

SANIXUSA HAND SANITIZER- ethyl alcohol 75% gel
GUANGZHOU RAINHOME PHARM&TECH CO., LTD

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.

Sanixusa Hand Sanitizer

Active Ingredient

Ethyl Alcohol 75%

Purpose

Antiseptic

Uses

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available

Warnings

For External Use Only. FLAMMABLE. Keep away from heat or flame.

Do not use

In children less than 2 months of age. On open skin wounds.

When using this product

Keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor.

If irritation or rash occurs. These may be signs of a serious condition.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Place enough product on hands to cover all surfaces. Rub hands together until dry. Supervise children under 6 years of age when using this product to avoid swallowing

Other Information

Store between 15-30C (59-85F).

Avoid freezing and excessive heat above 40C (104F)

Inactive Ingredients

Water, C10-30 alkyl acrylate crosspolymer, aminomethyl propanol, glycerin, aloe barbadensis (leaf) extract

Sanixusa Hand Sanitizer



NEW FORMULA

SaniXUSA

75% Alcohol

PREMIUM HAND SANITIZER

WITH ALOE LEAF EXTRACT

kills 99.99% of germs*

- ✓ NEW effective formula
- ✓ No water required
- ✓ Color free
- ✓ Perfume free

www.sanixusa.com **16.90oz (500ml)**

SANIXUSA ANTIBACTERIAL HAND SANITIZER

Drug Facts	
Active Ingredient [s]	Purpose
Ethyl Alcohol 75% w/v	Antiseptic
Use [s]	
Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.	
Warnings	
For external use only. Flammable. Keep away from heat or flame.	
Do not use:	
• in children less than 2 months of age	
• on one skin wounds	
When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse thoroughly with water.	
Stop using and seek doctor if irritation or rash occurs. These may be signs of a serious condition.	
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	
Directions	
• Place enough product on hands to cover all surfaces.	
• Rub hands together until dry.	
• Supervise child 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 Water, C10-30 alkyl acrylate cross polymer, aminomethyl propanol, glycerin, aloe barbadensis (leaf) extract	



Made in China
 Manufacturer: Guangzhou Rainbow Pharmaceutical Co., Ltd.
 2/F, Building 15, No. 10, Yongheng Road, Wange Economic Zone, Guangzhou Development District, Guangzhou, Guangdong Province, P.R. China

Imported & Distributed by
 Chills Beverages USA
 6452 East Rogers Drive, Boca Raton, Florida 33487

* Conducted in laboratory setting and not actual use.

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www.sanixusa.com **33.81oz (1litre)**

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SANIXUSA HAND SANITIZER

ethyl alcohol 75% gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78 126-110
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
WATER (UNII: 059QF0K00R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78 126-110-03	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
2	NDC:78 126-110-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	

3	NDC:78 126-110-06	200 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
4	NDC:78 126-110-16	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
5	NDC:78 126-110-33	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 - GUANGZHOU RAINHOME PHARM&TECH CO., LTD (547018779)

Registrant - Chilla Beverages USA (078572585)

Revised: 5/2020

GUANGZHOU RAINHOME PHARM&TECH CO., LTD