LIDOCAINE HYDROCHLORIDE 2%- lidocaine hydrochloride gel
Aldermed Inc.

Lidocaine Hydrochloride 2% Gel

Active Ingredient

Lidocaine Hydrochloride 2%

Purpose

External Analgesic

Use(s)

■ For the temporary relief of pain associated with ■ Minor burns ■ Sunburn

Warnings

For External Use Only

Do not use

■ On wounds or damaged skin

■ In large quantities, particularly over raw surfaces or blistered areas

When using this product

■ Avoid contact with the eyes

■ Do not bandage tightly

Stop use and ask a doctor if

■ Condition worsens

■ Symptoms persist for more than 7 days

■ Symptoms 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.(1-800-222-1222)

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

Other Information

- Store at room temperature
- Tamper Evident. Do not use if seal is torn, cut, or opened.
- See sachet or cartonfor lot number and expiry date

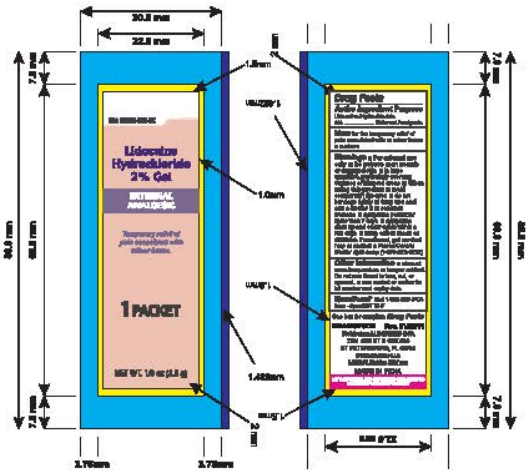
Inactive Ingredients

Acrylates/C10-30 Alkyl Acrylate Crosscopolymer, Carbomer, Glycerin, Imidurea, Methylparaben, Propylene Glycol, Propylparaben, Purified Water, Tea Tree Leaf Oil, Trolamine

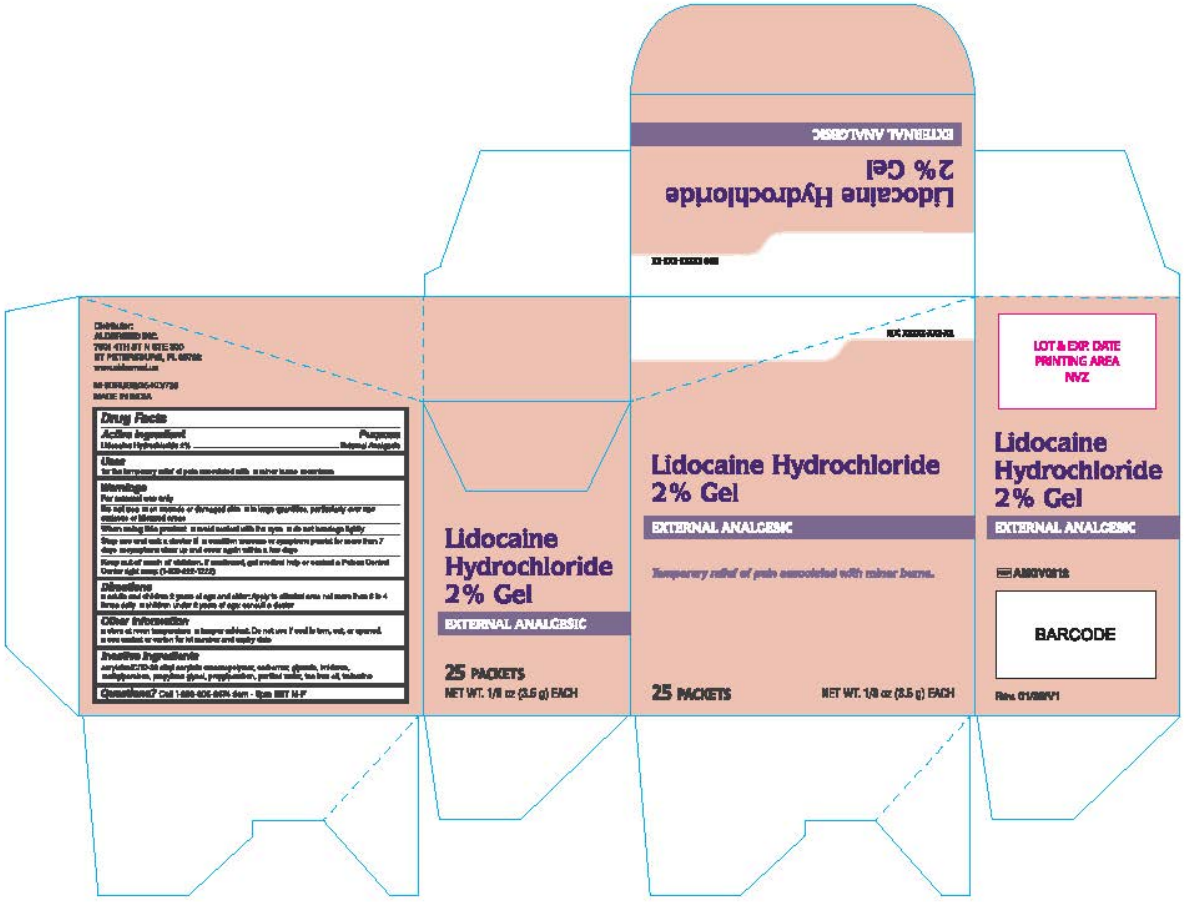
Questions?

1-888-396-2739 Monday - Friday, 9AM - 5PM EST

Label


Lidocaine Hydrochloride 2% Gel
1 PACKET

Size: (L) 73mm x (W)44mm x (H)88.9 mm



LIDOCAINE HYDROCHLORIDE 2%

lidocaine hydrochloride gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:87236-015
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE -	LIDOCAINE HYDROCHLORIDE	2 g

UNII:98PI200987)	ANHYDROUS	in 100 g

Inactive Ingredients

Ingredient Name	Strength
TROLAMINE (UNII: 9O3K93S3TK)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
GLYCERIN (UNII: PDC6A3C0OX)	
IMIDUREA (UNII: M629807ATL)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
TEA TREE OIL (UNII: VIF565UC2G)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
CARBOMER 940 (UNII: 4Q93RCW27E)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87236-015-35	25 in 1 BOX	01/01/2026	
1		3.5 g in 1 PACKET; 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	01/01/2026	

Labeler - Aldermed Inc. (144887712)

Revised: 1/2026 Aldermed Inc.