ALOCANE PLUS- lidocaine hydrochloride and benzalkonium chloride gel
Quest Products LLC

Alocane ® Plus

Drug Facts

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

Uses:

First aid to help prevent bacterial infection associated with contamination of the skin. Temporarily relieves pain and itching due to:

- Minor Burns
- Minor Cuts & Scrapes
- Minor Skin Irritations

Warnings

For external use only. Avoid contact with eyes.

Keep out of reach of children. In case of accidental ingestion, seek medical help or call a Poison Control Center right away.

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

Adults and children 2 years of age and older: Apply to affected areas not more than 3 to 4 times daily.

Children under 2 years of age: consult a doctor.

Other Information

Store at room temperature 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.

PRINCIPAL DISPLAY PANEL - 48 packet carton

ALOCANE PLUS

lidocaine hydrochloride and benzalkonium chloride gel

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:68229-300-06	48 in 1 CARTON	11/12/2020	03/31/2026
1	NDC:68229-300-05	3.55 mL in 1 DOSE PACK; 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	05/01/2020	

Labeler - Quest Products LLC (075402441)

Revised: 12/2025 Quest Products LLC