 1-888-287-1915.

## **EQUATE BURN RELIEF**

lidocaine hydrochloride gel

### **Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:79903-237
<b>Route of Administration</b>	TOPICAL		

### **Active Ingredient/Active Moiety**

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

## Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ALOE</b> (UNII: V5VD430YW9)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>CARBOMER 934</b> (UNII: Z135WT9208)	
<b>DISODIUM HEDTA</b> (UNII: KME849MC7A)	
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)	
<b>MENTHOL</b> (UNII: L7T10EIP3A)	
<b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>POLYSORBATE 20</b> (UNII: 7T1F30V5YH)	
<b>PROPANEDIOL</b> (UNII: 5965N8W85T)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	
<b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-237-08	226 g in 1 BOTTLE; Type 0: Not a Combination Product	11/15/2023	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/15/2023	

**Labeler -** Walmart Inc (051957769)

## Establishment

Name	Address	ID/FEI	Business Operations
Nantong Health & Beyond Hygienic Products Inc.		421280161	manufacture(79903-237)

Revised: 11/2023

Walmart Inc