

**ASSURED- lidocaine hydrochloride gel**  
**Zhejiang Jingwei Pharmaceutical Co., Ltd.**

-----  
**Drug Facts**

Inactive Ingredients aloe barbadensis leaf juice, carbomer 940, glycerin, menthol, methylparaben, polyethylene glycol 400, propylene glycol, propylparaben, triethanolamine, water

Active Ingredient

Lidocaine hydrochloride 2.5 %

Purpose

Topical analgesic

Uses for temporary relief of pain associated with

- minor burns
- sunburn
- minor cuts or scrapes
- 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

- if condition worsens, or if symptoms persist for more than 7 days or clear up and then occur again within a few days, discontinue use of this product and consult a physician

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away

Directions

adults and children under 2 years of age:

- clean the affected area and gently dry
- apply to the affected area not more than 3 to 4 times daily

children under 2 years of age: consult a physician

Other Information

store at room temperature 59 - 86 F (15 C - 30 C )



COMPARE TO  
AFTERBURN<sup>®</sup>

**ASSURED** **ASSURED**

# Burn Relief Gel

• Lidocaine hydrochloride 2.5%  
Topical Anesthetic

**Quickly Cools & Soothes on Contact**

- Used for:**
- Sunburns
  - Windburns
  - Kitchen Burns
  - Other Minor Burns, Cuts & Scrapes

NET WT  
0.7 OZ (20 g)

**Drug Facts**

Active ingredient	Purpose
Lidocaine hydrochloride 2.5%	Topical Anesthetic

**Uses** for temporary relief of pain associated with:

- minor burns • scrapes
- minor cuts or abrasions • minor skin irritations

**Warnings**  
For external use only.

**Do not use**

- in large quantities, particularly over raw surfaces or lacerated 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 then occur again within a few days, discontinue use of this product and consult a physician

**Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.**

**Directions**  
adults and children over 2 years of age

- apply to affected area and gently rub
- apply to the affected area with thick coats 3 to 4 times daily
- children under 2 years of age, consult a physician

**Other information**  
Store at room temperature for -10° F (5° F - 30° F).

**Inactive ingredients** also contains: butadiene, butyl alcohol, carbomer 974, glycerin, menthyl, methylparaben, polyethylene glycol 400, propylene glycol, propylparaben, triethanolamine, water

23003 951  
DISTRIBUTED BY  
GENESSEE  
MEDICAL SUPPLY, INC.  
208 VINEY PARKWAY  
GIESSEN, VA 22088  
MADE IN CHINA

100% WATER SOLUBLE  
NO RUBBER BANDS  
NO ADHESIVE  
NO PAIN

3927763511

WITH ALDE VERB WITH ALDE VERB WITH ALDE VERB

**Burn Relief**  
**Burn Relief**  
**Burn Relief**

ASSURED ASSURED ASSURED

152.4 mm

63.5 mm

25.4 mm

127 mm

8mm

# ASSURED

lidocaine hydrochloride gel

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:55621-010
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	2.5 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
<b>CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED)</b> (UNII: 4Q93RCW27E)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>MENTHOL, UNSPECIFIED FORM</b> (UNII: L7T10EIP3A)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>WATER</b> (UNII: 059QF0KO0R)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55621-010-01	1 in 1 BOX	10/30/2014	
1		20 g in 1 TUBE; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	10/30/2014	

**Labeler** - Zhejiang Jingwei Pharmaceutical Co., Ltd. (530876549)

## Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Jingwei Pharmaceutical Co., Ltd.		530876549	manufacture(55621-010)

Revised: 10/2023

Zhejiang Jingwei Pharmaceutical Co., Ltd.