

ASSURED™

25%
MORE
FREE

Instant Hand Sanitizer

Kills 99%
of germs!



COMPARE TO ACTIVE INGREDIENTS IN
Purell® Hand Sanitizer*

Special Value 25% more free

10 fl oz / 295 ml

Drug Facts

Active ingredient

Ethyl Alcohol 62.0%.....Antimicrobial

Purpose

Uses • for handwashing to decrease bacteria on the skin
• recommended for repeated use

Warnings

For external use only.

Flammable, 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.

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 • briskly rub hands together until dry • supervise children under 6 years in the use of this product

Other information

- store at 20°C to 25°C (68° to 77°F)
- may discolor certain fabrics

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

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

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DISTRIBUTED BY:
GREENBRIER INTERNATIONAL, INC.
500 VOLVO PARKWAY, CHESAPEAKE, VA 23320
MADE IN CHINA

ASSURED ALOE

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58503-081
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	62 g in 100 g	
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: ZY81Z83H0X)				
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:58503-081-01	295 g in 1 BOTTLE; Type 0: Not a Combination Product	07/02/2015	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	07/02/2015		

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

Establishment			
Name	Address	ID/FEI	Business Operations
China Ningbo Shangge Cosmetic Technology Corp.		529287434	manufacture(58503-081)

Revised: 9/2019

China Ningbo Shangge Cosmetic Technology Corp.