

APOLLO ANTISEPTIC HAND SANITIZER- ethyl alcohol gel
Seber Design Group, Llc

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

apollo™ Antiseptic Hand Sanitizer

Drug Facts

Active ingredient

Ethyl alcohol 70% v/v

Purpose

Antiseptic

Uses • hand sanitizer to decrease bacteria on the skin • recommended for repeated use • for use when soap and water are not available

Warnings

Flammable, keep away from fire/flame

For external use only

Do not use • in children less than 2 months of age • on open skin wounds

When using this product • do not get into eyes. In case of contact, rinse eyes thoroughly with water

Stop use and ask a doctor if • irritation and redness develop • condition persists for more than 72 hours

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

Directions • wet hands thoroughly with product and allow to dry without wiping • supervise children under 6 years of age when using this product to avoid swallowing

Other information • store between 15-30°C (59-86°F) • avoid freezing and excessive heat above 40°C (104°F)

Inactive ingredients aloe vera (Aloe barbadensis) leaf juice, carbomer, triethanolamine, glycerin, propylene glycol, tocopheryl acetate, water

Questions or comments? +1-877-584-1434

You may also report serious side effects to this phone number. Mon-Fri 9:00 AM - 5:00 PM

Antiseptic Hand Sanitizer

- **Moisturizing**
- **Ethyl alcohol 70%**
- **Topical solution**

Distributed by: Seber Design Group, LLC, 964 E. Badillo St, #501, Covina, CA 91724


COUNTRY OF ORIGIN: CHINA

11-236-B

(L) 102004001 EXP 04/2022

Packaging



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APOLLO ANTISEPTIC HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:77279-164

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
TROLAMINE (UNII: 9O3K93S3TK)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77279-164-64	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/01/2020	

Labeler - Seber Design Group, Llc (078349592)

Revised: 5/2020

Seber Design Group, Llc