ASSURED ALOE HAND SANITIZER 8 OZ.- ethyl alcohol gel China Ningbo Shangge Cosmetic Technology Corp.

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.

Assured Aloe Hand Sanitizer 8 oz.

Active Ingredient Purpose

Ethyl Alcohol 70.0% v/v..... Antimicrobial

□Uses

- for handwashing to decrease bacteria on the skin
- recommended for repeated use

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

□Assured

Instant Hand Sanitizer

Kills 99% of germs!

Aloe Vera & Moisturizers

Compare to Active Ingredients in Purell® Hand Sanitizer*

Special Value 25% more free

8 fl oz / 237 ml

*This product is not manufactured or distributed by Gojo Industries Inc, distributor of the registered trademark Purell

|| Directions

- wet hands thoroughly with product
- briskly rub hands together until dry
- supervise children under 6 years in the use of this product

□ Warnings

IFor external use only.

IFlammable, keep away from heat and flame.

□Do not use in the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation and redness develop and persist for more than 72 hours.

Inactive Ingredients

water, carbomer, triethanolamine, glycerin, propylene glycol, fragrance, aloe barbadensis leaf juice, fd&c blue no. 1, fd&c yellow no. 5.





ASSURED ALOE HAND SANITIZER 8 OZ.

ethyl alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:58503-109

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
WATER (UNII: 059QF0KO0R)	
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)	
TROLAMINE (UNII: 9O3K93S3TK)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging						
# Item Code	Package Description	Marketing Start Date	Marketing End Date			
1 NDC:58503-109-01	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/0 1/20 18				
Marketing Information						
Marketing Catego	ry Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not fin	nal part333E	12/0 1/20 18				

Labeler - China Ningbo Shangge Cosmetic Technology Corp. (529287434)

Establishment							
Name	Address	ID/FEI	Business Operations				
China Ningbo Shangge Cosmetic Technology Corp.		529287434	manufacture(58503-109)				

Revised: 11/2018 China Ningbo Shangge Cosmetic Technology Corp.