GHOPE HANDS SANITIZER- ethyl alcohol liquid
Gns Usa Inc.

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.

***Ghope* Hands Sanitizer**

Drug Facts

Active ingredients

Ethyl Alcohol 70.0%

Purpose

ANTISEPTIC

Uses

for hand-washing to decrease bacteria on the skin. only when water is not available

Warnings

Flammable. Keep away from fire and flames. For external use only.

When using this product • Do not get into eyes.

• If contact occurs, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or redness develops.

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 • For children under 6, use only under adult supervision.

Inactive Ingredients

Purified Water, Aloe Extract, Glycerin, Sodium Hyaluronate, Carbomer, Butylene Glycol, Triethanolamine, Flavoring

Questions?

+1-703-562-1020 or visit http://ghope-ppe.com

Leaves Hands Feeling Soft

Kills 99.9% of germs with alcohol formula

Ethanol 70% Advanced Refresh Hand Sanitizer

GNS USA INC Centreville, VA 20121

Manufactured for

GNS USA INC

MADE IN KOREA

Packaging

GHOPE HANDS SANITIZER

ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77521-500
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)	
ALOE (UNII: V5VD430YW9)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77521-500-01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/11/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/11/2020	

Labeler - Gns Usa Inc. (080435020)

Revised: 7/2020 Gns Usa Inc.