

ALOCANE PLUS- lidocaine hydrochloride and benzalkonium chloride gel
Quest Products, Inc.

Alocane[®] Plus

Drug Facts

Active Ingredient	Purpose
Lidocaine HCL 4%	Topical Analgesic
Benzalkonium Chloride 0.13%	Topical Antiseptic

Uses

For the temporary relief of pain and itching associated with: Sunburn, Minor Cuts & Scrapes, Minor Skin Irritation, Minor Burns

Warnings

For external use only. Avoid contact with eyes.

Keep out of reach of children.

Stop use and ask doctor: if condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days.

If pregnant or breast-feeding, ask a health care professional before use.

Directions

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Apply to affected area not more than 3 to 4 times daily. Do not use in large quantities, particularly over raw surfaces or blistered areas.

Children under 2 years of age, consult a physician.

Other Information

Store at Room Temperature of 15° - 30°C (59° - 86°F)

Inactive Ingredients

1,3-Propanediol, Aloe Barbadensis (Aloe) Leaf Juice, Caprylyl Glycol, Chlorphenesin, Dimethyl Isosorbide, Hydroxyethyl Cellulose, Phenoxyethanol, Tocopheryl Acetate (Vitamin E), Water.

DISTRIBUTED BY

QUEST PRODUCTS LLC

PRINCIPAL DISPLAY PANEL - 59 mL Tube Carton

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ALOCANE PLUS

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68229-300
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 g in 100 mL
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
PROPANEDIOL (UNII: 5965N8W85T)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
DIMETHYL ISOSORBIDE (UNII: SA6A6V432S)	
HYDROXYETHYL CELLULOSE (5500 MPA.S AT 2%) (UNII: M825OX60H9)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68229-300-02	50 in 1 BOX	12/11/2013	07/12/2021
1	NDC:68229-300-01	3.55 mL in 1 DOSE PACK; Type 0: Not a Combination Product		
2	NDC:68229-300-03	1 in 1 CARTON	03/01/2020	
2		59 mL in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:68229-300-04	946 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	07/09/2021	07/15/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	12/11/2013	

Labeler - Quest Products, Inc. (075402441)

Establishment

Name	Address	ID/FEI	Business Operations
Fill Tech USA		926433855	manufacture(68229-300)

Revised: 7/2025

Quest Products, Inc.