APOLLO INSTANT HAND SANITIZER- instant hand sanitizer gel Sbu Group L.P.

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 Instant Hand Sanitizer

Active ingredient

Alcohol 75%v/v

Purpose

Antiseptic

Uses:

Hand sanitizer to help decrease bacterial on the skin, when water, soap& towel are not available.

Recommended for repeated use.

Warnings

For external use only. Flammable. Keep away from fire or flame.

Do not apply around eyes. Do not use in ears & mouth.

When using this product,

Avoid contact with eyes. In case of contact flush eyes with water.

Stop use and ask a doctor

If redness or irritation develop and persist for more than 72 hours

Keep out of reach of children.

Children must be supervised in use of this product.

Directions

Place enough product into your palms and thoroughly spread on both hands.

Rub into skin until dry.

Other information

Store below 110°F 43°C

May discolor certain fabrics or surfaces.

Inactive ingredients

Water, Glycerin, Triethanolamine, Acrylates/C10-30 Alkyl Acrylate Crosspolymer.

Package Label

apollo

Instant Hand Sanitizer

Anti Bacterial

Contains 75 % Alcohol

2 FL OZ (60ml)

Kills 99% Germs

NDC: 78303-001-01





Drug Facts

Active ingredient Purpose: Alcohol 75% v/v.....Antiseptic

Uses • Hand sanitizer to help decrease bacterial on the skin, when water, soap & towel are not available. • Recommended for repeated use.

Warnings: For external use only.

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Distributed by Apollo Distributors 10510 Rockley Road Houston, Texas 77099 Phone 281-564-6464 EXP :

Batch Number :

For emergency contact: +12815646464 Manufactured by: GUANGZHOU XUELEI COSMETIC CO.,LTD.

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Package Label

apollo

Instant Hand Sanitizer

Anti Bacterial

Contains 75 % Alcohol

1.29 Gallons

5,000 ML

Kills 99% Germs

NDC: 78303-001-02



APOLLO INSTANT HAND SANITIZER

instant hand sanitizer gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78303-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL	

Inactive Ingredients			
Ingredient Name	Strength		
TROLAMINE (UNII: 903K93S3TK)			
CARBOMER COPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 809Y72KV36)			
WATER (UNII: 059QF0KO0R)			
GLYCERIN (UNII: PDC6A3C0OX)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78303- 001-01	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/06/2020	
2	NDC:78303- 001-02	5000 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/06/2020	

Marketing Information				
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	05/06/2020		

Labeler - Sbu Group L.P. (841530095)

Registrant - GUANGZHOU XUELEI COSMETIC CO.,LTD. (526885026)

Establishment			
Name	Address	ID/FEI	Business Operations
GUANGZHOU XUELEI COSMETIC CO.,LTD.		526885026	manufacture(78303-001) , pack(78303-001) , label(78303-001)

Revised: 11/2021 Sbu Group L.P.