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 TM Antiseptic Hand Sanitizer

Drug Facts

Active ingredient

Ethyl alcohol 75% 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 aqua, acrylates/c10-30 alkyl acrylate cross polymer, glycerin, tocopherol.

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

- Moisturizing
- Topical solution

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

COUNTRY OF ORIGIN: CHINA

Packaging



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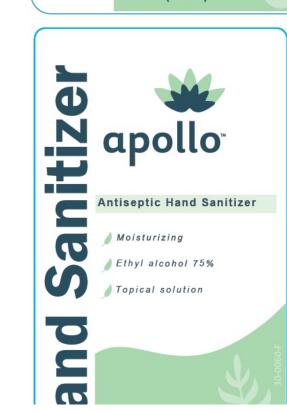
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i apollo **Antiseptic Hand Sanitizer**

Moisturizing

3.99 fl oz (118 mL)

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COUNTRY OF ORIGIN: CHINA







Antiseptic Hand Sanitizer

Moisturizing

8.45 fl oz (250 mL)

Drug Facts

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COUNTRY OF ORIGIN: CHINA



apollo

Antiseptic Hand Sanitizer

Moisturizing

33.8 fl oz (1L)

Drug Facts

Active ingredient

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APOLLO ANTISEPTIC HAND SANITIZER

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77279-375	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
GLYCERIN (UNII: PDC6A3C0OX)	
TOCOPHEROL (UNII: R0ZB2556P8)	

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:77279-375-20	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020		
2	NDC:77279-375-21	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020		
3	NDC:77279-375-22	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020		
4	NDC:77279-375-23	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020		
5	NDC:77279-375-24	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020		
6	NDC:77279-375-28	1000 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