GUSTAR HAND SANITIZER (ETHANOL)- alcohol gel GuStar 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.

Drug Facts

ACTIVE INGREDIENT

Ethyl alcohol 70%

INACTIVE INGREDIENTS

Aloe Vera Gel, Grapefruit Seed Extract, Disodium EDTA, Butylene Glycol, Aminomethyl Propanol, Carbomer, Phenoxyethanol, Purified Water

PURPOSE

Antiseptic

WARNINGS

- For external use only.
- Flammable. Keep away from heat and Flame.

When using this product

- Keep out of eyes, ears and mouth.
- If contact occurs, rinse eyes thoroughly with water

Stop use and ask a doctor

■ If irritation and rash occurs.

KEEP OUT OF REACH OF CHILDREN

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

Uses

Hand sanitizer to help reduce bacteria on the skin. Only when water and soap is not available.

Directions

- Wet hands thoroughly with product and rub hands together until dry.
- Supervise children under 6 years of age when using

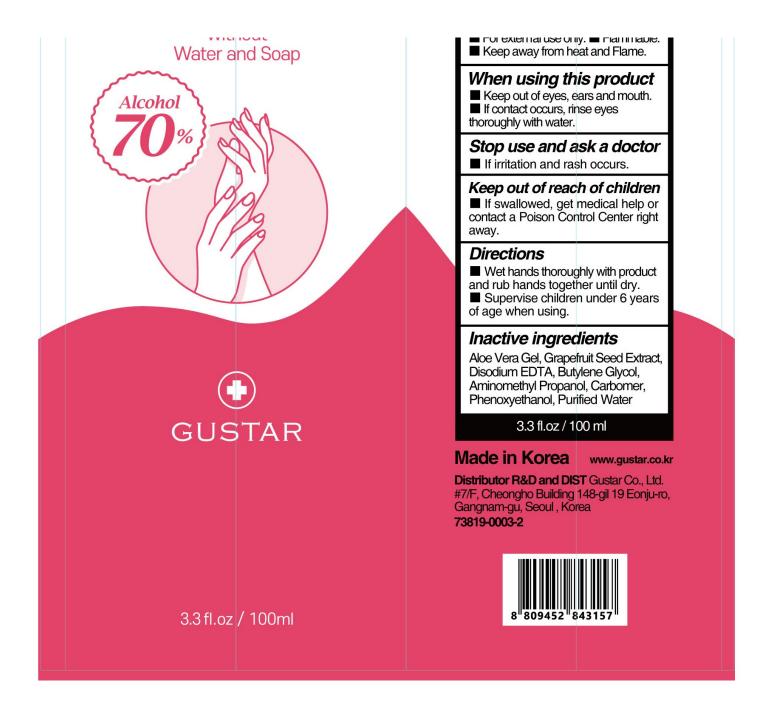
Package Label: GUSTAR HAND SANITIZER 40mL





Package Label: GUSTAR HAND SANITIZER 100mL





Package Label: GUSTAR HAND SANITIZER 500mL



GUSTAR HAND SANITIZER (ETHANOL)

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73819-0003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 g in 100 mL	

Inactive Ingredients			
Ingredient Name	Strength		
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)			
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)			
WATER (UNII: 059QF0KO0R)			
GRAPEFRUIT SEED OIL (UNII: 598 D944HOL)			

l	Packaging			
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:73819-0003-1	40 mL in 1 POUCH; Type 0: Not a Combination Product	03/21/2020	
l	2 NDC:73819-0003-2	100 mL in 1 TUBE; Type 0: Not a Combination Product	03/21/2020	
l	3 NDC:73819-0003-3	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/21/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	03/21/2020		

Labeler - GuStar Co Ltd (694234583)

Registrant - GuStar Co Ltd (694234583)

Establishment			
Name	Address	ID/FEI	Business Operations
THE SUN CO.LTD		694433104	manufacture(73819-0003)

Revised: 5/2020 GuStar Co Ltd